

Methodology for Deriving Standards for Contaminants in Soil to Protect Human Health

New Zealand Government

This report may be cited as: Ministry for the Environment. 2011. *Methodology for Deriving Standards for Contaminants in Soil to Protect Human Health*. Wellington: Ministry for the Environment.

Published in June 2011 by the Ministry for the Environment Manatū Mō Te Taiao PO Box 10362, Wellington 6143, New Zealand

ISBN: 978-0-478-37237-3

Publication number: ME 1055

Other publications in this series include: Toxicological Intake Values for Priority Contaminants in Soil

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This document is available on the Ministry for the Environment's website: www.mfe.govt.nz



Acknowledgements

This report was prepared for the Ministry for the Environment by Graeme Proffitt (Pattle Delamore Partners Limited). The sections of the report relating to plant bioconcentration factors for each contaminant, and produce consumption (Appendix 3), were derived from reports prepared by Jo-Anne Cavanagh (Landcare Research).

The advice and assistance of the Ministry's Technical Advisory Group for this report is acknowledged. The members were:

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Executive Summary

This technical report sets out a risk-based methodology for deriving soil contaminant concentrations protective of human health. Together with the *Toxicological Intake Values for Priority Contaminants in Soil* (MfE, 2011), this document serves as a technical reference in support of the National Environmental Standard for Assessing and Managing Contaminants in Soil to Protect Human Health and should be read in conjunction with it (see http://www.mfe.govt.nz/laws/standards/contaminants-in-soil/index.html).

This report addresses a question that is fundamental to the science of contaminated land management: What contaminant concentration in soil can people be exposed to and yet not be subject to an appreciable risk of harm? The answer to this question varies internationally because each jurisdiction frames its response to fit within unique risk policy and legislative frameworks. So, although the technical approach to risk assessment of contaminated land is shared broadly by most countries, there are significant differences in detail such that a standard adopted by one country may not suit another.

As an alternative to adopting standards from another country, the Ministry for the Environment has examined the science of risk assessment and compiled a derivation methodology for healthbased standards to apply to soil contaminants in New Zealand under the Resource Management Act 1991. This initiative comes at a time when it is appropriate also to review the soil guideline values contained within the Ministry's existing suite of contaminated land guidelines. The 'soil contaminant standards' contained in this technical report are intended to supersede the 'soil acceptance criteria' used in previous New Zealand guidelines; the new methodology also resolves technical differences between them. The Ministry intends to apply the same approach when reviewing the petroleum hydrocarbon contaminants.

The term 'soil contaminant standards' to protect human health, or $SCSs_{(health)}$, specifically refers to soil contaminant concentrations that are mandatory, under the *National Environmental Standard for Assessing and Managing Contaminants in Soil to Protect Human Health.* $SCSs_{(health)}$ may be applied as Tier 1 or screening criteria; as conservative clean-up targets, to inform on-site management actions; or to trigger further investigation within a Tier 2 assessment. When talking about generic numerical values in guidelines or foreign jurisdictions, or for soil contaminant concentrations that are derived on a site-specific basis according to this methodology, the term soil guideline values (SGVs) is used.

If $SCSs_{(health)}$ or $SGVs_{(health)}$ are exceeded, it may result in health effects that are more than minor for some people, and as such are unacceptable. Conversely, if actual soil concentrations are less than or equal to the $SCSs_{(health)}$ or $SGVs_{(health)}$ then this is judged to be acceptable, because any adverse effects on human health for most people are likely to be no more than minor.

The methodology in this document is government policy and has two applications. First it demonstrates how the standards for contaminants in soil were derived for use within the NES regulation as 'soil contaminant standards', $SCSs_{(health)}$. Secondly, it sets out the basis for deriving soil contaminant concentrations protective of human health for exposure scenarios that lie outside of the generic exposure scenarios under the NES. The purpose of this second application of the methodology is to guide the site-specific derivation of soil guideline values, $SSGVs_{(health)}$, when there is good reason to use a site-specific risk assessment.

It is important to note that the methodology and the derived standards are intended to be protective of human health only, and do not apply to other environmental receptors. When it is relevant to protect other valued elements of the environment, separate consideration of appropriate values to achieve this is required.

This report presents:

- a national risk-based methodology for deriving soil contaminant concentrations protective of human health
- a suite of numerical criteria for priority contaminants that are legally binding as gazetted under the *National Environmental Standard for Assessing and Managing Contaminants in Soil to Protect Human Health*
- background information on the risk assessment methodologies and exposure parameters.

 $SCSs_{(health)}$ are derived for the following priority contaminants: arsenic, boron, cadmium, chromium, copper, inorganic lead, inorganic mercury (but not elemental mercury), benzo(a)pyrene (representing the carcinogenic polycyclic aromatic hydrocarbons), DDT (as the sum of DDT, DDD and DDE), dieldrin, dioxin (as 2,3,7,8-tetrachlorodibenzo-*p*-dioxin (TCDD) and dioxin-like polychlorinated biphenyls (PCBs)), and pentachlorophenol (PCP).

The SCS_(health) calculations draw on toxicological intake values and background exposures set out in the companion document, *Toxicological Intake Values for Priority Contaminants in Soil* (MfE, 2011). Numerical values are calculated for five generic land-use scenarios, and utilise standardised receptors and exposure parameters.

A summary of the $SCSs_{(health)}$ derived is presented in Tables ES1 and ES2. Contaminated-land practitioners are referred to the more detailed version of these tables set out in section 7 of this report, in which additional residential sub-scenarios have also been derived.

	Arsenic Boron		Cadmium	Chromium		Copper	Inorganic	Inorganic
			(pH 5) ⁻	III ²	VI		lead	mercury
Rural residential / lifestyle block 25% produce	17	NL	0.8	NL	290	NL	160	200
Residential 10% produce	20	NL	3	NL	460	NL	210	310
High-density residential	45	NL	230	NL	1,500	NL	500	1,000
Recreation	80	NL	400	NL	2,700	NL	880	1,800
Commercial / industrial outdoor worker	70	NL	1,300	NL	6,300	NL	3,300	4,200

 Table ES1: Summary of soil contaminant standards – SCSs_(health) – for inorganic substances (mg/kg)

1 Default value is for pH 5. Values increase with increasing pH (see Appendix 2).

2 The SCSs_(health) for boron, chromium III and copper represent levels well in excess of concentrations that would affect the health of plants.

NL = No limit. This is where the derived values exceed 10,000 mg/kg.

Scenario	BaP DDT Dieldrin PCP		PCP	Dioxi	Dioxin (µg/kg TEQ)	
					TCDD	Dioxin-like PCBs
Rural residential / lifestyle block 25% produce	6	45	1.1	55	0.12	0.09
Residential 10% produce	10	70	2.6	55	0.15	0.12
High-density residential	24	240	45	110	0.35	0.33
Recreation	40	400	70	150	0.6	0.52
Commercial / industrial outdoor worker	35	1,000	160	360	1.4	1.2

Table ES2: Summary of soil contaminant standards – SCSs_(health) – for organic compounds (mg/kg¹)

1 All values in mg/kg dry weight except dioxins which are in μ g/kg.

1 Introduction

1.1 Background

The Ministry for the Environment (MfE) has determined the need for a better policy framework for managing contaminated land in New Zealand. After extensive public consultation, the Ministry published a position paper in September 2007 (MfE, 2007). The paper identified, among other things, an inconsistent and variable use by contaminated-land practitioners of numerical contaminated land guidelines used to assess the risk that contaminated soil might pose to human health. The Ministry then determined that, as a matter of priority, it would develop:

- a national risk-based methodology for deriving soil contaminant concentrations protective of human health
- a suite of numerical criteria for priority contaminants as examples of the national methodology
- site management options and actions that follow from applying the above criteria; ie, the numerical criteria may:
 - serve as Tier 1 or screening criteria to assess whether there is a potential risk to human health
 - when the numerical criteria are exceeded, serve as conservative clean-up targets for many situations, ie, where further investigation or site-specific risk assessment is not warranted or economic
 - inform on-site management actions to reduce the potential for adverse effects
 - trigger further investigation to better assess the risk and/or determine site-specific criteria as a Tier 2 assessment.

The Ministry intends that the derivation methodology be incorporated by reference into the *National Environmental Standard for Assessing and Managing Contaminants in Soil to Protect Human Health* (NES), as was proposed in the Discussion Document (*Proposed National Environmental Standard for Assessing and Managing Contaminants in Soil*, MfE, 2010a). While the technical detail behind the newly derived soil contaminant standards is presented within this separate methodology document, the soil contaminant standards will be applied in accordance with the NES and within the context of the derivation methodology.

This technical report introduces and sets out the risk-based methodology the Ministry has adopted as government policy for deriving SCSs_(health): soil contaminant standards to protect human health. SCSs_(health) are derived for a limited group of priority contaminants that are of primary concern in New Zealand. These are specifically: arsenic, boron, cadmium, chromium, copper, inorganic lead, inorganic mercury, benzo(a)pyrene, DDT, dieldrin, pentachlorophenol, and dioxin (as 2,3,7,8-tetrachlorodibenzo-*p*-dioxin, and dioxin-like polychlorinated biphenyls). These calculations draw on toxicological values and background exposures set out in a companion document *Toxicological Intake Values for Priority Contaminants in Soil* (MfE, 2011).

The numerical values are restricted to consideration of human-health risks and are based on conservative standard land-use scenarios, using standardised receptors and exposure parameters.

The consideration of exposure pathways includes (following MfE, 2007):

- estimating the levels of contaminants in the media (eg, air, water, food) that potentially convey the contaminants from soil to people (the 'receptor')
- identifying the typical physical characteristics of New Zealanders (such as area of skin, weight, air breathed, food and water ingested) that collectively determine a standard exposure model in association with the exposure pathways. This requires consideration of the:
 - groups of people (eg, children, adults, workers) who are potentially exposed, as well as the group considered the most sensitive to the toxic effect of each contaminant
 - time periods over which exposure occurs
 - age-related contact rates such as ingestion and inhalation rates
 - consumption of produce from their own home gardens.

The United Kingdom, Australia and elsewhere have recently reviewed the partitioning and vapour migration models used to derive guidelines for the inhalation of volatile contaminants. Therefore, this pathway is not considered in detail and no standards for volatile contaminants have been derived. It is beyond the scope of, and premature for, this document to review or derive more appropriate inhalation models. Further research is being carried out internationally on applying such models.

A note on the use of terms:

- The term 'soil contaminant standards', SCSs_(health), specifically refers to those numerical concentrations that have regulatory status under the National Environmental Standard for Assessing and Managing Contaminants in Soil to Protect Human Health.
- For ease of reference, SCSs_(health) are often referred to in this document simply as SCSs, or soil contaminant standard(s). Where a health protection context is thought useful, the term SCSs_(health) is retained.
- SCSs_(health) may be applied as: Tier 1 or screening criteria; as conservative clean-up targets, to inform on-site management actions; or to trigger further investigation within a Tier 2 assessment.
- When talking about generic numerical values in guidelines or foreign jurisdictions, or for soil contaminant concentrations that are derived on a site-specific basis according to this methodology, the term 'soil guideline values' (SGVs) is used.

1.2 Purpose

This technical report sets out in a transparent manner the methodology as to how the suite of soil contaminant standards was derived. $SCSs_{(health)}$ are soil contaminant concentrations appropriate to five generic land-use exposure scenarios at which the exposure to levels equal to or less than the SCSs is judged to be acceptable, because any adverse effects on human health for most people are likely to be no more than minor.

In addition to the five standard scenarios, some additional residential sub-scenarios have been derived for illustrative purposes. The additional residential scenarios are for proportions of home-grown produce other than the standard 10 and 25 per cent. The additional derivations show the effect on the derivations for zero and 50 per cent home-grown produce and may be used where site-specific derivation of soil guideline values is warranted.

SCS_(health) derivations are given for:

- seven elements arsenic, boron, cadmium, chromium (in trivalent and hexavalent forms), copper, inorganic lead and inorganic mercury
- five organic compounds, or groups of compounds benzo(a)pyrene (to represent the carcinogenic polycyclic aromatic hydrocarbons); DDT (as the sum of DDT and its metabolites DDD and DDE); dieldrin (or aldrin or the sum of aldrin and dieldrin); pentachlorophenol, and dioxin (including tetrachlorodibenzo-*p*-dioxin (to represent dioxin mixtures) and dioxin-like polychlorinated biphenyls).

1.3 Document organisation

This document is divided into the following parts: After this introduction (section 1), background information on the principles of the soil quality guidelines derivation is presented in section 2. Section 3 reviews and recommends generic exposure scenarios. Soil guideline derivation equations are provided in section 4. The proposed exposure pathways, and specific parameters for these pathways for each exposure scenario, are given in section 5, with toxicological values. The equations and parameters, together with contaminant-specific values from MfE (2011) are then used to derive soil contaminant standards for selected priority contaminants in section 6. A summary of the derived standards is provided in section 7. Limitations of the use of the soil contaminant standards are given in section 8. An explanation of the methodology for site-specific risk assessments is provided in section 9.

The details pertaining to $SCS_{(health)}$ calculations for each individual contaminant in accordance with exposure assumptions are set out in appendix 1, with pH-dependant details provided for cadmium in appendix 2, and the details for dioxin with egg consumption in appendix 4. Appendix 3 sets out the produce consumption estimates made. Details of SGV derivation methodologies for selected overseas jurisdictions, and specific derivation methodologies for existing New Zealand guidelines, are given in appendix 5. Appendix 6 contains topsoil datasets for arsenic and cadmium used to calculate the national background levels. This is followed by a glossary and a comprehensive references section.

2 General Approach to Deriving Soil Guideline Values

2.1 Concepts in risk assessment

Soil guideline values are generic quality standards adopted in many countries to regulate the management of contaminated land. They are usually in the form of concentration thresholds (mg/kg soil dry weight) of contaminants in soil, above which certain actions are recommended or enforced. The implications of exceeding the soil guidelines vary according to the regulatory framework of the particular national or regional jurisdiction. They range from the need for further investigations to the need for remedial actions. Given the different purposes in national regulatory frameworks, soil guidelines have been given various names (translated as appropriate): trigger values, reference values, target values, intervention values, clean-up values, cut-off values, and many others (Carlon, 2007).

Generic risk-based human-health soil guideline values are fundamental to the risk assessment process for contaminated land. Risk assessment is a process in which information is analysed to determine if an environmental hazard might cause harm to exposed persons and ecosystems (US EPA, 2004a). In this case the environmental hazard is land contamination. The risk assessment process is a multi-step process (Defra and EA, 2002a) consisting of:

- 1. defining the conceptual exposure model
- 2. collecting human exposure characteristics and contaminant fate and transport data
- 3. selecting contaminant concentration in soil from review of site investigation data
- 4. quantitative modelling of exposure
- 5. comparing predicted human exposure with health criteria values, based on contaminant toxicity
- 6. evaluating significance of risk to human health where exposure is close to or exceeds health-based reference values.

The above process is complex and time-consuming if carried out from first principles on every site. Multiple-exposure scenarios, receptors, exposure mechanisms and chemicals must be evaluated to determine their significance. Thousands of pages of guidance have been written by authorities such as the United States Environmental Protection Agency (US EPA) to ensure consistency of process. Most sites do not warrant the detail or expenses of a full risk assessment, which is why many jurisdictions have developed 'screening level' assessment processes using standardised exposure scenarios. The basis of screening assessments (otherwise known as Tier 1 assessments) is the use of SGVs derived for these standardised exposure scenarios.

The assessment process then becomes a much simpler process of taking soil samples (in accordance with, for example: MfE, 2004), analysing them for relevant contaminants and comparing the results with an appropriate soil guideline value. If the sample results are below the SGV, then the risk to human health is deemed acceptable; if the results exceed the SGV, a potential risk to human health is indicated and some form of action or management may be necessary. For most sites the assessment process stops there: the site has been dealt with as appropriate. However, for some sites it is worth making a more accurate assessment, using the more detailed site-specific assessment process (often called Tier 2 assessment).

The generic values are deliberately conservative to be protective of the great majority of potentially exposed people (often defined as the 95th percentile of the population). However, for some large sites this could lead to very expensive remediation or other forms of site management to limit exposure. In these situations the additional expense of a more detailed and less conservative assessment process may be far outweighed by the cost saving in remediation or management.

It is important that the SGVs_(health) are not too conservative, as they may be used as clean-up guidelines where the cost of further assessment is not warranted. If screening guidelines are too conservative, then more clean-up than is warranted could occur. The US EPA uses the concept of reasonable maximum exposure (RME) in combining upper-bound and average exposure factors to arrive at exposure scenarios that are protective but reasonable (US EPA, 1989b, 2004a). Under this approach, some intake variables may not be at their individual maximum values, but in combination with other variables will result in estimates of the RME. However, the US EPA has been criticised for this approach on the basis that the RME approach combines too many upperbound parameter estimates, resulting in unrealistic over-conservative assessments (US EPA, 2004a).

Key components of the generic human health guidelines include standard human exposure scenarios relevant to a variety of land uses (eg, residential, commercial, parkland) and exposure through a variety of pathways (eg, inhalation, ingestion, skin absorption). The soil guideline values are then calculated to ensure that some pre-determined allowable daily intake of the contaminant is not exceeded. The allowable daily intakes (otherwise known as a reference health standard – RHS) are discussed in greater detail in MfE (2011). These scenarios and exposure pathways are explored in subsequent sections.

The Canadian Council of Ministers of the Environment has developed a useful set of guiding principles for the development of SGVs (CCME, 2006). These are set out, in modified form, below:

- 1. There should be no appreciable risk to humans from a contaminated site. For each specified land use, there should be no restrictions as to the extent or nature of the interaction with the site. All activities normally associated with the intended land use should be free of any appreciable health risk.
- 2. Guidelines are based on defined, representative situations (exposure scenarios or land uses). Deriving numerical guidelines necessitates defining specific scenarios within which the exposure likely to arise on the site can be predicted with some degree of certainty.
- 3. Guidelines are derived by considering exposure through all relevant pathways.
- 4. A critical human receptor is identified for each land use. To ensure that the guidelines do not limit the application of a site within the intended land-use category, the defined exposure scenarios are usually based on the most sensitive receptor to the chemical, and the most critical health effect.
- 5. Guidelines should be reasonable, workable and usable. Guidelines are developed by applying scientifically derived information, backed by professional judgement where data gaps occur. Occasionally, defined exposure-based procedures produce numerical guidelines far below background levels of contamination occurring naturally in the soil. When this occurs, guidelines cannot be below background levels.

2.2 Generic equations

Soil guideline values for the protection of human health are typically based on generic assumptions about exposure incorporated into standard equations. These equations are essentially identical, regardless of jurisdiction, although the detail can vary considerably. The equations for a particular contaminant and exposure pathway i, follow the form:

Intake_i = soil concentration \times contact rate_i \times exposure time

Eqn 1

The intake is usually normalised to an intake rate per unit of body weight (BW, in kilogram) and unit of time (day) by dividing by body weight and an averaging time. In addition, the exposure time is typically represented as exposure frequency in days per year multiplied by exposure duration in years, resulting in:

Intake rate_i =
$$\frac{\text{soil concentration} \times \text{contact rate}_i \times \text{exposure frequency} \times \text{exposure duration}}{\text{body weight} \times \text{averaging time}}$$
 Eqn 2

The intake rate (in kg BW/day) is then compared with some acceptable intake rate for the substance (the reference health standard) and, for some substances, the particular pathway – with a human health risk indicated for exposure to that particular soil concentration and pathway if the intake rate exceeds the acceptable intake rate.

The acceptable intake is either the tolerable daily intake (TDI) for threshold compounds, or the dose that yields a specified increased cancer risk (the risk-specific dose). In New Zealand the specified acceptable cancer risk for non-threshold compounds is one additional cancer in 100,000 people (10^{-5}). This is discussed in greater detail in MfE (2011).

Contact rate may be for soil ingestion, inhalation (particulates and vapours), dermal absorption, produce uptake, contaminated water and a variety of other things, depending on the policy of the particular jurisdiction. Contact rate may be modified by absorption, or by matrix factors to account for different modes of absorption into the body from contaminated soil, or by enrichment factors that account for the enriched concentration of contaminants on fine particles that are more likely to be inhaled or stick to the skin. The averaging time is the length of time over which exposure is averaged to give an average daily rate.

An allowable soil concentration (a soil guideline value) can be back-calculated by setting the intake rate to the acceptable intake rate and rearranging the equation so that the soil concentration is on the left hand side:

Soil guideline value_i =
$$\frac{\text{acceptable intake}_i \times \text{body weight} \times \text{averaging time}}{\text{contact rate}_i \times \text{exposure frequency} \times \text{exposure duration}}$$
 Eqn 3

Similar values can be derived for each exposure pathway considered relevant. Some jurisdictions then choose the lowest value from the various pathway values, as the generic guideline for the substance in question. If there is more than one significant exposure pathway, the chosen soil guideline will result in an intake greater than the allowable intake, unless the allowable intake is factored down somehow to take the multiple pathways into account. As seen in appendix 5, Canada and some New Zealand guidelines adopt this approach.

A more usual approach is to back-calculate the guideline value combined over the relevant pathways. This is achieved by equating the sum of the hazard quotients (HQ) for each pathway with 1 (unity). The hazard quotient for a particular pathway is simply the ratio of the intake rate over the allowable intake.

intake rate ₁	intake rate ₂	$+$ intake rate ₃ $+$ $\dots =$ HO $+$ HO $+$ HO $+$ $\dots =$ 1	Ean A
acceptable intake ₁	acceptable intake ₂	acceptable intake ₃ + \cdots = $\Pi Q_1 + \Pi Q_2 + \Pi Q_3 + \cdots = 1$	Lqii 4

The soil guideline value is then calculated from equation 4 by substituting equation 2 into equation 4 for each exposure pathway, and then redefining the soil concentration (assumed to be the same for each pathway) as the desired generic guideline value. After rearranging to bring the soil guideline value to the left hand side, the equation becomes:



This is the basic algorithm used by the US EPA (1996a; 1996b) to calculated generic soil screening levels (SSLs) for multiple pathways and in the Ministry's 'Timber Treatment Guidelines' (MfE and MoH, 1997), and 'Sheep-dip Guide' (MfE, 2006a). The earlier equation 3 takes a slightly different form for each exposure pathway, being expanded with additional factors so the individual contact rates, which are generically expressed in mass of contaminant per day (mg/day), can be calculated from the exposure pathway-specific contact rates.

This additive approach is only valid if the acceptable intake has the same value for each pathway (typically defined by the oral pathway), or if different for one or more pathways (eg, the acceptable intake is different for the dermal pathway), so that all pathways have the same mode of toxic action. If the mode of action is different the pathway(s) with the different mode(s) of action must be considered separately, because the effects cannot be assumed to be additive.

Similarly, the additive approach is only valid if exposure to the soil in question can physically occur through multiple pathways. For example, it is not valid to combine exposure in a residential scenario to soil and produce ingestion, dermal absorption and inhalation of volatiles for deeper soil (eg, hydrocarbons on the water table at 2 metres) – because it is not physically possible for a person to be exposed to deeper soil through all these pathways simultaneously. In this instance, surface soil would need to be treated separately from deeper soil; and volatile inhalation is likely to be the only relevant pathway for deeper soil.

2.3 Generic exposure scenarios

As noted above, there is a considerable similarity to the basic approaches taken around the world to defining generic exposure scenarios, based around land use. Universally there is some form of residential exposure and generally some form of commercial or industrial exposure. Some jurisdictions consider an agricultural scenario – not for protection of produce, but for the protection of workers and/or farm residents. Some jurisdictions have also developed generic guidelines for various recreational scenarios.

Table 1 summarises exposure scenarios used in New Zealand's industry-based guidelines and several countries that New Zealand has traditionally looked to for guideline values, when a New Zealand guideline does not exist: specifically Australia, the United States, the United Kingdom, Canada and the Netherlands. All these countries have well-developed contaminated land assessment frameworks and guideline derivation methodologies.

The various scenarios presuppose a variety of exposure pathways and critical receptors. These are discussed in greater detail in subsequent sections.

Country	Scenario	Source documents
New Zealand 'Timber Treatment Guidelines'	Agricultural / horticultural ¹ (includes home-grown produce) Residential (high and typical home-grown produce intake) ² Industrial – paved, unpaved Subsurface maintenance workers	MfE and MoH (1997)
New Zealand 'Gasworks Guidelines'	Agricultural / horticultural ¹ (includes home-grown produce) Standard residential High-density residential (limited soil contact, no vegetable gardens) Commercial / industrial Parkland / recreational Maintenance workers	MfE (1997)
New Zealand 'Oil Industry Guidelines'	Agricultural / horticultural ¹ (includes home-grown produce) Residential Commercial / industrial Maintenance Protection of groundwater for potable use	MfE (1999)
New Zealand 'Sheep-dip Guide'	Lifestyle block (includes home-grown produce) Standard residential (includes home-grown produce) High-density urban residential Parks / recreation Commercial / industrial (unpaved)	MfE (2006a)
Australia	Standard residential (includes children's daycare centres, kindergartens, preschools, and primary schools) Residential with minimal soil contact (includes dwellings with fully and permanently paved yard space, e.g. high-rise apartments and flats) Parklands / recreational land use Commercial / industrial land use	NEPC (1999a)
US EPA Soil Screening Levels	Residential Industrial: indoor worker, outdoor worker Construction	US EPA (1996a and 2002a)
US EPA Regional preliminary remediation goals	Residential Industrial	US EPA (2008)
Canada	Agricultural Residential / parkland Commercial Industrial	CCME (1996, 2006)
UK	Allotments Residential with and without plant uptake Commercial / industrial	Defra and EA (2002a, 2002b)
The Netherlands	Residential with garden (standard scenario) Children's play areas Residential with kitchen garden Agriculture (equivalent to standard residential) Nature Green areas (parks and recreational areas) Other green areas, buildings and industry	VROM (2000), Brand et al (2007)

 Table 1:
 Scenarios for generic numeric values in various jurisdictions

1 Agricultural and agricultural / horticultural is used interchangeably in these documents.

2 The high value for home-grown produce is described as typical of 'rural residential'.

2.3.1 Exposure pathways

There is sufficient commonality around the world with respect to exposure pathways that some can be considered generic. Similarly, there is, in general, a common approach internationally to quantitative risk assessment. Soil ingestion is universally considered, as is one or both of dermal exposure and inhalation of particulates and/or vapours. Beyond these exposure pathways, there is considerable variation between countries as to what particular pathway should be included. For example, the United States uses few pathways for generic screening values with relatively conservative parameters, leaving further consideration to site-specific assessment; by contrast, the UK and the Netherlands consider multiple pathways using spreadsheet-based models. Further detail is described for these countries in appendix 5.

2.3.2 Contaminant characteristics

The characteristics of a particular contaminant determine, which exposure pathways may be significant, the form of the exposure equations, and (in some instances) the critical receptor. All jurisdictions considered as part of this study divide contaminants into:

- threshold and non-threshold in terms of their toxicological behaviour
- volatile or non-volatile with respect to the inhalation pathway.

New Zealand guideline documents differentiate between genotoxic and non-genotoxic carcinogens with the latter treated as threshold contaminants. Threshold and non-threshold contaminants are discussed in greater detail in MfE (2011).

Threshold contaminants are considered to manifest toxic effects if exposure exceeds a threshold concentration, and conventionally (including in New Zealand), are considered to include both non-genotoxic carcinogens and non-carcinogens. For these contaminants, a reference health standard (commonly as a tolerable daily intake) is typically established, being the estimated daily amount that can be taken into the body without any detrimental health effects occurring.

Non-threshold contaminants are conventionally considered to include genotoxic carcinogens, and are considered to have effects at all levels of exposure. The potency of non-threshold contaminants is typically expressed as a slope factor or maximum likelihood estimate (both represent the increased risk per daily dose), or as a risk-specific dose or index dose (analogous to a tolerable daily intake for a minimal and acceptable human health risk). The risk-specific or index dose is obtained by dividing the acceptable increased risk level (ie, 10⁻⁵ in New Zealand) by the slope factor (MfE, 2011).

The volatility of a contaminant affects whether the vapour pathway needs to be considered in addition to, or instead of, inhalation of contaminants attached to particulates (airborne dust). Inhalation of volatile organic compounds may be a significant part of overall exposure, whereas inhalation of particulates is typically not (and is ignored by some jurisdictions). Although of the same general form, the equations required to calculate exposure from particulates and volatiles have quite different subsidiary equations to derive the exposure rates. Subsidiary equations for particulates typically determine the amount of dust that is entrained into the air by the wind; equations for volatiles rely on the physico-chemical properties of the contaminant and soil, soil cover and building characteristics, to establish vapour concentrations that migrate to indoor or outdoor air spaces.

2.1.5 Critical receptor

Calculation of an $SGVs_{(health)}$ requires defining a critical receptor, with associated body weight, exposed skin areas, breathing rates and exposure rates for soil ingestion and dermal absorption, inhalation of particulates, and the like.

The critical receptor depends on land use / activity, and whether the contaminant is considered to be threshold or non-threshold. The critical receptor for the residential situation for threshold contaminants is almost universally a child (with associated low body weight and greater exposure rates to soil ingestion and dermal contact). The critical receptor in the industrial setting for a threshold substance is an adult worker.

For non-threshold substances, exposure over a lifetime is typically assumed: the critical receptor becomes a combination of childhood and adult exposure.

Further detail of this and other scenario-specific parameters is set out in section 5.

3 Exposure Scenarios

3.1 Scenarios adopted

Health risk assessment in contaminated-site practice is based on the assumption that individuals are exposed to contaminated soil while going about their normal activities. This exposure can occur in a number of ways: these are typically called exposure pathways and include soil ingestion, dermal absorption, and consumption of vegetables grown in contaminated soil. The exposure pathways used in the SCS_(health) derivation are explained in greater detail in section 4.1.

An exposure scenario is a combination of exposure pathways typical of a particular activity in which exposure to soil contaminants is likely to occur: the intent is to estimate the intake of a contaminant for that scenario. For simple risk assessment, a number of generic scenarios are used, with a standard combination of exposure pathways for each scenario. The generic scenarios used for the SCS derivations are intended to be typical of the great majority of situations in which ordinary New Zealanders may be exposed to soil contaminants. They include three residential scenarios – standard, rural and high-density – to cover the range of residential situations in which most people live; parks / recreation to cover active play or sporting activities; and a commercial / industrial scenario to cover outdoor workers at work.

This section describes the rationale for the five exposure scenarios that were chosen in the methodology and comments on some other potential scenarios that are not used.

3.1.1 Residential scenarios – rural residential / lifestyle block, standard residential, high-density residential

The protection of humans in a residential setting is a major driving factor behind the development of soil contaminant standards in New Zealand. All jurisdictions have SGVs for some form of residential setting. Ingestion of contaminated soil and consumption of home-grown produce are typically the most significant pathways for inorganic compounds, while inhalation of volatile organic compounds and dermal absorption of some organics can add further significant pathways. The residential scenarios are characterised by a child being the critical receptor (because of their lower body weight and relatively higher exposure to soil than adults) and inclusion of home-grown produce consumption, for all but the high-density residential scenario. Not all overseas jurisdictions provide for produce consumption in generic guidelines but as growing vegetables at home is quite common in New Zealand, this pathway can make a significant difference to the derived value for a particular contaminant, depending on whether it is taken up into plants.

Standard residential

On average, separate houses form 73 per cent of private dwellings in main urban areas (eg, cities), 79–82 per cent in other urban areas, and 83–88 per cent in rural areas in New Zealand in 2001 (Statistics New Zealand, undated). Particularly in outer urban areas and rural areas, separate houses are likely to contain gardens, and therefore have the potential to grow vegetables. Given the high proportion of separate houses and the potential to consume home-

grown vegetables from these dwellings, there needs to be a residential exposure scenario that considers consumption of home-grown produce.

This is supported by some local government research in Hamilton, Christchurch and Hastings. Work in Hamilton, reported in Cavanagh (2004a), indicates that about one-third of residential houses are likely to contain a garden in which home produce is grown. Similarly, in Christchurch, about 20 per cent of residential houses in recent subdivisions (less than 10 years old) contained a garden in which home produce is grown. A survey of residents' vegetable gardening habits and size of gardens carried out in Hastings and Havelock North found about two-thirds of the 121 households surveyed grew at least some of their vegetable consumption (Philip McKay, Hastings District Council, pers. comm).

High-density residential

The remainder of private dwellings in New Zealand are predominantly multiple-unit dwellings, multi-unit townhouses, blocks of flats and high-rise apartments. Almost 20 per cent of private dwellings in main urban areas were multi-unit dwellings in the 2001 Census; that proportion has probably risen since. Single-storey multi-unit dwellings are less likely to have gardens than separate houses and the gardens that do exist will tend to be small ornamental gardens, limiting the opportunity for soil contact. Significant growing of vegetables is not expected. The high-density residential scenario will therefore have lower soil ingestion rates than standard residential, and not include home-grown vegetable consumption.

A special case of high-density housing is inner city apartments, including high-rise developments. Residents would not be expected to have direct contact with soil, the only possible exposure being inhalation of volatiles for ground-floor residents where the floor is in direct contact with the ground. Ground-floor residences are akin to an industrial / commercial indoor worker scenario, except for the greater exposure frequency and being longer on site each day. Unless volatile contaminants are present there will be no soil ingestion, particulate inhalation or dermal contact – and therefore no risk. Risk from volatiles for ground-floor residents would be as calculated for the indoor worker, factored up to consider the greater exposure frequency and duration exposure to volatile compounds is not considered in this document.

Rural residential / lifestyle blocks

Increasingly, land that was previously in agricultural or horticultural use is being subdivided into lifestyle blocks. This land may be contaminated from the historical widespread use of persistent agrichemicals and/or disused sheep dips. There is also a greater potential for a higher proportion of home-grown vegetables (eg, compared to urban residents) to be grown on a lifestyle block, so the significance of consuming contaminated produce may be greater than in the residential scenarios considered above.

However, there is no information on how much produce might be grown for own use within lifestyle blocks or rural areas. Depending on the circumstances, 10 per cent home-grown produce may be appropriate (ie, as for standard residential), whereas 50 per cent is expected to be towards the high end of a more self-sufficient lifestyle that some rural dwellers may adopt. The 100 per cent produce value used in the 'Timber Treatment Guidelines' (MfE and MoH, 1997) is considered unrealistic for most people. In the absence of information for rural-dwellers, but in response to submissions that advocated for a higher proportion than the proposed 10 per cent in the discussion document (MfE 2010a), a produce proportion of 25 per cent has been adopted as the default. For illustrative purposes, soil guideline values have also been derived for

a higher proportion, 50 per cent, and for a lower proportion, 10 per cent. Ideally, the percentage of produce grown at home should be considered on a case-by-case basis for rural sites where a high proportion of produce is home-grown – particularly for those practising a 'self-sufficiency' lifestyle – and the SGV derived accordingly. The derived 50 per cent produce SGV may be suitable as a first approximation where the assessor deems it appropriate.

Poultry or grazing animals are often raised on a lifestyle block or farm, and consuming products from these animals (eggs, milk and meat) can constitute additional pathways of exposure to contaminants, particularly for lipophilic organic contaminants that bioaccumulate through the food chain. These pathways are highly variable in occurrence and highly influenced by site-specific considerations (contaminant, soil type, type of produce, degree of resident self-sufficiency), and should therefore be considered on a site-specific basis.

The rural residential scenario is not intended to cover the productive areas of agricultural land but is intended to be applicable to the immediate vicinity of the farmhouse or staff houses, an area of perhaps several hundred to a few thousand square metres. This will include kitchen garden areas, areas where children might routinely play and perhaps the 'home paddock', with the intention of protecting the farming family and any staff and their families. This is as agreed by the Technical Review Group for the Proposed National Environmental Standard for Assessing and Managing Contaminants in Soil, which suggested the scenario be called 'rural' so as to include residences on farms (MfE, 2005).

Locations for storing farm chemicals and fuel or for housing implements are generally similar to an unpaved commercial / industrial scenario with respect to human health effects, and should be considered accordingly.

3.1.2 Parks / recreation scenario

Some overseas jurisdictions including the Netherlands and Australia have park or recreational scenarios. Canada combines this scenario with residential use. The US EPA policy is to regard the use as site-specific. The scenario exists in two current New Zealand guidelines, the 'Gasworks Guidelines' (MfE, 1997) and the 'Sheep-dip Guide' (MfE, 2006a).

With New Zealanders' relatively outdoor lifestyle, use of parks and urban green spaces is common. Further, inadvertent or deliberate use of contaminated land for park area or reserve areas is not uncommon in New Zealand: for example, many playing fields in Wellington were former landfills, and the soil under Victoria Park in Auckland contains gasworks wastes. Subdivision of former agricultural or horticultural land for new urban development typically includes setting some land aside for reserve areas and public rights-of-way. Some district councils are permitting disposal of contaminated ex-orchard soil into such reserve areas during the development process, provided the soil complies with a recreational soil guideline value.

Recreational activities are diverse. They range from walking through a park, where very little soil exposure occurs, to playing contact sports such as rugby, where players can end up being caked with mud during wet conditions. An intermediate scenario might be a child in a playground where the grass is worn (resulting in soil contact). A generic guideline has difficulty covering such a wide range of potential exposures, with selection of exposure parameters for the extremes described above resulting in quite different soil guideline values. Conservatively, a residential non-produce scenario could be used for parks / recreation, but typically the exposure frequency will be less than that compared to residential scenarios, and is not recommended.

An analysis of alternative scenarios is presented in section 5.3.3. The analysis shows that, if soil ingestion is the main consideration (true for most substances but not necessarily all organic contaminants), then the parks / recreation scenario can reasonably cover suburban reserves within residential areas and a sports field both for children and adults, and also a secondary school playing field. However, a primary school playing field should be the subject of site-specific assessment, given the lower body weight of the children. The scenario is too conservative for parks and reserves used for passive recreation (eg, walking in a park or public garden), but could be used as a first screening. Children's playgrounds are so variable in their layout and use that the scenario might be conservative or non-conservative, and therefore site-specific assessment is recommended. The risk to park maintenance staff should be assessed separately using the commercial / industrial unpaved scenario (see subsection 3.1.3 below).

Park areas are often large and have multiples uses. It is necessary to firstly develop a good conceptual model of the site so that appropriate receptors and pathways are assigned to relevant sub-areas for screening assessment. It could be that a particular park is amenable to screening with SCSs over parts of the site, but site-specific assessment is necessary for other parts that potentially have much higher or lower soil contact than this scenario envisages. Table 2 gives a summary of parks / recreational subscenarios and the recommended approach.

Subscenario	Approach
Playing field	Included in scenario. Check occupational exposure for maintenance staff using commercial / industrial unpaved
Residential reserve where children play frequently	Included in scenario
Secondary school playing field	Included in scenario
Primary school playing field	Site-specific
Public green areas, reserves and gardens used for passive recreation	Scenario is very conservative but could be used for first screening, otherwise site-specific. Check exposure for park maintenance staff using commercial / industrial unpaved
Children's playground	Site-specific

 Table 2:
 Parks / recreation scenarios – recommended approaches

3.1.3 Commercial / industrial scenario

Contamination of industrial land is common. This land may remain as industrial land or may be converted to commercial uses such as shopping centres, warehousing and office parks; and sometimes for residential use. Some industrial land may have considerable areas of exposed soil, while much commercial land is almost or completely paved or covered in buildings. This results in quite a wide range of potential exposures. Workers in factories or commercial buildings will have little, if any, direct exposure to soil, but may have exposure to volatiles.

The US EPA, in developing its soil screening levels, differentiated between an indoor worker, who spent most of his or her time indoors with little soil exposure, and an outdoor worker involved in outside maintenance activities (effectively the site caretaker). This approach has some attraction. Existing New Zealand guidelines have differentiated between paved and unpaved, but have not discussed how the paved scenario might apply to indoor workers. The industrial paved scenario implies virtually no exposure, even to volatiles, except where soil was exposed during excavation. However, the infrequency of subsurface maintenance suggests maintenance should not be a controlling factor in a commercial or industrial scenario.

The revisions to the Dutch CSOIL 2000 model (reported in Brand et al, 2007) put industry, infrastructure (which includes roads) and buildings all in the same category, with the same contact frequency and soil ingestion rates as the recreation scenario – in effect acknowledging the common low exposure of these land uses. The UK rejects soil exposure on fully paved sites as implausible, reserving the commercial / industrial scenario to indoor workers in single-storey buildings such as factories and warehouses.

Canadians separate the commercial and industrial scenarios (CCME, 2006), to differentiate between where activity is primarily commercial (eg, a shopping mall) and industrial, which is specifically for production or manufacturing of goods. It would appear the Canadian industrial scenario is for an unpaved site, which would make it equivalent to the existing commercial / industrial unpaved scenario. New Zealand commercial / industrial paved could then be considered akin to the Canadian commercial scenario, with low exposure.

Cavanagh (2004a) suggested that the paved commercial land-use scenario (eg, shopping malls, retail shops), for which there is no exposure to surface soil, and an unpaved industrial land use in which buildings (eg, factories) are located within an otherwise unpaved site, provide useful generic descriptions of common industrial land uses. However, Cavanagh rejected the paved commercial scenario as not providing useful values for metals and semi-volatile contaminants, and proposed the unpaved industrial scenario as covering all commercial / industrial scenarios. However, this is overly conservative for the commonly encountered indoor commercial and industrial situations. The lack of direct soil exposure with which to derive values for metals and semi-volatiles (which will have no concentration limit) does not invalidate such a scenario. It is therefore proposed that two commercial / industrial scenarios be continued with, but that they are renamed and redefined as follows:

Commercial / industrial (indoor worker) represents factory workers on commercial or industrial sites with little exposed soil, where workers spend the majority of their time indoors carrying out relatively low-intensity tasks. Direct exposure to soil is limited or zero but exposure to volatiles migrating to indoor air is possible. There would be no concentration limit for most metals and semi-volatile organics, given the protection by concrete floors.

Nevertheless potentially contaminated sites that are being redeveloped for commercial or industrial use should be investigated. While there may be little or no risk in the long term, there will be some risk to workers during construction (see next paragraph); and potential risk to the wider environment from disposal of surplus soil generated during redevelopment. Disposal must be guided by sampling results.

Commercial / industrial (outdoor worker or unpaved) represents the outdoor worker who carries out maintenance activities involving soil exposure to surface or near-surface soil through gardening and other landscaping activities, and occasional shallow excavation for routine underground service maintenance activities. Exposure to soil is less intensive and/or less frequent than would occur during construction or extensive excavation works, but occurs over a longer period.

A separate case is exposure to volatiles outdoors on an unpaved site, where workers not engaged in activities likely to incur direct soil exposure, may be exposed to vapours in going about their general work. Such exposure could be for most days at work, which is more frequent than that envisaged for direct soil exposure for an outdoor worker. The current 'Oil Industry Guidelines' (MfE, 1999) suggest that this scenario is unlikely to be critical, as the guideline values are generally high and concentrations exceeding the guideline values over wide areas (the assumption behind the derivation of the values) are not likely to occur in practice for most situations. The few situations where exposure could be significant, eg, major petroleum or

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chemical installations, are candidates for site-specific assessment. For completeness, this scenario should be considered on a chemical-specific case-by-case basis when guidelines for volatiles are reviewed.

3.2 Scenarios not adopted

3.2.1 Maintenance / excavation scenario

Existing New Zealand guidelines have a scenario for maintenance / excavation workers, to deal with the greater exposure to soil contaminants than for ordinary outdoor workers. Cavanagh (2004a) recommended this scenario as applicable to subsurface maintenance works and construction activities, being applicable for all land uses, including the assessment of commercial (completely paved) land. This scenario was also noted as the only scenario relevant for soil contamination at depth.

The exposure parameters for the current New Zealand maintenance / excavation scenario are unrealistic. The typical commercial / industrial site simply does not get dug up on 50 occasions each year, every year for 20 years, involving the same personnel. Even if excavations were carried out on a number of occasions on a site, such excavation would typically be by contractors using different personnel. Therefore it can be assumed that exposure of an individual would be no more than a few occasions per year, suggesting the current guidelines are conservative by a factor of perhaps 10 for threshold substances – and much more than that for non-threshold substances, for which the duration affects the final value.

The Technical Review Group agreed after lengthy discussion that such a scenario should not be part of the NES (MfE, 2005). Sites would not be cleaned up to this standard. The Review Group considered it was more appropriate that exposure be limited through the site-specific controls that are required under health and safety legislation. This is similar to the UK, where maintenance / excavation activities are considered to be covered by occupational health protection legislation (Defra and EA, 2002b).

Under New Zealand legislation, when a site is known to be contaminated there is an onus on the employer to be aware of the potential hazard. The Health and Safety in Employment (HSE) Act (1992) is intended to protect the safety of individual workers and requires the employer (and individuals) to take steps to identify and eliminate, isolate or minimise hazards. Carried out properly, this will reduce the exposure of excavation workers to acceptable limits. In any case, most excavations will be of short enough duration that exposure will not be great, and if the ground is particularly contaminated (eg, gasworks waste or leaking underground storage tanks) it is often sufficiently obvious that workers and supervisors could be expected to notice and take precautions.

Particular individuals might, on occasion, be exposed to contaminated soil on more than one site as part of their work, but it is not reasonable to base the assessment of soil on all commercial / industrial sites on this relatively rare, and person-specific, situation. Most sites are not significantly contaminated and most workers would not move from one contaminated site to another. Again, the HSE Act is intended to protect the safety of the individual workers and provides the appropriate approach here.

There is a small subset of workers involved in specialist maintenance and soil removal tasks involving site contamination. Given the limited duration and frequency of most excavation, exposure to volatiles or liquid contaminants is generally the most likely risk-creating scenario. A particular case is redevelopment and re-tanking of service station sites, where exposure to volatile compounds is common. This is already dealt with by employers under the HSE Act by the writing of safety plans and compliance with confined-space regulations which require the measurement of vapours. Such measures are routinely carried out during tank pit excavations. Generic soil guideline values are of little use in such situations, given the wide variety of soil conditions and site circumstances.

In summary, it is proposed to dispense with the commercial / industrial excavation worker / maintenance scenario, and leave this situation to health and safety legislation.

3.2.2 Agricultural scenario

The 'Timber Treatment Guidelines' (MfE and MoH, 1997), the 'Gasworks Guidelines' (MfE, 1997) and 'Oil Industry Guidelines' (MfE, 1999) include an agricultural / horticultural land use scenario to protect the general public from concentrations of contaminants in produce that would pose a concern to public health. The scenario also protects the health of residents at any farm property from exposure via consumption of home grown livestock and produce, and through direct contact with contaminated soil. However, the more recent 'Sheep Dip Guidelines' (MfE, 2006a) dropped the agricultural scenario in favour of the lifestyle block scenario which removed consideration of protecting the productive capacity of land and exceeding the maximum residue levels in food.

The rural residential / lifestyle block scenario, proposed in this document, is intended to be applicable to the immediate vicinity of the farmhouse or staff houses with the intention of protecting the health of the farming family and any staff and their families.

The rural residential scenario is therefore not intended to cover the commercially productive areas of agricultural land. It is considered that the commercial uses of farm properties are either outside the scope of this proposal (ie, not directly related to effects on human health) or are dealt with by other legislation protecting public health:

- Plant growth and the health of soil micro organisms, while beneficial to maintaining productive capacity, are not directly related to human health effects.
- The safety of food produced for the general public is subject to the joint New Zealand Australian Food Standards. Testing under this jurisdiction is a more direct measure of determining whether this land is safe for human use.
- Farm worker's exposure is subject to the provisions of the Health Safety and Employment Act 1992. The requirements of an employer under this legislation are more fully discussed under the maintenance and excavation scenario described above.

In summary, it is proposed to dispense with the agricultural scenario, since the protection of farming families is addressed by the rural residential / lifestyle block scenario. The considerations relating to the productive parts of agricultural land being left to the food safety and health and safety legislation.

3.2.3 School or childcare facility scenarios

From time to time, schools and/or childcare facilities have been proposed as another generic scenario (MfE, 2005). Few international protocols specifically mention this land use, and those that do typically include it with a residential scenario (eg, NEPC, 1999a). However, typical exposure frequencies and durations are less than for a residential scenario; produce consumption does not generally apply (or probably will not apply at the rates assumed for the residential scenario – daily produce consumption – even if the school or kindergarten has a vegetable garden for demonstration or other purposes).

Schools are typically required to be open for between 190 and 200 days, considerably less than the 350-day exposure frequency for the residential scenario in existing New Zealand guidelines. Also, a pupil is usually indoors for most of the day and there is typically a smaller proportion of exposed soil in a school, compared with a residential site, suggesting less opportunity for exposure by direct contact. While duration is not relevant for threshold substances, the duration for non-threshold substances of 30 years for residential exposure is greater than would be expected for the school scenario. The body weight typically assumed for childhood exposure (typically 13–15 kg, depending on jurisdiction) is clearly too small for the average school child, being increasingly conservative as the child grows.

The greatest exposure is likely for a childcare centre, where attendance may be every day that the parent goes to work (say up to 250 days a year) from as early as being a toddler (ie, the 13–15 kg body weight applies) and could involve more outdoor play where soil exposure might occur, than a typical school situation. On this basis a child at a childcare centre is probably at least twice more exposed (on a weight-normalised basis) than a child just starting primary school (assumed 20 kg body weight) and at least five times more exposed than a child just entering high school (assumed 50 kg body weight), without fully accounting for differences in the proportion of exposed soil and opportunity for contact in the two sorts of facilities.

Cavanagh (2004a) suggested that an alternative to providing a school or childcare facility scenario as a generic scenario would be to provide explicit information (eg, appropriate exposure frequencies and duration) to enable the ready derivation of a site-specific value. The Technical Review Group (MfE, 2005) suggested that early childhood centres should be included under either the residential or high-density residential land uses, depending on how much paving the site had (while acknowledging that the residential land-use scenario includes produce consumption), but that calculations should be performed to determine the best fit.

Given the wide range of situations that a school or childcare scenario has to cover – early childhood, primary school, secondary school – with a wide range of body weights and potential exposure, a single generic scenario is not considered feasible. Instead, apart from the proposed use of the parks / recreation scenario for secondary school playing fields (section 3.2.2), a site-specific risk assessment is recommended as the appropriate approach for schools and childcare centres. However, providing there is no significant growing and consumption of vegetables (ie, a typical child is not receiving 10 per cent or more of its daily vegetable intake from site-grown vegetables), residential no-produce guidelines could be used as a conservative initial screening. Alternatively, the parks / recreational scenario could be used as an initial screening value for areas of secondary and primary schools other than playing fields, if justified on a case-by-case basis. Generally the exposure parameters for the parks / recreational scenario would be conservative for general areas of schools. In the small number of childcare centres where there is significant site-grown vegetable consumption, the standard residential guideline could be used as an initial screening value, provided research showed actual site-grown consumption fell within that assumed for the residential scenario.

Non-produce residential guideline values have been calculated (see detailed calculations in section 6) to facilitate site-specific assessment such as for schools, but it should be noted that non-produce residential scenarios are not part of the generic exposure scenarios and should not be used in other than site-specific assessment.

3.2.4 Groundwater

Groundwater is widely used in some areas of New Zealand for drinking-water, irrigation and stock water. Groundwater also discharges to the aquatic environment of surface water. Groundwater is included in the derivation of soil guideline values in some overseas jurisdictions, and soil guideline values protective of groundwater for human consumption are derived in the 'Oil Industry Guidelines' (MfE, 1999). These values are considered separately from values derived from soil ingestion, inhalation, produce ingestion and dermal exposure.

The development of soil guidelines for the protection of groundwater is currently beyond the scope of this document. It requires modelling the partitioning of the contaminant from soil to groundwater, and requires assumptions about soil type, area, depth and thickness of contamination, and hydrogeological properties of the underlying aquifer. Apart from the large variation in the way contaminated sites are contaminated, New Zealand is so varied geologically that selection of sensible generic parameters is a difficult task.

The consideration of groundwater contamination should be treated as a site-specific issue. If significant contamination is found on a site, and if groundwater is used there for consumption locally, then monitoring wells should be installed to measure groundwater concentrations directly.

4 **Derivation Equations**

4.1 General

All the jurisdictions reviewed for this report use essentially the same basic exposure equations, with variations in the:

- specific way they are applied
- subsidiary equations required to calculate indirect exposure from inhalation of particulates and volatiles, etc
- detail of the exposure scenarios and pathways that contribute to those scenarios.

Soil guideline values are based on assessing the intake of a particular contaminant and exposure pathway. Generic equations have been developed in section 2.1.1. In summary, for each pathway i the SGV_(health) is:

Soil guideline value_i = $\frac{\text{acceptable intake}_{i} \times \text{body weight} \times \text{averaging time}}{\text{contact rate}_{i} \times \text{exposure frequency} \times \text{exposure duration}}$

and the combined soil guideline value is obtained from:



Existing New Zealand guidelines use the following exposure pathways:

- soil ingestion
- produce consumption (residential scenarios only)
- dermal exposure
- inhalation of particulates
- inhalation of volatiles.

Only the first three pathways are considered in this document for the derivation of $SCS_{(health.)}$ Inhalation of particulates is a minor pathway and can be dispensed with unless the toxicity of the contaminant of concern via the inhalation route is much greater than via the oral route. The particulate inhalation pathway typically contributes considerably less than one per cent to the total exposure, and is therefore well within the uncertainty of calculation of the other pathways; hence it can be safely ignored. It should be checked where the inhalation toxicity is much higher than the oral toxicity, or where site-specific conditions suggest dust is an unusually large contributor.

While the inhalation pathway is important for volatiles, the derivation of SCSs for volatiles is beyond the scope of the current document. This issue is discussed further in section 4.7.2.

Some jurisdictions consider other exposure pathways in their generic derivations. However, in the New Zealand context it has been decided that for inorganic and semi-volatile contaminants, the three chosen pathways form the great majority of exposure for typical situations.

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The two receptor groups adopted are the same as those used in existing New Zealand guidelines (eg, MfE and MoH, 1997). This follows on from US EPA practice. The age-groups are 1-6 years and 7-30 years. The 1-6-year-old child will be the critical receptor for non-threshold substances for residential and recreational scenarios. An 'adult', 7-30 years, will be the critical receptor for worker scenarios.

Following existing practice, exposure is combined across age groups for non-threshold substances, in the form of age-adjusted contact rates.

4.2 Background exposure

It is common in some jurisdictions to subtract background exposure from the reference health standard and using the residual to calculate the soil guideline value. This is only applied to threshold substances, because intakes of non-threshold contaminants are considered on the basis of an increase in risk, which is irrespective of background exposure. Not subtracting background exposure for threshold substances would theoretically permit exposure in excess of the reference health standard (RHS) at soil concentrations equal to or slightly less than the SGV.

Some existing New Zealand SGV derivations follow the practice of subtracting the background whereas others do not. It has been determined (as reported in MfE, 2011) that an allowance for background exposure should be made for deriving New Zealand soil contaminant values for threshold substances. MfE (2011) provides details on how background exposure should be determined. This report also provides recommended values for background exposure for the priority contaminants considered in this document.

It is possible for background exposure to exceed the RHS, in which case an SGV cannot be derived. The adopted approach around this problem is a variation of that adopted in the UK. In the SGV derivation protocol for the UK (EA, 2008a), when background exposure comprises greater than 50 per cent of the RHS then the background exposure is taken to be 50 per cent of the RHS, ¹ Rather than using a fixed percentage when background is greater than 50 per cent of the RHS, the method adopted in this document is for the Toxicological Advisory Group of government toxicologists to consider the proportion allocated to exposure is negligible (defined as less than 5 per cent of the RHS) or no data on background exposure exists, then a maximum of 95 per cent of the RHS should be allocated to exposure from soil. This is to provide a slight degree of precaution for substances for which determining the background exposure may be problematic (MfE, 2011).

¹ Until the recent updating of the UK protocol, the requirement was 20 per cent of the tolerable daily intake (TDI) (Defra and EA, 2002b) when the background intake exceeded 80 per cent of the TDI. This was changed as it was considered to result in excessively low SGVs.

4.3 SGV_(health) derivation equations for each pathway

The main $SGV_{(health)}$ derivation equations for each exposure pathway are presented below. Exposure parameters values for each equation are provided in section 5. Common terms are listed below:

- SGV_i = soil guideline value for pathway i (mg/day)
- RHS = contaminant-specific reference health standard (mg/kg BW/day)
- BI = background intake (mg/kg BW/day)
- ED = exposure duration (years)
- EF = exposure frequency (days/year)
- AT = averaging time ED \times 365 days for a threshold substance
 - = lifetime (75 years) \times 365 = 27,375 days for non-threshold substance
- BW = body weight (kg).

4.4 Soil ingestion

Threshold substance

$$SGV_{ing} = \frac{(RHS - BI) \times BW \times AT \times 10^{6}}{IR \times EF \times ED} mg/kg$$
Eqn

6

where: IR = soil ingestion rate (mg/day)

As $AT = ED \times 365$, this reduces to:

$$SGV_{ing} = \frac{(RHS - BI) \times BW \times 365 \times 10^6}{IR \times EF} mg/kg$$
Eqn 7

Non-threshold substance

$$SGV_{ing} = \frac{RHS \times 27375 \times 10^{6}}{IR_{adi} \times EF} mg/kg$$
Eqn 8

with IR_{adj} being represented by:

$$IR_{adj} = \sum \frac{IR_i \times ED_i}{BW_i}$$
Eqn 9

IR _{adj}	=	the age-adjusted soil ingestion rate (mg year/kg day)
Σ		signifies summation over receptor groups $i = 1$ to n
IR _i	=	soil ingestion rate for receptor group i (mg/day)
$\mathbf{B}\mathbf{W}_{\mathrm{i}}$	=	body weight for receptor group i (kg).
	$\begin{array}{l} IR_{adj} \\ \sum \\ IR_i \\ BW_i \end{array}$	$\begin{array}{ll} IR_{adj} & = \\ \sum \\ IR_i & = \\ BW_i & = \end{array}$

4.5 Dermal absorption

Threshold substance

$$SGV_{d} = \frac{(RHS - BI) \times BW \times AT \times 10^{6}}{AR \times AH \times AF \times EF \times ED \times EV} mg/kg$$
where: AR = skin area of relevant exposed parts of the body (cm²)
$$AH = soil adherence factor (mg/cm2 - event)$$

$$AF = chemical specific dermal absorption factor$$

$$EV = events/day.$$

As $AT = ED \times 365$ and for the default of one event per day, this reduces to:

$$SGV_{d} = \frac{(RHS - BI) \times BW \times 365 \times 10^{6}}{AR \times AH \times AF \times EF} mg/kg$$
Eqn 11

Non-threshold substance

$$SGV_{d} = \frac{RHS \times 27,375 \times 10^{6}}{AD_{adj} \times AF \times EF} mg/kg$$
Eqn 12

$$AD_{adj}$$
 being represented by: $AD_{adj} = \sum_{i} \frac{AR_i \times AH_i \times ED_i}{BW_i}$ Eqn 13

 $\begin{array}{lll} \text{where:} & AD_{adj} = & \text{the age-adjusted dermal absorption factor (mg year/kg)} \\ & \sum & \text{signifies summation over receptor groups } i = 1 \text{ to n} \\ & AR_i = & \text{skin area of relevant exposed parts of the body for receptor group i (cm²)} \\ & AH_i = & \text{soil adherence factor for receptor group i (mg/cm² event)} \\ & BW_i = & \text{body weight for receptor group i (kg).} \end{array}$

The approach to dermal absorption is that of the US EPA (2001a; 2002a), which superseded US EPA's original approach used in the existing New Zealand guidelines. The approach is based on absorption per event, rather than being exposed for some proportion of the day, with the assumption that the soil adheres to the skin long enough for the contaminant to be absorbed into the body.² The soil adherence factor varies according to the type of exposure or activity, and varies according to the body part exposed. A single body-part-area-weighted adherence factor is used. The default is for there to be one dermal absorption event per day for residential or outdoor work situations. The equations are written for a single event.

² This assumption may not accord with the way the dermal absorption factor is derived. Some dermal absorption factors are derived on the basis of 24-hour exposure. Few people would not wash for 24 hours, therefore a 24-hour dermal exposure factor should be adjusted by a factor to take into account the period between washing. For the residential situation this is taken as 12 hours and for the occupational situation this is taken as 8 hours (from the *Sheep-dip Guide:* MfE, 2006a).

4.6 **Produce ingestion**

The equations for produce ingestion depend on the types of produce considered. As noted in section 5.4.2, only vegetable produce is considered and this has been divided into above-ground (leafy) and below-ground (roots and tubers), as contaminant uptake may be different for the different types of vegetable.

Threshold substance

Pg

$$SGV_{P} = \frac{(RHS - BI) \times BW \times AT}{IP \times Pg \times ED \times EF[(BCF_{root} + SL_{root}) \times p_{root} + (BCF_{tuber} + SL_{tuber}) \times p_{tuber} + (BCF_{leafy} + SL_{leafy}) \times p_{leafy}]}$$
Eqn 14

where: 1

IP = produce ingestion rate (kg dry weight/day)

- proportion of total daily produce consumption that is home-grown produce (dimensionless: no units)
- BCF = contaminant-specific bioconcentration factor (dry weight) (dimensionless)
- SL = produce-type-specific soil loading factor (dry weight) for soil attached to produce (no units)
- p = the proportion of total daily produce consumption for each produce type (dimensionless)

the subscripts leafy, root or tuber refer to above-ground edible vegetation and below ground roots (eg, carrots) and tubers (eg, potatoes), respectively.

As $AT = ED \times 365$, this reduces to:

$$SGV_{P} = \frac{(RHS - BI) \times BW \times 365}{IP \times Pg \times EF[(BCF_{root} + SL_{root}) \times p_{root} + (BCF_{tuber} + SL_{tuber}) \times p_{tuber} + (BCF_{leafy} + SL_{leafy}) \times p_{leafy}]}$$
 Eqn 15

Where BCFs are determined from empirical data (eg, for metals) the soil loading factor is zero as the measured BCFs will include attached soil. Where BCFs are derived theoretically, SL is taken to be 0.001 for roots and tubers (dry weight) and 0.0002 for leafy vegetables, following EA (2008a).

Where no distinction is made between the type of vegetable, equation 10 reduces to:

$$SGV_{P} = \frac{(RHS - BI) \times BW \times 365}{IP \times Pg \times EF \times (BCF + SL)} mg/kg$$
Eqn 16

and SL = 0 or 0.0008, as appropriate, the latter calculated as a weighted average, assuming dry weight daily consumption for a child of 7.6 grams of root and tuber vegetable and 2.9 grams of leafy vegetables, respectively (see section A3). The adult ratio of root to leaf consumption is sufficiently similar that the same average SL can be used.

Non-threshold substance

 $SGV_{p} = \frac{RHS \times 27375}{IP_{adj} \times Pg \times EF[(BCF_{root} + SL_{root}) \times p_{root} + (BCF_{tuber} + SL_{tuber}) \times p_{tuber} + (BCF_{leafy} + SL_{leafy}) \times p_{leafy}]} mg/kg Eqn 17$

or where there is no vegetable type distinction:

$$SGV_{P} = \frac{RHS \times 27375}{IP_{adj} \times Pg \times EF \times (BCF + SL)} mg/kg$$
Eqn 18
with IP_{adj} being represented by: IP_{adj} =
$$\sum \frac{IP_i \times ED_i}{BW_i}$$
 Eqn 19

Specific exceptions in the use of the produce pathway equations exist for some contaminants, notably copper and boron. In the case of copper, at soil concentrations that do not constitute a health human risk, applying a BCF would theoretically result in plant tissue concentrations that would kill the plant, meaning it could not be harvested. This being the case, the use of the produce pathway equations is not appropriate. Instead, a maximum tissue concentration is assumed and an additional notional background intake subtracted from the reference health standard on the assumption that the produce is consumed at that concentration. A similar approach has been adopted for boron because reliable bioaccumulation factors could not be derived.

The details are provided in sections 6.2 (boron) and 6.5 (copper).

4.7 Inhalation

In this document, the inhalation pathway has not been used in calculating the $SCSs_{(health)}$. The equations are given here for completeness.

4.7.1 Contaminated particulates

Contaminated airborne particulates may be generated from bare soil on a contaminated site and be inhaled by residents or workers. As discussed previously, this pathway is minor and will not normally be a component of a generic soil guideline value. Given the minor role the particulate inhalation pathway plays, no differentiation between the contaminant concentration of indoor dust and outdoor dust is proposed (unlike the CLEA model for example: EA, 2008a).

Threshold substance

$$SGV_{ih} = \frac{(RHS - BI) \times BW \times PEF \times AT}{IH \times EF \times R \times ED} mg/kg$$

Eqn 20

where: IH = inhalation rate (m^3/day) R = proportion retained in lung

R = proportion retained in lungs (dimensionless)

PEF = particle emission factor³ (m^3/kg).

³ The PEF is equivalent to the inverse of the proportion of the respirable (<10 micron) dust concentration coming from a contaminated source.

Non-threshold substance

$$SGV_{ih} = \frac{RHS \times PEF \times 27375}{IH_{adj} \times EF \times R} mg/kg$$
 Eqn 21

with IR_{adj} being represented by:

$$IH_{adj} = \sum \frac{IH_i \times ED_i}{BW_i}$$
Eqn 22

where:

 IH_{adj} = the age adjusted inhalation rate (m³ year/kg day)

 $\sum_{i=1}^{n}$ signifies summation over receptor groups i = 1 to n

 IH_i = inhalation rate for receptor group i (m³/day)

 BW_i = body weight for receptor group i (kg).

For some exposure scenarios (eg, occupational) a person will only be exposed for part of the day. This can be treated in two ways in the equations, either the inhalation rates are adjusted to reflect the shorter exposed period, or daily rates are used with the addition of a factor to reflect the proportion of the day spent exposed (eg, eight hours out of 24).

4.7.2 Volatiles inhalation

Exposure by inhalation of volatile organics presents a particular difficulty because the risk occurs indirectly from breathing vapours that have partitioned from contaminated soil at some depth below the surface. Figure 1 is a conceptual diagram of this process.

An equation similar to equation 11 can simply be written by replacing the concentration of contaminated particles being inhaled, with a vapour concentration. Two situations must be considered: volatilisation to outdoor air and volatilisation to indoor air.

Calculating soil guideline values for outdoor air inhalation is relatively straightforward. The vapour concentration to be put into the equation can be determined by applying a volatilisation factor to the soil concentrations. The volatilisation factor depends on generic soil properties, depth of contamination, contaminant-specific partition coefficients, diffusivity in air, and Henry's Law coefficients. Volatilisation factors can be calculated using the model by Jury et al (1983, 1984, 1990). Equations are presented in US EPA (2002a) with input factors for those equations. These factors, some of which are specific to US meteorological and geological conditions, can probably be applied to New Zealand conditions, if specific values are carefully selected.



Figure 1: Pathway for subsurface vapour intrusion into indoor air

Source: US EPA, 2004b.

Volatilisation to indoor air is a different proposition, however. There seems little point in calculating soil guideline values for outdoor inhalation without also addressing the more critical indoor air situation.

Volatilisation to indoor air requires modelling the migration of the vapour from the contaminant source, up through the soil and then through cracks or gaps in the building to the inside of the building. This has typically been carried out using a simple vapour migration model such as the Johnson and Ettinger (1991) model.

However, the vapour concentration is a non-linear function of the soil concentration, which means the equation must be solved for the soil concentration (the soil guideline value) in an iterative fashion. In other words, it is not just a matter of rearranging the equations to solve for the soil concentration, as it is with the other pathways. A further difficulty is that the vapour migration models have tended to be inaccurate, often by several orders of magnitude.

Davis et al (2004) provide a useful summary of some of the factors influencing vapour migration, each of which can contribute to inaccuracies if not model inadequacy:

- volume and location of the contamination relative to the surface and the groundwater table
- volatilisation and portioning from soil and groundwater
- diffusion
- sorption onto organic matter in the soil
- biodegradation
- soil properties such as soil moisture and permeability
- soil stratigraphy and layering
- temperature and barometric effects
- pressure effects due to wind
- density differences.

In addition, where migration into buildings is concerned, the design and configuration of the building has a large influence: for example whether it has a basement, is slab-on-grade, or is on piles with a crawl space. To date in New Zealand, only the slab-on-grade scenario has been considered. Yet houses with crawl spaces are just as common, if not more so. Attempting to model these two types of construction is beyond the scope of the present study and has proved difficult elsewhere.

As noted earlier, the UK Environment Agency (Evans et al, 2002) has reviewed a number of vapour migration models for use with both slab-on-grade and crawl space construction. They concluded that none of the models was sufficiently accurate to be recommended. They proposed further work on the issue.

Australia's **Commonwealth Scientific and Industrial Research Organisation** (CSIRO) has also reviewed vapour models with particular reference to Australian construction (Davis et al, 2004), which is not dissimilar to New Zealand's. CSIRO initially concluded that there were unacceptably large discrepancies between the models, with a lack of validation against field measurements. However, Davis et al (2008) have re-examined the Johnson and Ettinger (1991) model and recommended a modified form of it (not yet generally available) as the most suitable for Australian conditions when generating health screening levels for petroleum hydrocarbons, including for house with crawl space construction.

It is recommended that the Australian and UK work be examined as to its applicability for the New Zealand situation.

5 Exposure Parameters

5.1 General

The parameters used in the derivation of existing New Zealand soil guideline values are primarily based on US EPA data and some Australian data (eg, produce consumption). Some of these parameters have been updated since the guidelines were derived, for example, the US EPA has updated its approach to dermal exposure (US EPA, 2001a), further research has been carried out on soil ingestion (reviewed in US EPA, 2006a; Paustenbach et al, 2006; Van Holderbeke et al, 2007), and recent studies have become available on produce uptake (EA, 2006; Swartjes et al, 2007). In addition, information that is relevant to New Zealand is available and could be used instead of international data (eg, Russell et al, 1999). It is therefore appropriate to review the parameters currently in use.

In this report the parameters used to estimate exposure are divided into the following parameters:

- general exposure
- pathway-specific
- contaminant-specific (as a subset of pathway-specific).

General exposure parameters are dependent on the relevant receptors and the exposure scenario, but are independent of the pathway of exposure: they are common to all pathways. These include exposure frequencies, durations, averaging times and body weights.

Pathway-specific parameters are of two types. The first type is related to the type of receptor and activity (land use) in defining rates of exposure, eg, rates of soil ingestion, inhalation and produce consumption. The second type is dependent on the specific chemical and defines uptake rates, whether absorption rates through the skin or uptake into plants. These depend on the physico-chemical properties of the individual contaminants.

This section provides recommendations for the general exposure parameters, and then parameters used in individual exposure pathways.

5.2 Degree of conservatism

In selecting the appropriate values of individual parameters to derive guideline values, the degree of conservatism inherent in the individual parameters as well as in the derivation process, must be considered. This will ensure that the derived guideline values are not over- or under-conservative. Parameter values used internationally are influenced by different policy approaches.

As noted earlier, the US EPA (from which many of New Zealand's current parameter values are obtained) adopts a 'reasonable maximum exposure' approach. This is nominally aimed at providing a reasonable worst-case exposure scenario (that is, no more than 5 to 10 per cent of the population would be likely to exceed these exposures) and is based on a combination of average and upper-end exposures.

The US EPA has been criticised for being too conservative (US EPA, 2004a) because of combining too many upper-end estimates but has defended this approach as appropriate. Similar criticism has been levelled at the UK approach (Defra, 2006a), and the derivation process and associated input parameters were recently reviewed (Defra, 2008a; EA, 2008a).

Most countries use high-end estimates for exposure duration (this is only relevant for nonthreshold substances because exposure duration drops out for threshold substances). High-end estimates are also used for exposure frequency, but average estimates are used for body weight, skin areas and inhalation rates. This is the existing New Zealand practice, based on US EPA practice and it is proposed to continue this.

Given reference health values also tend to be conservative (with some exceptions): the combination of high-end durations (for non-threshold substances) and frequencies should ensure adequate conservatism. Soil guideline values will frequently be used as clean-up values when the cost of site-specific assessment cannot be justified. Over-conservative generic SGVs could result in unnecessary remediation (with consequent cost: turning useful resource into a waste and unnecessarily using up landfill space if the remediation involved 'dig and dump') or in unnecessary abandonment of projects, when the human health risk was actually acceptable. It is therefore proposed that the remaining parameters be central estimates to avoid overconservatism. Many of these parameters are contaminant-specific, and because of varying a central estimate will, for some particular site conditions, be non-conservative. Accordingly, use of central estimates for contaminant-specific parameters must be used with care.

5.3 General exposure parameters

The general exposure parameters include age ranges and body weights for those age ranges. These are dependent on the receptors considered important for a particular scenario. Averaging time, exposure frequency and exposure duration are scenario-dependent and in some cases also receptor-dependent.

General exposure parameters are required for children in all residential scenarios (lifestyle block, standard residential, high-density residential) and the parks and recreational scenario. General exposure parameters for adults are also required for all residential scenarios and the parks and recreation scenarios, and in addition for the commercial / industrial scenarios.

For children, the parameter values are primarily based on the age range over which soil and dust ingestion via inadvertent mouthing of non-food objects is important. The most recent Australian draft values (NEPC, 2010) are based on a zero to six-year-old child, Canadian values are based on a toddler (six months to four years), whereas the US EPA (and New Zealand) values are nominally based on a child aged one to six years. In reality, the 15 kg body weight currently adopted in the United States, suggests a two to three-year-old child.

As noted earlier, for threshold contaminants the exposure duration is not important (other than being long enough for the exposure to be considered chronic) as it occurs in both numerator and denominator of the derivation equations, thus cancels. The most important general parameters are body weight, which is directly related to age in the case of children, and exposure frequency. However, for non-threshold contaminants, the defined childhood and adult exposure periods are used to calculate age-adjusted parameters in determining lifetime-average doses. In these cases, childhood parameters are used for the defined age range for a child, and adult parameters are used for the remainder of exposure duration.

Combining high-end estimates for exposure frequency and duration has the risk of greater conservatism for non-threshold substances than for an equivalent threshold calculation. However, given that the exposure duration is less than the 75-year⁴ averaging time for a non-threshold substance, the potential over-conservatism is somewhat mitigated.

A summary of the age ranges and body weights of receptors used in existing national and international derivation protocols for deriving soil numeric values is given in Table 3.

Country	New Zealand	Australia ¹	US	Canada	Netherlands
Receptor age (years): • child • adult	1–6 7–30	0-6	1–6 7–30	6 months– 4 years	0–6 7–70
Body weight (kg): • child • adult	15 70	15.5 75	15 70	13 70	15 70

Table 3:Age ranges and body weights of receptors considered in national and
international derivation protocols

na = not applicable.

1 = revised NEPM draft 2010.

5.3.1 Body weight

The difference in the selected age range of concern for children is primarily reflected in the variable body weights used in different countries, which range from 13 to 15.5 kg in table 3. Fifteen kilograms is the approximate weight of a three-year-old child, or the average of a zero to six-year-old based on data provided in the *Exposure Factors Handbook* (US EPA, 1997), whereas a body weight of 13 kg is used for a two-year-old child. The differences in the applied body weight of a child would give rise to a variation of about 5–15 per cent in a derived value for a threshold contaminant based on soil ingestion, if all other parameter values were identical.

It is proposed to reduce the 15-kg body weight for a child used in existing New Zealand soil guidelines to 13 kg on the basis that this represents the median weight of the youngest child (a two-year-old) likely to be routinely exposed to dust indoors and soil outdoors. In other words, 15 kilograms is close to the 97th percentile for a two-year old and therefore does not represent a central tendency value for that age. Children younger than two will be less exposed to outdoor soil than a two-year-old, on average, as more parental or caregiver control of play activities is expected to be exercised.

For adults, a body weight of 70 kg is typically used by most countries. The exceptions are Australia, which uses 75 kg being the rounded average weight of adult males and females combined based on the latest *Australian exposure factor guidance* handbook (2011 *in press*) and the UK (which bases the adult receptor on a female and uses a distribution with a mean weight of 68.5 kg. Apart from the US, Australian and UK protocols, there seems to be little use of country-specific data for body weight. The rationale for selecting 70 kg as adult body weight appears to be that it provides a reasonable approximation of average adult body weight, as opposed to being based on more specific data. For example, the US *Exposure Factors Handbook* recommends that a body weight of 71.8 kg be used in the US. This is derived as a mean of the average male body weight of 78.1 kg and the average female body weight of 65.4 kg based on data collected over 1976–1980 (US EPA, 1997).

⁴ Increased from the current 70 years, see section 5.3.2.

The 1997 New Zealand National Nutrition Survey (Russell et al, 1999) established an average body weight for New Zealanders of 74.5 kg, based on the average of the average male and female body weights of 80.7 kg and 68.7 kg, respectively. These weights are for people aged 15 years and over, and would therefore over-estimate an average 'adult' weight from age seven. A body weight of 70 kg is used by the Ministry of Health for setting the *Drinking-water Standards* (MoH, 2008) and by the New Zealand Food Safety Authority for setting maximum residue limits in different foodstuffs.

The Toxicological Advisory Group decided to continue using an adult body weight of 70 kg for consistency with other New Zealand health guideline setting authorities. While this is conservative (results in lower soil guideline values) for the average adult body weight, it is justified on the basis that it is close to the average female adult weight.

5.3.2 Averaging time

The averaging time selected depends on the type of toxic effect being assessed. When evaluating exposures to developmental toxicants, intakes are calculated by averaging over the exposure event (eg, a day or a single exposure incident). For acute toxicants, intakes are calculated by averaging over the shortest exposure period that could produce an effect, usually an exposure event or a day. When evaluating longer-term exposure to threshold toxicants, intakes are typically calculated by averaging intakes over the period of exposure (ie, subchronic or chronic daily intakes). For non-threshold toxicants, intakes are calculated by pro-rating the total cumulative dose over a lifetime (ie, chronic daily intakes, also called lifetime average daily intake). This approach for carcinogens is based on the assumption that a high dose received over a short period of time is equivalent to a corresponding low dose spread over a lifetime (US EPA, 1989a).

To this point the New Zealand guidelines have adopted the US EPA approach. For threshold contaminants the averaging time (in days) is typically the exposure duration (in years) for the critical receptor multiplied by 365, the number of days in a year. For the residential setting the critical receptor is typically a child. Given that exposure duration cancels out in the exposure equations for threshold substances, the time dependence of the exposure reduces to the proportion of the year exposed (ratio of exposure frequency and number of days in a year). However, averaging time is important for non-threshold contaminants if the US EPA approach to non-threshold contaminants is adopted. The convention is almost universally to use an averaging time of a 70-year lifetime, expressed as days, resulting in an estimate of exposure as an annual average daily rate. An exception is the Dutch protocol, which uses an averaging time of 70 years for all contaminants and therefore results in higher soil guideline values. The *New Zealand Drinking-water Standards* (MoH, 2008) use 70 years as a lifetime for non-threshold substances.

It has been decided to continue using the averaging time conventions as adopted in all the existing New Zealand guidelines; however, it is appropriate to consider whether the 70-year lifetime should continue – or whether the averaging time should be increased to reflect the increased life-expectancy enjoyed by New Zealanders. Seventy years appears to have been based on now-outdated statistics.

Despite the generic United States guidance still using 70 years, the *Exposure Factors Handbook* (US EPA, 1997) recommends using more recent statistics of life expectancy. The US life expectancy across males and females was 70.8 years in 1970 whereas it had increased to 75.5 years at the time of the 1993 Census. US EPA (1997) recommends 75 years as an appropriate value to reflect the average life expectancy of the general population in the US.

Life expectancy in New Zealand has also increased. Figure 2 shows the male and female life expectancy at birth in New Zealand for the years 1876 to 2002. A life expectancy at birth of 70 years for the general population was achieved in about 1955, with 75 years being achieved in about 1988. The latest Census has shown a further increase, to an average of 81.9 and 77.9 years, respectively, for a girl and boy born in 2007 (Statistics New Zealand, 2008).

Continuing to use an averaging time of 70 years will result in a guideline value for a nonthreshold substance 12.5 per cent lower (more conservative) than if a life expectancy of 80 years was used, or 6.25 lower than if a life expectancy of 75 years was used.

Using an 80-year life expectancy is appropriate for people born now but not appropriate for the majority of the current population. Accordingly, based on these statistics it is recommended that the averaging time for non-threshold contaminants be increased to 75 years, to reflect the increased life expectancy of the current population.



Figure 2: Life expectancy at birth, 1876–2002

Source: Statistics New Zealand, 2006.

5.3.3 Exposure frequency

Exposure frequency is typically expressed as number of days per year, and relates to a given land-use scenario. The majority of countries use an exposure frequency of 365 days per year for residential (including parks / recreational exposure scenarios), and between 230 and 240 days per year for commercial / industrial scenarios. An exposure frequency of a little less than a year is used for residential scenarios in the existing New Zealand and US EPA protocols (350 days), based on US EPA data of an upper-end estimate of the time spent at home (95th percentile). The existing New Zealand protocols use 240 days per year for commercial / industrial scenarios. Note that the use of 365 days for residential was recently criticised in the UK on the basis that families routinely took holidays away from home every year; however, this was rejected on the basis of survey information (Defra, 2006a).

Residential

Limited data is available on time use by New Zealanders. The *New Zealand Time Use Survey* (Statistics New Zealand, 1999) provides the most extensive set of data on New Zealanders' activities. Based on the data in this report, New Zealanders, on average, spend about 70 per cent (males 66 per cent, females 75 per cent) of their time at home,⁵ which equates to about 260 days per year. However, this is not a very helpful average, as it says nothing about how many days are actually spent at home. Derivation of soil guideline values does not require the assumption of 24 hours at home for a soil ingestion or dermal absorption to occur. It just requires sufficient time for a dermal absorption or soil ingestion event to occur, which could be in the few hours after school, in the case of a child.

In the absence of better data, and given that a high-end estimate is desired, it is appropriate that the current residential exposure frequency of 350 days remains unchanged. This represents about two weeks out of the home, not necessarily as whole days but sufficiently long on the days when away from the home that soil exposure is unlikely to occur. This exposure frequency represents a 10 per cent 'discount' over the practice in Australia, many European countries and Canada, but is considered a high-end estimate.

Parks / recreation

No information on the frequency and duration of visiting parks and recreational facilities was found for New Zealand. A frequency of 350 days per year is used in the New Zealand 'Gasworks Guidelines' (MfE, 1997) and 'Sheep-dip Guide' (MfE, 2006a): this seems excessive, given the likelihood of bad weather preventing recreational activity for much greater than 15 days per year.

As noted in section 3.1.2, a wide range of activities can be fitted into the scenario, from simply walking in a park with little risk of soil exposure, through to frequent contact with potentially muddy ground while playing a sport such as rugby. The different possibilities will have different exposure frequencies and different exposure rates. Unfortunately it is not possible to determine an exposure frequency in isolation from the exposure-creating activity.

Possible scenarios include:

- An adult keen on both summer and winter sport who practises and plays on the same playing field perhaps two or three days a week in both summer and winter, for up to 50 weeks of the year (a maximum of 150 days per year). Soil ingestion rates are likely to be high-end estimates for both winter and summer sports, in winter being higher than summer.
- A small child plays junior sport on playing fields for limited summer and winter seasons, perhaps 26 weeks total. Winter soil ingestion rates would be greater than for residential but summer ingestion rates lower. The child then goes on to play more serious sport as an older child and teenager (say to age 20) on the same playing field for 40 weeks per year with games and practices two or three times per week. Soil ingestion rates are likely to be highend estimates.

⁵ Derived by Cavanagh (2004a) from data contained in Table 25 of the New Zealand Time Use Survey (Statistics New Zealand, 1999). Retrieved from http://www.stats.govt.nz

• A child up to early teenage years plays on a grass-covered suburban reserve near home several days a week, more frequent in summer than in winter, say a total of 200 days a year (roughly equivalent to six days a week in summer, two days a week in winter and four days a week for the rest of the year). There is anecdotal evidence that with decreasing residential lot sizes and increasing urban infill, suburban reserves are increasingly being used in place of residential backyards. Opportunity for soil contact will be limited by the grass cover, but a child would likely get dirtier after rain and/or more active play than during dry conditions and/or more passive play. As a matter of professional judgement, soil ingestion rates would be half to a quarter that of residential rates.

The question then arises as to what typical combination might be critical: frequent exposure at lower rates of more passive recreation; or less-frequent exposure at higher rates of more active recreation (eg, playing a common contact sport such as rugby in winter and sports such as cricket or softball, still involving some ground contact, in summer). For the purposes of calculating an initial comparative contaminant exposure, it can be assumed that most exposure will be through soil ingestion (although for some organic contaminants dermal absorption may also be important). Using the child residential soil ingestion rate of 50 mg/day and 100 mg/day as a high-end estimate for both children and adults (see section 5.4.1, table 7) and factoring these values up or down for the various scenarios and activities as a matter of professional judgement, weight-normalised daily soil ingestion rates (mg/kg BW/day) can be calculated.

The comparison is shown in table 4 for threshold and non-threshold substances. Age-adjusted exposure is necessary for non-threshold substances based on exposure durations. The proposed defaults for residential use have been assumed (see section 5.3.4), ie, six years exposure for a child and 14 years for an adult (ie, playing sport actively for 14 years on the same field). Soil intake for the equivalent standard residential scenario is shown for comparison.

Receptor	Child (13 kg) / teenager (70 kg)	Child (13 kg) / teenager (70 kg)	Adult (70 kg)	Child (13 kg) / adult (70 kg)	
Scenario	Junior rugby and cricket: 26 weeks x 1 day/week, then older child and teenager up to age 20 serious rugby and cricket for 40 weeks/ 3 days/week	Suburban reserve most days up to age 14 (older child only relevant for non- threshold contaminant)	Serious rugby and cricket player	Residential	
Days/year	26 / 120	200	150	35	50
Soil ingestion mg/day	100 rugby, 50 cricket	25	100 rugby, 50 cricket	50 25	
Rate mg/kg BW/day (threshold)	0.41	1.05	0.44	3.69	
Rate mg/kg BW/day (non-threshold)	0.10	0.11	0.08	0.	36

 Table 4:
 Comparison of daily soil intake for recreational scenarios

The suburban reserve has a weight-normalised daily soil intake significantly higher than the other recreational scenarios for threshold substances, and all recreational scenarios are significantly less than the residential scenario for both the threshold and non-threshold scenarios. For non-threshold substances, the three recreational scenarios are all similar. In other words, the chosen recreational scenarios, despite being quite different, are not so different that it is unreasonable to take the highest value and also apply this to the alternative scenarios. The suburban child scenario is the critical scenario for both threshold substances and non-threshold substances with respect to soil ingestion.

It is therefore recommended that the default parks / recreation scenario be that of a child up to age 14 years (see section 5.2.4) for 200 days per year with soil ingestion at a half the rate of standard residential. The exposure frequency is considered a high-end estimate and will also cover the scenario of high-end sports field use by a child or an adult. Appropriate dermal factors for sporting activity will also be required.

Commercial / industrial

Exposure frequency for commercial / industrial scenarios is typically based on a five-day working week, for 46 to 48 weeks (230–240 days per year). The exposure frequency for commercial / industrial scenarios in existing New Zealand industry-based guidelines is 240 days (48 weeks) per year. However, there are about two weeks of statutory holidays and, following the recent increase, an entitlement to four weeks annual leave. This equates to six weeks when most workers will not be at work, or 230 days when at work. It is therefore appropriate to reduce the current 240 days to 230 days. This is likely to represent a high-end estimate for those who work five days a week, as intended, but will be less conservative for those who frequently work a sixth day.

School scenario

While a school scenario is not proposed as a generic scenario, it is a scenario that concerns the community from time to time. For the purposes of site-specific assessments, an exposure frequency of 200 days per year is suggested. New Zealand primary and intermediate schools are required to be open for instruction for 394 half-days (197 days) and secondary and composite schools for 380 half-days (190 days). A frequency of 200 days per year should provide an upper-end estimate of exposure frequency for schools.

Early childhood education and childcare centres are more variable, ranging from a child attending perhaps two or three half days a week during the school term at a playcentre, through to the equivalent of a working year for a child in full-time care at a childcare centre while the parents work.

5.3.4 Exposure duration

Exposure duration is normally taken to be the length of time in years over which exposure occurs. However, the term is also used to describe the proportion of the day that any particular exposure event can occur.

Exposure duration – years

For threshold contaminants where children are the critical receptors (eg, residential and parks / recreation scenario), the duration of exposure is not important, as the averaging time in years is the same as the exposure duration. However, the critical age range is used to select other parameters appropriate to that age, which is taken as ages one to six.

Residential

The age ranges are important for the age-adjusted exposure calculations for non-threshold contaminants. As noted earlier, existing New Zealand and US protocols use an exposure duration of 30 years (6 years as a child and 24 years as an adult), which nominally represents a maximum time a resident will spend on one property, based on US data.

Census information on the length of time spent on a residential property is available for New Zealand. Statistics New Zealand has collected information on internal migration, from which some useful numbers can be derived. The 2006 Census provides information (available online at www.stats.govt.nz) on the number of continuous years at the same residence, broken down by age bracket. Within an age-bracket 'x–y' years, it is possible to determine the percentage of people who have inhabited the same residence for the full length of that age-bracket. Conservatively, this percentage is a high estimate of people in that age range who have inhabited the same residence all their lives. These statistics cannot help determine the number of people who have lived at a property as a child and perhaps teenager, leave home and then returned to live at the same house as an adult after a period away (possibly a relatively rare occurrence); nevertheless the data give some sense of what proportion of the population have lived continuously at one house both as a child and an adult.

For 20–24 years, 5.3 per cent of that age range had lived at a single residence for as long as their age in 2006. This translates to only 0.36 per cent of the total population; in other words, 99.6 per cent of the population had spent less than their first 20–24 years at a single residence. For 25–29 years, the closest age range for the currently assumed continuous exposure of 30 years, the same calculation produces 1.3 per cent having lived at the same residence almost all their lives. This group represents only 0.08 per cent of the total population. Only in the 15–19 age range does the percentage of the total population residing in the same house for their whole life rise to 1 per cent; still a small number.

Only 3.2 per cent of children aged four years have stayed all their lives in the same residence, so 96.8 per cent have not. Although data does not exist for up to six years of age (relevant to the childhood exposure scenario), the data on 0–4-year-olds suggests a similarly low percentage of children who have lived at the same house all their lives.

Considering the adult exposure scenario, the 2006 Census data shows that 6.2 per cent of the total population have stayed continuously in the same residence for 25 or more years; 9.7 per cent for 20 or more years; and 15.5 per cent for 15 or more years. Conversely, 84.5 per cent of the total population has lived in the same house fewer than 15 years, and 57.7 per cent fewer than 4 years.

It is not possible to determine from the 2006 Census the percentage of the population having resided in the same house both as a child and an adult. Clearly, continuous residence from birth to late teenage years is a small percentage, with residence as a child for up to six years being estimated at only 3 per cent of the population. In addition, the available information suggests only a very small percentage of adults are likely to reside in their childhood home for that length of time: most likely a small subset of rural dwellers who have been raised on a farm, and remain or return there as an adult. The current assumption of residing in the same house for 30 years, including as a child, therefore seems excessively conservative.

Given that New Zealand's population is mobile as demonstrated, it is appropriate for the standard residential scenario that the length of residence as an adult and child is reduced to 20 years: 6 years as a child and 14 years as an adult. This will still be a high-end estimate. The same duration is proposed for the high-density residential and parks / recreation scenarios.

A study by Sanson et al (2004) of several thousand lifestyle blocks in New Zealand, using census and land ownership records, found that 35.5 per cent of lifestyle blocks with a dwelling remained in the same ownership for more than 30 years. For the rural residential scenario, it is therefore proposed to retain the current 30-year duration (6 years as a child and 24 years as an adult).

It is worth examining whether some site-specific situations exist with extended adult residence being the critical case. This would only be for non-threshold contaminants where exposure durations affect the SGV_(health) calculation. The 2006 Census found only 4.5 per cent of the general population stayed in the same house for more than 30 years. This suggests a duration of 30 years as an adult is a reasonable high-end estimate for the standard residential scenario. The Sanson et al (2004) study suggests that perhaps 40 years as an adult is a high-end estimate for the rural / lifestyle block scenario.

Check calculations using these extended adult durations show that the child / adult combination still remains critical, although the derived SGVs for extended adult durations are only marginally higher than the SGVs for the child / adult combination for some contaminants. If, in a site-specific study, the various default exposure parameters in this document were changed, then extended adult-only durations could become critical for some non-threshold substances; this would be particularly so if the ratio of adult to child rates of any of soil ingestion, produce ingestion, or dermal soil adherence was increased. A further check shows that increasing the adult-only duration to 35 years results in the derived SGVs for the standard residential scenario becoming the critical (lowest) value for some substances. This indicates that scenarios involving extended durations need to be checked in site-specific analyses.

Parks / recreation

Continuous recreational exposure to the same location is quite likely to extend beyond the childhood years of one to six. This is not important for threshold substances but is important for non-threshold substances. There is no information available for the likely exposure a person would have to the same sports field or suburban reserve as a child and a teenager or older. It is conceivable that a person residing in the same town from childhood to adulthood has played active sport as a child, teenager and young adult on the same playing field. However, as noted in section 5.3.3, the critical recreational scenario appears to be more likely that of a suburban reserve. It seems unlikely that a person would be exposed to soil in a suburban reserve from childhood through to adulthood, but it is conceivable that a child would play in a suburban reserve up to their teenage years. There will be an age, however, when a teenager is more likely to have interests other than active play in a suburban reserve, and the frequency of play will rapidly diminish. As a matter of professional judgement this age has been set at 14. This age was used in section 5.3.3 to determine which of several possible recreational scenarios was critical. The derivation of the SCS is not very sensitive to the cut-off age for an older child, with a year either way making only a marginal difference (a few per cent) to the SCS.

Commercial / industrial

For commercial / industrial scenarios, the exposure duration is based on the length of time in one job, and ranges from 20 years in New Zealand for current guidelines to 43 years in the UK. The 20-year exposure duration used in New Zealand is based on US EPA data for the 90th percentile for time spent in one job (MfE and MoH, 1997). US EPA protocols for deriving generic soil guideline values use an exposure duration of 25 years for commercial / industrial scenarios based on the 95th percentile for time spent in one job.

There is limited data available on job tenure in New Zealand. The question has not been included in the five-yearly census. Statistics New Zealand's Linked Employer-Employee Data (LEED) matches tax data to statistics collected from employers; this provides limited job tenure information since the 1999 tax year. The most recent compilation, for the 2006 tax year (available online at www.stats.govt.nz), allows the percentage of employees who have worked seven years or greater with the same employer to be calculated. The data is limited in that tenure is not necessarily continuous and work is not necessarily full-time: the data only indicates whether work was performed in a particular month for an employer, not how much work. In addition, while the statistics are broken down to a regional level, they do not necessarily indicate work at a particular site. However, 12 per cent of workers have worked for the same employer for seven years or greater. Given the limitations, the percentage of workers working continuously at the same site is likely to be lower. This suggests that a 20-year job tenure is greater than the 90th percentile. In the absence of better information, it is proposed that the exposure duration for industrial / commercial land uses remain at 20 years, as a high-end estimate.

Exposure duration – hours per day

Exposure duration expressed as hours per day may also be used for some exposure scenarios or exposure pathways. Soil and dust ingestion is normally taken as a daily rate, based on a combination of indoor and outdoor exposure with no specified contact period during the day. While the assumption is that ingestion is spread out over the day, in reality it would consist of a number of individual hand-to-mouth contact events of only a few minutes duration. In addition, the available studies for estimating ingestion rates have been carried out over a number of days or weeks, and the rates are daily averages (see section 5.4.1); these render the duration of the activity which resulted in the soil ingestion irrelevant.

For dermal exposure, the UK protocol originally used a 12-hour exposure duration in a residential setting (Defra and EA, 2002a) but later changed that (EA, 2005a, 2008a) to the US EPA approach: this uses a 24-hour exposure duration, on the basis that soil is attached to the skin during an event and is then not washed off before it is absorbed (US EPA, 2001a). The default assumption is one exposure event per day. The Dutch CSOIL model (Brand et al, 2007) approach is to assign a period in hours for each of indoor and outdoor for both children and adults, because a dermal adsorption rate per hour is used in the model. Different durations are specified for different exposure scenarios. The approach of US EPA (2001a) and a later update, US EPA (2004c), are used in the current study. This may not be appropriate for all substances, depending on how the particular absorption rate is derived, and may need to be varied on a contaminant-specific basis.

For inhalation exposure, the durations of exposure are inextricably linked with inhalation rates, with an inhalation rate of m^3/day specified for the particular scenario and receptor, taking into account the duration. This is the approach adopted here, and is discussed in the next section.

In summary, the proportion of the day exposed is not important for soil ingestion and dermal exposure, if dermal absorption coefficients are based on event-based experiments. However, if the dermal absorption coefficient for a particular contaminant is based on 24-hour exposure, then the proportion of the day exposed has to be factored into the calculation. Following the existing 'Sheep-dip Guide' (MfE, 2006a), for the residential situation this is taken as 12 hours and for the occupational situation 8 hours. For the inhalation pathway, residential exposure is assumed to be 24 hours and occupational exposure 8 hours.

5.4 Pathway-specific parameters

The pathway-specific parameters are either exposure rates or combination of parameters that make up an exposure rate. These are the time-dependent soil ingestion, produce consumption and inhalation rates, and soil adherence values and skin surface areas necessary to calculate dermal soil loadings. For each pathway there are several scenario / receptor combinations, each of which must have a set of parameters – meaning a large number of parameters need to be decided on.

Soil ingestion and produce consumption are typically the pathways contributing most to residential guideline values, although dermal absorption can be significant for some contaminants. Soil ingestion and sometimes dermal absorption are the important pathways for high-density residential and the non-residential scenarios.

5.4.1 Soil ingestion

Soil ingestion can occur in both indoor and outdoor settings as a result of deliberate sucking and mouthing of objects by children, inadvertent hand-to-mouth transfer by children and adults, and ingestion of soil attached to produce. Early estimates of the amount of soil ingested were largely activity-based, whereby soil ingestion rates were estimated from factors such as time spent outside or doing certain activities, the number of hand-to-mouth events, and the degree of hand-soiling – whether measured or predicted. More recent studies have predominantly used tracer elements found in soil (typically aluminium, silicon and titanium, but also barium, manganese, vanadium, yttrium and zirconium) to provide a more direct measurement of soil ingestion. In these studies, soil ingested daily and the levels in urine and faeces. The better-designed studies took into account other sources of these tracers in food, medicines and consumer products.

Stanek et al (2001) have outlined the improvement in soil ingestion estimates from tracer studies as estimation techniques improved. Initial estimates of soil ingestion were based on individual trace elements (Binder et al, 1986; Calabrese et al, 1989; Davis et al, 1990). Since the estimated distributions from different trace elements often differed substantially, subsequent work focused on ways of identifying more reliable estimates (Stanek and Calabrese, 1991a; Calabrese and Stanek, 1991). This work led to the use of trace element food / soil ratios as a means of identifying potentially reliable trace elements, and the use of the median trace element estimate from among a subset of reliable elements (Stanek and Calabrese 1991b). The results of this work contributed to improvements in study designs for soil ingestion estimation via the inclusion of additional trace elements, longer study designs, and use of special diets.

While the relative understanding of soil ingestion has improved, average rates of ingestion in relation to specific activities and land-use remain uncertain. Limited data exists for child ingestion rates in a residential setting from a few reliable studies. Very little data exists for adult ingestion rates for a residential setting. Estimating ingestion rates for non-residential scenarios relies on professional judgement as no reliable tracer studies exist.

A large range of soil ingestion rates is used by individual agencies in deriving guideline values (table 5), depending on the particular agency's philosophy (worst-case / high-end estimate versus best estimate) and which studies have been used to derive the estimates.

Receptor	New Zealand	Australia	US	Canada	Netherlands	UK
Child	100	100ª	200	80	100 ^c	100 ^d
Adult – resident	25	25 ^b 50 ^a 12.5 ^b	100	20	50	na
Adult – worker	25	25	50 indoor 100 outdoor	20	na	50

 Table 5:
 Soil ingestion rates used in national and international protocols

a Low density residential according to revised NEPM draft (2010).

b High density residential according to revised NEPM draft (2010).

c Reduced from 150 mg/day in 2001 (Brand et al, 2007).

d Originally a probability distribution with mean 100 mg/day and 95th percentile of 300 mg/day (Defra and EA, 2002a), subsequently modified to a single-point estimate of 100 mg/day (EA, 2008a).

na = Not applicable.

Residential soil ingestion - children

Children have been the primary focus for soil ingestion studies due to their inclination to mouth objects (hands, toys) and ingest dirt. Combined with low body weight, childhood soil ingestion is typically a major component of residential soil guideline values. Generally, it is considered that intensive mouthing diminishes after two to three years of age and negligible soil ingestion occurs after the age of six to seven (Paustenbach, 2000), making soil ingestion at later ages relatively less important.

The value of 100 mg/day for soil ingestion by children used in current New Zealand protocols (eg, MfE and MoH, 1997) comes from ANZECC (1992) and is the same as that used in the derivation of the Australian health investigation levels as proposed by Langley and El Saadi (1991), based on a review by Taylor (1991). The value was considered to be a conservative value at the time but drew on work that pre-dated more recent tracer studies. The 100 mg/ day is still used in Australia for children in the low density residential scenario, following a re-check and endorsement of the USEPA factors in 2010.

The *Exposure Factors Handbook* (US EPA, 1997) states a soil ingestion rate of 100 mg/day represents an average estimate for exposure assessments, but recommends as a high-end estimate a rate of 200 mg/day in risk assessment. Generic soil guidance in the US (eg, US EPA 1996a, 2002a) is based on 200 mg/day.

A report commissioned by the US EPA (Versar, 2001) considered the then available studies on childhood soil ingestion. The report noted that a number of studies had weaknesses and concluded that the best estimates were from several mass-balance studies using metal tracers conducted in the United States. Two Dutch studies (Clausing et al, 1987; Van Wijnen et al, 1990) were rejected because of study-design limitations. Versar (2001) also noted the possibility of different cultural practices between Dutch and US child rearing that might affect behaviour and soil ingestion.

The draft *Child Specific Exposure Factors Handbook* (US EPA, 2006a) reviews the science to that point and arrives at a mean childhood soil ingestion rate of 90 mg/day and median soil ingestion rate of 35 mg/day, with a 95th percentile value of 236 mg/day. Because of the skewed nature of the distribution, the mean is larger than the 75th percentile value. US EPA (2006a) arrives at these estimates by applying (unstated) weighting factors to the results of five 'key' mass-balance tracer studies to calculate a weighted average distribution: four original studies and one study re-analysing data from two of the other studies – in effect double counting some of the data in the weighted average. US EPA (2006a) acknowledges this double counting but does not explain why, nor discuss what effect this might have on the overall estimate. It is also not clear why some studies were considered 'key' whereas later studies re-analysing the same original data using improved statistical techniques were not considered 'key'. The later analyses generally resulted in lower soil ingestion estimates.

The four original tracer studies considered robust enough to be included in the weighted average of the US EPA (2006a) review were all conducted in the United States. These were:

- a study of 104 children from an semi-arid, three-city area in southeast Washington State (Davis et al, 1990)
- a further study a year later of a 19-child subset of the Washington study (Davis and Mirick, 2006)
- a study of 65 children from the Amherst area of Massachusetts (Stanek and Calabrese, 1995a, using data from Calabrese et al, 1989)
- a study of 64 children in Anaconda, Montana (Calabrese et al, 1997).

The fifth study included in the weighted average, Stanek and Calabrese (1995b), sought to provide better estimates by reanalysing Davis et al (1990) and Calabrese et al (1989) using the 'best tracer method' to correct for errors in the tracer input and output measurements, and error from ingestion of tracers from non-food and non-soil sources.

The review rejected some other original tracer studies because of study design limitations, particularly not accounting for tracers sources other than soil, for example in food and in medicines. Rejected studies included those of Binder et al (1986), Clausing et al (1987), and Van Wijnen et al (1990).

US EPA (2006a) goes on to recommend that the best estimate for mean child soil ingestion for ages one to seven years is 100 mg/day (rounding up of 90 mg/day) and further recommends a 95th percentile value for soil ingestion of 400 mg/day. It is notable that the latter recommendation is from soil and dust data, rather than soil alone, whereas the 95th percentile estimate for soil alone rounded to the nearest 100 would be 200 mg/kg. The recommendations in US EPA (2006a) essentially confirm earlier recommendations in the 2002 interim version of the same document (US EPA, 2002b) and the *Exposure Factors Handbook* (US EPA, 1997).

The double-counting of two of the earlier studies will have tended to bias the weighted averages calculated in US EPA (2006a). It is not possible to determine the precise bias because the weighting factors are not given (other than the incorporated sample size and 'other statistical factors'). However, if a simple average is used for the mean and median soil ingestions across the five studies, the values are 80 and 33 mg/day, compared with the weighted averages of 90 and 35 mg/day. If Davis et al (1990) and Stanek and Calabrese (1995a) are then not included in the average (as reanalysed in Stanek and Calabrese (1995b)), the simple averages of mean and median soil ingestion for the smaller number of key studies are 69 and 29 mg/day, respectively. This suggests the double-counting has biased the US EPA (2006a) result upwards by perhaps 10–20 mg/day for the mean and perhaps 5 mg/day for the median. The US EPA's

rounded-up recommendation of 100 mg/kg for the mean soil ingestion therefore appears high by 20-30 mg/kg.⁶

It is notable that US EPA (2006a) has apparently not taken into account the view of one of the leading investigators in this field, Edward Calabrese: he was the principal author of the 1989 Amherst study and a collaborator with Edward Stanek in the later Anaconda study plus the many re-analyses of both studies. In a letter to the General Electric Company, Calabrese (2003) expressed the view that the 1989 Amherst study overestimated child soil ingestion. Calabrese instead favoured the reanalysed results of the Anaconda study (Stanek and Calabrese, 2000), in recommending a central tendency rate of 20 mg/day (based on a median) and an upperbound (95th percentile) rate of 100 mg/day.

Paustenbach et al (2006) reviewed the literature as part of determining input values for a probabilistic risk analysis for dioxins. They concluded that Stanek et al (2001) provided the most robust data set for determining a probability distribution for childhood soil ingestion. Stanek et al (2001) reanalysed earlier data from the Anaconda study (Calabrese et al, 1997), extrapolating the short-term measurements to long-term estimates, and concluded that earlier estimates (eg, the recommendations in US EPA, 1997) were too high. Their method was aimed at eliminating bias resulting from uncertainty in the daily estimate, or variability in soil ingestion from day to day – that bias overestimating soil ingestion for upper percentiles and underestimating it for lower percentiles. Paustenbach et al (2006) suggested the analysis of Stanek et al (2001) was a vast improvement over the original analysis of the studies considered to have an adequate design (Davis et al, 1990, Calabrese et al, 1989, Calabrese et al, 1997), with the improved method resulting in lower values.

Van Holderbeke et al (2007) provided a good summary of the various issues around attempting to estimate soil ingestion. They reviewed the literature to that time, including mass-balance tracer studies, behavioural hand-to-mouth and hand-soil loading-based studies, and various studies based on biomonitoring. Their aim was to arrive at soil ingestion estimates relevant to a project in the Kempen region of Belgium and the Netherlands. With respect to the tracer studies, they went over similar ground as US EPA (2006a) in determining ranges and arithmetic average across the various studies for the soil ingestion rate mean and percentiles. Like US EPA (2006a) they included in their calculations data from Davis et al (1990), Davis and Mirick (2006) and Calabrese et al (1997); but unlike US EPA (2006a) rejected the Amherst study (Calabrese et al, 1989) and subsequent re-analyses of that data. Data from the Amherst study was left out on the basis of the principal author subsequently stating that the results overestimated soil ingestion rates (Calabrese, 2003). Unlike US EPA (2006a) they also included results from Stanek and Calabrese (2000), a re-analysis of the Anaconda study – in effect double-counting that study – and also included data from Clausing et al (1987) and Van Wijnen et al (1990). The latter two studies were apparently included on the basis that they are European studies and therefore more relevant to Belgium, despite their design limitations.

⁵ Since this section was written, the US EPA has released the final version of the *Child Specfic Exposure Factors Handbook* (EPA/600/R-06/096F, National Center for Environmental Assessment, Office of Research and Development, Washington, September 2008). Although this document arrives at a similar conclusion with respect to soil ingestion rates as the 2006 draft, and presents a similar range of scientific studies from the international literature, little detail is given to justify the recommended ingestion rates.

Van Holderbeke et al (2007) concluded that the best central tendency estimates of child ingestion rates from tracer studies were, for the median value, 27 mg/day (average of medians ranging from 17–42) and for the mean value, 63 mg/day (average of means ranging from 31 to 120 mg/day).

As noted earlier, Van Holderbeke et al (2007) also reviewed the literature for studies other than mass-balance tracer studies. Such other studies are generally considered less reliable. Without going into the details here, these authors determined that average child ingestion rates from hand loading studies range from 7 to 60 mg/day, from biomonitoring studies 50–100 mg/day, and from empirical relationships 20–70 mg/day. These ranges are generally consistent with the mass-tracer studies.

A summary of the child ingestion values proposed in the various references are set out in table 6. They all draw on essentially the same information, with different interpretations as to which original studies or reanalyses of these studies should be considered.

Apart from Van Holderbeke et al (2007), the various reviews place little reliance on the European tracer studies, favouring the US studies because of their better study design. Van Holderbeke et al appear to have included the other studies only because they were European and hence might provide better values for European children, regardless of study design. This raises the issue of how relevant any of the studies might be for New Zealand conditions and children.

Reference	Age range (years)	Mean (mg/day)	Median (mg/day)	75th percentile (mg/day)	Upper bound (mg/day)	Comment
Versar (2001)	1–2	30	24	_	100 (90th percentile)	
	3–5	30	20	_	150 (90th percentile)	
	6–10	71	37	-	187 (90th percentile)	
Calabrese (2003 ⁷)	1–4		20	-	100 (95th percentile)	From Stanek and Calabrese (2000)
US EPA (2006a)	1–7	100ª	35	78	400 ^b (95th percentile)	Weighted averages from five 'key' mass balance studies
Paustenbach et al (2006)	1–4	31	24	42	91 (95th percentile)	From Stanek et al (2001)
Van Holderbeke	1–7	60	30	_	195	Tracer studies
et al (2007)		7–60	-	-	-	Behavioural studies
		50–100	-	-	-	Modelling / biomarkers
		20–70	_	-	_	Empirical relationships

 Table 6:
 Summary of child soil ingestion rate recommendations as reviewed

a Rounded up from the weighted mean value of 90 mg/kg.

b Includes soil and dust. Value is rounded from average of 95th percentiles of the reviewed studies of 449 mg/kg. Average of 95th percentiles for soil alone was 236 mg/kg.

⁷ Personal letter, not peer reviewed.

Many physical and societal factors affect the opportunity for soil and dust exposure and a child's behaviour in potentially being exposed. The factors tend to be interrelated, and include:

- climate opportunity for outdoor play, likelihood of soil sticking to shoes and being tracked inside, generation of dust
- style of residential development house construction and dust-tightness, types of indoor floor coverings (and whether they gather dust), presence of gardens or other bare soil
- lifestyle time spent outdoors, type of outdoor play, popularity of gardening activities, societal attitudes to how clean a house should be
- parental attitudes attitude to allowing outside play and whether children are allowed to get dirty, insistence on hand washing.

Given these factors, and without information to quantify any differences, it cannot be determined how well the United States values translate to another location. At best it is a matter of judgement: in a general sense the New Zealand style of housing and lifestyle is possibly closer to those in the United States than Europe, suggesting the opportunity for outside play (and therefore exposure to soil) might be similar. For example, details provided in the US studies suggest a relatively high proportion of properties have gardens or unpaved driveways, providing the opportunity for soil contact. In addition, all the US tracer studies were carried out in summer or autumn with the particular locations' summer temperatures generally higher than is typical for New Zealand's temperate maritime climate. But in so far as a favourable temperature indicates a greater likelihood for outside play, the US studies perhaps represent a reasonable year-round estimate for New Zealand. This estimate is probably conservative because, even in New Zealand's temperate climate, outdoor activities will most likely be reduced in winter relative to summer and provide less opportunity for soil contact than the US summer studies would suggest.

If it is accepted that US tracer studies provide the best values for New Zealand and given that single-point estimates are to be used, a value must be chosen that avoids compounding conservatism. The necessary conservatism of SGVs is already obtained by using high-end estimates of other parameters (eg, exposure frequency and duration), and the conservatism of the substance-specific toxicity values. This means that the chosen soil ingestion values should be representing central tendency. The work of Van Holderbeke et al (2007) and others suggests the child ingestion rate of 100 mg/day used in current New Zealand guidelines is rather larger than a central estimate, lying perhaps between the 80th and 95th percentiles.

Central tendency is normally represented by the mean or median. In the case of soil ingestion, the distribution of values is quite skewed in the positive or right direction, with a long tail of a few high values. This results in the mean being considerably larger than the median, with consequent uncertainty as to which best represents central tendency. The median is often chosen as the best measure of central tendency with right-skewed distributions (eg, for house prices or income within a group of people) and is possibly appropriate here. The available studies suggest the median is in the range 20–30 mg/day, say 30 mg/day, with the mean being about twice that value (60 mg/day).

As a pragmatic compromise, a value halfway between the median and mean has been employed as the residential childhood rate, that is, 50 mg/day (rounded up to the nearest 10 mg/day). This falls between the 75th and 80th percentiles of the distribution calculated by Stanek et al (2001) of 42 and 53 mg/day, respectively.

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Soil ingestion – pica children

Deliberate soil ingestion by so-called 'pica children'⁸ is typically not taken into consideration in the estimated soil ingestion rates for children, because this eating disorder is considered to be rare rather than a chronic effect (Paustenbach, 2000). More recently, Paustenbach et al (2006) suggested that the upper bound estimates proposed by Stanek et al (2001) would account for the vast majority of typical children as well as mild or infrequent pica behaviours in 'a conservative risk assessment'.

It is proposed that pica behaviour is not taken into account, as behaviour modification is a more appropriate response. This was the view of the Technical Review Group for the NES (MfE, 2005). However, it may be appropriate to allow for pica behaviour in some site-specific situations, in which case choosing an upper-bound estimate for childhood soil ingestion (say 100 mg/day) may be appropriate for mild cases, but otherwise higher rates should be chosen. Acute toxic effects, which are beyond the scope of this document, may be more likely for pica children.

Residential soil ingestion - adults and older children

There is little data for adult soil ingestion rates; development of appropriate soil ingestion rates is difficult (Paustenbach et al, 2006). Values ranging from 12.5 to 100 mg/day have been used by various regulatory agencies. US EPA (1997) recommends an adult ingestion rate of 50 mg/day as a reasonable central estimate, based on a review of three available studies. Considering the uncertainties in the central estimate, US EPA (1997) was unable to recommend an upper-bound estimate. This work noted that many past US EPA assessments had used 50 mg/kg for industrial settings and 100 mg/day for residential settings. The revised NEPM draft (NEPC, 2010) recommends 50 mg/day is used for adults in low density residential settings, and 12.5 mg/day in high density residential settings. Existing New Zealand guidelines use 25 mg/day for residential scenarios.

The few available studies indicate that adult soil ingestion is likely to be in the order of 5 to 25 mg/day. For example, Paustenbach (2000) concluded that average daily soil ingestion rates of 5 to 25 mg/day were reasonable, based on a review of the limited data. Similarly, Stanek et al (1997) estimated, using tracer studies that mean daily soil ingestion rates (over a four-week period) by adults was 6 mg/day. However, it should be noted that Otte et al (2001) suggested that this average was skewed by a low average in the fourth week of the four-week study, and that taking only the first three weeks' data was more appropriate. This would give rise to an average daily soil ingestion rate of 53 mg/day.

Calabrese's (2003) letter to the General Electric Company, referred to above, also commented on adult ingestion rates. He recommended an upper-bound adult soil ingestion rate of 50 mg/day and a central estimate of 10 mg/day, based on work by him and Stanek (Stanek et al, 1997). Calabrese noted that the US EPA had based their estimates on his earlier work reported in 1990, but that the more recent work was an improvement with more participants (20 instead of 6), a longer study period (28 days instead of 14), and better study design and analysis.

⁸ Pica is a eating disorder characterised by an appetite for substances largely non-nutritive (eg, soil, clay, ash, metal, etc).

Paustenbach et al (2006) reviewed the available studies and found a log-normal distribution of adult soil ingestion appropriate for their probabilistic risk assessment, with a range of 10-100 mg/day and a mean of 30 mg/day. They viewed this distribution as appropriately conservative to address uncertainties with respect to exposure from adult activities with heavier soil contact (eg, gardening or construction).

Van Holderbeke et al (2007), also reviewing the available studies, suggested a median value of 25 mg/day and a mean value of 45 mg/day for the residential scenario with gardens.

Given the uncertainty, it has been decided to retain the adult soil ingestion rate for the residential scenario used in existing New Zealand guidelines, that is, 25 mg/day. This value falls towards the centre of the various estimates cited above.

Soil ingestion – high-density residential

Little or no data exists for soil ingestion for land uses other than the standard residential scenario. Professional judgement must be resorted to.

The high-density residential scenario has a reduced opportunity for soil contact relative to the standard residential scenario: it relates to a multi-unit or townhouse type of development with only small areas of land around the dwelling, and yards being largely paved or grassed, but perhaps with small ornamental gardens allowing some soil contact. The scenario does not include apartment-type developments for which no soil contact is expected. Few overseas jurisdictions provide for the high-density residential scenario. An exception is Australia (NEPC, 2010, revised NEPM draft), which uses 25 mg/day for children and 12.5 mg/day for adults in high density residential settings with limited access to soil.

Two existing New Zealand guidelines have high-density residential scenarios – the 'Gasworks Guidelines' (MfE, 1997) and the 'Sheep-dip Guide' (MfE, 2006a). The former reduced the childhood soil ingestion rate by a factor of four and the adult rate by a factor of five compared with standard residential, for unknown reasons. However, the 'Sheep-dip Guide' used the same rates as the standard residential scenario.

A reduction by a factor of four seems too high for the limited but definite potential exposure contemplated by the definition of the high-density residential scenario. It is therefore proposed as a matter of professional judgement to reduce the standard residential ingestion rates by a factor of two, rounded to the nearest five units. The child soil ingestion rate then becomes 25 mg/day and the adult rate 15 mg/day.

Soil ingestion – parks / recreation scenario

Section 3.1.2, in exploring exposure frequency for the parks / recreation scenario, noted that it is impossible to discuss exposure frequency (or contact rate) without discussing soil ingestion, as likely sub-scenarios tended to be combinations of low ingestion rates and high frequency, or high ingestion rates and low frequency. This approach is repeated here.

As noted in section 3.1.2, Australia and the Netherlands have scenarios similar to the adopted parks / recreation scenario for New Zealand. In the Australian open space scenario the assumption is that adults ingest 25 mg soil/dust per day and children 50 mg/day (NEPC, 2010, draft).

Brand et al (2007), describing proposed revisions to the Dutch CSOIL model, proposed 20 mg/day and 10 mg/day for a child and adult, respectively, for the 'Greens' scenario. These rates are one-fifth of the standard Dutch residential with garden rates. It is notable, however, that the Dutch contact rates (days/year) are much less than typically employed elsewhere, with the combined ingestion and contact rate for Greens being a 25th of the residential combined ingestion and contact rate.

Two existing New Zealand documents, the 'Gasworks Guidelines' (MfE, 1997) and 'Sheep-dip Guide' (MfE, 2006a) have parklands scenarios. Both use 50 mg/day for a child and 10 mg/day for an adult, with exposure for most days (350) of the year. This translates to a combined contact and ingestion rate of about half that for the standard residential scenario.

Section 3.1.2 examined what are considered more realistic exposure scenarios for active recreational activities. An exposure frequency of 350 days per year is excessive for typical active recreational activities where a child might get dirty; upper-bound ingestion rates that might be typical of playing contact sports such as rugby will occur much less often than 350 days per year, even for the keen sportsperson. As a matter of professional judgement, for childhood play activities in green spaces near home (eg, grass-covered road berms and suburban green spaces) it seems reasonable to use an average childhood soil ingestion rate of a half of the standard residential rate with an exposure rate of 200 days per year for threshold substances. This results in soil ingestion rate of 25 mg/day for children. For simplicity, this rate is also applied to older children up to the age of 14 for non-threshold substances. Applying the same rate to older children is probably conservative. As an alternative scenario for non-threshold substances, again as a matter of professional judgement, a reasonable scenario is an adult exposed for 150 days per year (including practices) at an ingestion rate of 75 mg/day to reflect potentially high ingestion rates during high-contact sports such as rugby. The combinations of ingestion rates and exposure frequencies are between about one-third and one-fifth of the residential combination.

Soil ingestion - commercial / industrial scenario

The commercial / industrial indoor worker has no soil ingestion, but has moderate exposure on most working days (up to 230 days per year) while carrying out maintenance activities. This is considered conservative for workers on an unpaved site and also covers occasional (a few times a year) excavation activities associated with site maintenance at higher exposure rates.

Current New Zealand guidelines do not have a similar scenario. The excavation / maintenance scenario in the 'Timber Treatment', 'Gasworks' and 'Oil Industry Guidelines' (MfE and MoH, 1977; MfE, 1977, 1999) is for relatively high soil ingestion of 100 mg/day for 50 days per year. This ingestion rate is from GRI (1988) and is the same as the default recommended for outdoor workers in US EPA (2002a). The rationale for the value in US EPA (2002a) is not given other than it is to reflect higher exposure than for indoor workers, who are assumed to ingest 50 mg/day from indoor dust containing soil tracked in from outside. The latter appears conservative for typical indoor areas. The Dutch assume 10 mg/day on a few days per year for their proposed scenario covering buildings, infrastructure and industry (Brand et al, 2007).

The standard New Zealand scenario for commercial / industrial sites, whether indoor, or outdoor is currently 25 mg/day (sourced from ANZECC, 1992). The revised NEPM draft (NEPC, 2010) also assumes 25 mg/ day for workers on commercial/industrial premises.

The recommended soil ingestion for adult workers involved in maintenance activities that involve routine contact with soil (eg, gardening) is 50 mg/day. This is twice the current New Zealand value but reflects the assumption of greater routine soil contact; it is half the current 100 mg/day excavation / maintenance ingestion rate which assumes regular excavation work. The higher rate could be used for site-specific studies where long-term excavation work is being carried out. The values are based on professional judgement, rather than on scientific studies.

Summary of soil ingestion rates

A summary of the default soil ingestion rates is set out below.

	•					• ·	
Scenario	Rural read	esidential sidential	High-c resid	density ential	Parks / I	recreation	Commercial / industrial outdoo
Receptor	Child	Adult	Child	Adult	Child	Adult	Adult
Central tendency for generic guidelines	50	25	25	15	25	75	50
High-end estimate for site- specific assessment	100	_	50	-	100	100	100

 Table 7:
 Default soil ingestion rates for child and adult (mg/day)

The recommended values for the residential and parks / recreational scenarios are lower than used in the current guidelines. The general effect will be to increase SGVs, ie, be less conservative, although the effect is complicated by revising the toxicological values employed in the derivations for particular contaminants. Some toxicological values have increased and others reduced: this sometimes results in reduced soil ingestion values being outweighed by a reduction in the allowable daily intake of that contaminant. In addition, the effect of the changed soil ingestion rate may be emphasised or muted: this depends on whether the soil ingestion are also significant. Given that the parameters affecting the other pathways have also been revised, there is no direct relationship between SGVs calculated with the soil ingestion rate of 100 mg/day employed in current New Zealand guidelines, and SGVs_(health) calculated with the revised parameters.

For scenarios where home-grown produce consumption is a major pathway, **and** for contaminants that tend to be taken up into plants, the soil ingestion rate has relatively little influence on the SGVs. This is particularly so for scenarios with a high proportion of home-grown produce (see next section). Where the produce ingestion pathway is particularly dominant (eg, for cadmium at low pH), the soil ingestion rate has no influence. However, for the high-density residential scenario without produce consumption, the derived SGV is generally directly or nearly directly proportional to the soil ingestion rate. This applies unless the dermal pathway dominates, in which case the soil ingestion pathway also has little or no influence on the final value.

5.4.2 Produce consumption

Produce consumption can be a significant exposure pathway in residential scenarios, depending on how much of a particular substance is taken up by edible home-grown plants and how much home-grown produce is consumed. A simple calculation can be made using the common value of home-grown produce taking up one per cent of the soil concentration of a contaminant in dry weight terms. If a young child obtains 10 per cent of a typical 10 g/day dry-weight vegetable consumption from this produce, then this is the equivalent of ingesting an additional 10 mg of contaminated soil per day. If 25 per cent of daily produce consumption is home-grown, the same one per cent contaminant uptake equates to 25 mg/day of soil ingestion. These rates are significant relative to the proposed 50 mg/day residential soil ingestion rate for a child.

The produce consumption pathway must consider four parameters:

- the total amount of produce consumed
- the proportion of that consumption from home-grown produce
- the vegetable-type-specific soil loading on the outside of produce
- the chemical-specific uptake factors for contaminants taken up into the produce.

The first three of these parameters are considered below. A general discussion on contaminantspecific produce uptake factors (or bioconcentration factors) is provided in section 5.5.1, while the basis for the specific bioconcentration factors used in the $SCS_{(health)}$ derivations is given in the relevant sections for each contaminant in section 6.

Produce consumption rates

Produce consumption rate is the most country-specific pathway in relation to the amount and type of produce consumed, and the proportion of home-grown produce. Current values for the amount of produce consumed in different countries (where used for deriving guideline values) range from 65 to 151 g/day for a child (table 8).

For adults, current industry-based guidelines in New Zealand use values for the amount and type of produce consumed from the Australian National Dietary Survey 1983. Estimates based on daily nutrient requirements for toddlers (one to three years old) are provided in Langley (1993). These values are also shown in table 8.

 Table 8:
 Produce consumption rates (g FW/day) used in international protocols

Receptor	New Zealand ^a		Australia ^b	US	Canada	Netherlands	UK
	TTG, OIG, GWG	SDG					
Child	130 °	77	243.2	na	125	151	65 ^d
Adult	450 ^b	254	410.5	na	250	295	97 °

a TTG = 'Timber Treatment Guidelines' (MfE and MoH, 1997), OIG = 'Oil Industry Guidelines' (MfE, 1999), GWG = 'Gasworks Guidelines' (MfE, 1997), SDG = 'Sheep-dip Guide' (MfE, 2006a).

b Based on revised NEPM draft (2010).

c Divided into above-ground, roots and fruit.

d Values calculated from fresh weight data in EA (2008a) assuming body weights of 13 kg and 70 kg for 2–4 year toddler and adult, respectively.

na = not applicable

In the 'Timber Treatment Guidelines' (MfE and MoH, 1997) these amounts are divided into leafy vegetables, root vegetables and fruit in the proportion 31:29:40 (100 per cent total). Uptake into fruit is considered to be negligible. In the 'Gasworks Guidelines' (MfE, 1997) the amount of fruit consumption is ignored on the basis that fruit is grown by only a very small proportion of people, and the balance of produce consumption is divided 50:50 into leafy and root vegetables.

The 'Sheep-dip Guide' (MfE, 2006a) updates these consumption rates by considering a variety of vegetables and age-related consumption rates for root and leafy vegetables to produce a weighted-average rate. The approach was adopted from Cavanagh and Proffitt (2005), in which soil guideline values were derived for the Sandilands subdivision in Christchurch. The 'Sheep-dip Guide' continues the assumption that very few people grow their own fruit and therefore this contributes a negligible produce exposure, on average. This assumption would need to be revisited on a site-specific basis in the event of a large amount of home-grown fruit.

The consumption rates in the 'Sheep-dip Guide' are based on data from the 2003/2004 New Zealand Total Diet Survey (Vannoort, 2003). This survey provides consumption rates for a variety of common vegetables based on fresh weights. These have been converted to dry weights; weighted-average dry-weight produce consumption rates for leafy and root vegetables were then calculated for both children and adults. This approach is also used for the derivations in this document, except that root vegetables are divided into true roots (eg, carrot) and tubers (eg, potatoes) for which further details are provided in Appendix 3. Table 9 is a summary of the values used in the SCS_(health) calculations. Note that in the specific case of dioxins, the only vegetable considered is the cucurbit family (eg, marrows and pumpkins).

Produce type	Wet weight (g/day) and percentage		Dry weight (g/day) and percentage		
	Average adult	Average child	Average adult	Average child	
Tuber vegetable	92 (36%)	33 (435)	18.9 (56%)	6.6 (63%)	
Root vegetables	18 (7%)	9 (12%)	1.9 (6%)	1.0 (10%)	
Above-ground vegetables (not including cucurbits)	119 (47%)	25 (33%)	10 (30%)	2.4 (23%)	
Cucurbits (eg, pumpkin)	14 (6%)	6 (8%)	1.4 (4%)	0.46 (4%)	
Subtotal	243	73	32.2	10.46	
Unlikely to be grown at home	10 (4%)	3.6 (5%)	1.4 (4%)	0.57 (5%)	
Total	253	76.6	33.6	11.03	

Table 9:Default produce consumption rates: weights in grams with percentage of
total vegetables given in parenthesis

In dry-weight terms, the percentages of vegetable types likely to be grown at home have been rounded to tuber: 60 per cent; root: 10 per cent; and above-ground including cucurbits: 30 per cent. Where it has been important to differentiate cucurbits, their contribution has been taken as 4 per cent.

Proportion of home-grown produce

For the countries that take consumption of home-grown produce into consideration for their soil guideline values, the proportion of home-grown produce consumed ranges from 10 to about 28 per cent. This difference is largely related to whether average consumption is assumed, or whether the proportion of home-grown produce consumed is based on the small proportion of homes that do grow a significant proportion of their own vegetables. For example, survey data in the Netherlands show home-grown vegetables contribute 54.8 per cent and 13 per cent of the total consumption of vegetables and potatoes, respectively, for about 18 per cent of residents. Expressed as an average of the entire population, home-grown produce contributes 7.1 per cent of the total consumption. This is similar to the value of 10 per cent used in the CSOIL model (Otte et al, 2001; Brand et al, 2007) or the average of 10 per cent applied as a generic estimate for a standard residential scenario in the revised NEPM draft (2010).

A survey in the UK found that a similar proportion (15 per cent) of the population grew their own vegetables. However, in the old CLEA model (Defra and EA, 2002a), the proportion of home-grown produce is based on the six vegetables most commonly consumed by people growing their own. This equates to around 28 per cent of produce consumed for that group, which determines the soil guideline value for all as a high-end value. The updated CLEA model (EA, 2008a) divides the proportion of various produce types grown at home into subgroups by type of vegetable or fruit: leafy, root and tuber vegetables, and herbaceous and shrub fruit. Using survey data, EA (2008a) assumes that between two and six per cent of vegetables, depending on type, and six per cent of herbaceous fruit (eg, courgette, pumpkin, tomato) are grown in a typical home garden situation. However, for those households with allotments, EA (2008a) assumes greater proportions of 13 to 40 per cent for vegetables (depending on type) and 40 per cent for herbaceous fruits.

The *Exposure Factors Handbook* (US EPA, 1997) reports that about 40 per cent of households had a vegetable garden in 1986. Further data on the types of produce grown and consumed by the United States population is also given. This information is intended for use on a site-specific basis as opposed to a generic scenario (US EPA, 1997).

The applicability to New Zealand of produce consumption proportions from overseas is dubious, given the different cultures and opportunities for gardening. However, there is limited information on the proportion of residents who grow their own produce in New Zealand; what information there is comes from specific studies undertaken by local or regional councils. For example, surveys in Hamilton and Christchurch found between about one-fifth and one-third of households grew their own vegetables. One survey, conducted in five new substantial subdivisions near the periphery of Christchurch, found that garden areas where home-grown produce was cultivated ranged from 1 to 64 square metres, with the average slightly under 3 m²: 90 per cent were smaller than 10 m². Estimates from garden guides and personal knowledge suggests that the household garden size required to produce 50 per cent and 10 per cent of home-grown vegetables is of the order of 45–50 m² and 9–10 m², respectively. These areas are consistent with those reported in Defra (2006a) required to grow potatoes for a family.

A similar survey was carried out by the Hastings District Council. Residents of 121 out of 300 properties surveyed in Havelock and Hastings responded to a questionnaire requesting information on area of vegetable garden, types of produce grown, over what period crops were harvested, and an estimate of the percentage vegetables grown at home (Philip McKay, Hastings DC, pers. comm). While not a scientifically robust survey, the combination of area and harvest gave some check on residents' estimates. Sixty per cent of households claimed to grow vegetables, growing on average 23 per cent of their produce at home (although harvesting was not throughout the whole year). However, adjustment for their area of garden often showed there was unlikely to be sufficient garden area to produce the volume of vegetables claimed. After adjusting for garden area, the survey suggested perhaps 6 or 7 per cent of vegetable consumption was grown at home.

Current New Zealand guidelines use proportions of home-grown produce of 10 and 50 per cent for standard and rural residential, respectively. There is no substantial basis for selecting these proportions and further data should be collected. In the absence of this data, estimates of the proportion of home-grown vegetables are largely subjective. Based on limited data and anecdotal information, it is unlikely that more than 10 per cent of the produce consumed by the average urban resident is home-grown. Lifestyle blocks and farms enable a greater proportion of produce to be home-grown and consumed, yet it is debatable whether the 50 per cent proportion used in some earlier guidelines is representative of the current use of lifestyle blocks: it probably tends towards a high-end estimate. The 100 per cent home-grown produce in the agricultural scenario in the 'Timber Treatment Guidelines' (MfE and MoH, 1997) is excessive, being rarely if ever achieved in practice.

In the absence of more definitive data, it is considered appropriate to continue to use a fraction of 10 per cent produce for home-grown produce for the urban residential scenario but reduce the home-grown produce percentage for rural residential from 50 to 25 per cent. The latter is a policy decision rather than a science-based decision and reflects community expectation that rural dwellers have a greater opportunity to grow a higher proportion of their vegetables than urban dwellers. The value of 25 per cent is a compromise between what might be theoretically possible by those intent on following a self-sufficiency lifestyle and what is more likely to occur in reality for most rural-dwellers.

In practice, the site assessor should make a judgement based on site information as to whether some higher proportion is appropriate for a particular site. If a higher proportion than the standard urban or rural residential scenarios is appropriate, a site-specific derivation should be carried out. For conservative illustrative purposes, the derivation of SGVs with 50 per cent home-grown produce is presented in the appendices.

A 50 per cent home-grown produce value is considered sufficiently conservative to also cover uptake into home-grown eggs, except for lipophilic contaminants (eg, dioxins). For sites where such contaminants are at significant concentrations, exposure should be considered on a site-specific basis for home-grown products such as eggs, poultry and dairy.

Soil attached to produce

Soil attached to produce may be one source of inadvertent soil ingestion and should be included in the SGV derivation in some circumstances. There is no need to include attached soil if the contaminant-specific values of the bioconcentration factor (BCF) have been derived from empirical studies or measurements of contaminants in the produce and soil – whether for metals or organics. In that case the attached soil will be 'built in' to the derived BCF. However, consideration of attached soil needs to be included when BCFs are theoretically derived using partitioning relationships and the like, and therefore have not automatically considered attached soil. The following discussion relates to the latter case.

Soil may adhere to the skin of root vegetables (eg, carrots and potatoes) and on leafy vegetables, the latter from direct contact, rain splash or dust deposited on exposed surfaces. Although peeling and/or washing vegetables will reduce the amount of attached soil, it is likely that a residual amount remains. For example, Sheppard and Evenden (1992, cited in Defra and EA, 2002a) estimated the soil loading on thoroughly washed beet leaves was 2 mg soil per gram fresh weight beet (mg soil/g FW).

The old UK CLEA model (Defra and EA, 2002a) used soil loadings from 0.1 mg soil/g FW for root vegetables up to 1 mg soil/g FW produce for stem vegetables for the six vegetables considered in the exposure model. Conversion to dry weight results in soil loadings values of 0.1 per cent for root vegetables (which are assumed to have been peeled) and 1 per cent for leafy vegetables. The revised UK CLEA model (EA, 2008a) assumes dry weight soil loadings of 0.1 per cent for all produce types and then applies dimensionless 'preparation factors' ranging from 0.2 (green vegetables) to 1.0 (roots and tubers) to account for washing and peeling. The Dutch CSOIL model, following revision in 2001, assumes attachment of soil to leafy vegetables due to 'rain splash' amounting to one per cent of the vegetable dry weight. However, this is only applied for organic contaminants, as the empirically derived bioconcentration factors for metals are considered to include an allowance for attached soil (Otte et al, 2001; Brand et al, 2007).

Applying the current UK values to New Zealand produce consumption rates results in the equivalent of an additional soil ingestion rate of about 38 mg/day for an adult and 8 mg/day for a child, if 100 per cent of produce is grown at home (an extreme case), and correspondingly less for lower produce percentages. When converted to a body weight basis the incremental amount is virtually identical for adults and children. The amount of additional soil is significant for high percentages of home-grown produce but only marginally significant for 10 per cent produce (3.8 mg/day for an adult and 0.8 mg/kg for a child).

5.4.3 Dermal exposure

Dermal exposure to soil contaminants can result in acute effects (eg, dermatitis) on the skin, or it may contribute to cumulative exposure. The latter is the most common way to assess dermal exposure for exposure assessments. However, where acute effects occur, it may be appropriate to consider the dermal exposure pathway separately and derive soil guideline values to prevent these effects from occurring.

Three factors are typically used to estimate exposure via dermal absorption:

- the area of skin exposed
- the amount of soil that adheres to the skin
- the absorption rate of the individual contaminant.

The first two are discussed in more detail below and the contaminant-specific absorption factor is discussed in section 5.5.2.

Area of skin exposed

The area of skin exposed depends on the receptor considered (ie, child or adult) and on assumptions about what parts of the body are exposed. Table 10 provides a summary of the area of skin exposed, and the assumed exposed body parts used in national and international protocols. Most protocols assume exposure of the face or head, hands, forearms and lower legs of child residents, and exposure of fewer body parts for adult residents and workers.

The value for exposed skin surface area used in current New Zealand industry-based guidelines, except the 'Sheep-dip Guide', is based on Langley (1993), which in turn was based on ATSDR data (ATSDR, 1992). The ATSDR data assumes that 30 per cent of the average total skin surface area for a child aged 1–11 years is exposed, while 24 per cent of the skin surface area of an adult male is exposed. Older US EPA guidance (US EPA, 2001a) used a value of 2800 cm² for the exposed skin surface area for children, which corresponds to exposure of about 43 per cent of the total skin surface area for children aged 0–6 years (6560 cm², average of 50th percentile values for males and females). This value is based on data provided in the Exposure Factors Handbook (US EPA, 1997) for the skin surface area because the skin surface area of a three-year-old child is used to estimate the skin surface area for children aged less than three years. Skin area used in existing New Zealand and international protocols is summarised in table 10.

A revised approach to dermal exposure has more recently been taken by US EPA (2004c), where the amount of skin exposed depends upon the exposure scenario. Clothing is expected to limit the extent of the exposed surface area in cases of soil contact. All skin area estimates are 50th percentile values to correlate with average body weights used for all scenarios and pathways. Skin area is closely correlated with body weight.

Body-part-specific skin areas are presented in US EPA (2004c) calculated for an adult (>18 years old) and a child (<1 to <6 years old), based on data from the Exposure Factors Handbook (US EPA, 1997). No New Zealand data exists for skin area. Given skin area is related to weight and it has been decided to use a 13 kg child and a 70 kg adult, the US EPA data may be used to calculate skin areas. Any error between New Zealanders and Americans will be small compared with errors elsewhere in the dermal exposure pathway (such as the contaminant-specific absorption factors).

Pathway	Child		Ac	dult – resident	Adult – worker	
	Exposed skin surface area (cm ²)	Area exposed	Exposed skin surface area (cm ²)	Area exposed	Exposed skin surface area (cm ²)	Area exposed
New Zealand (based on Langley, 1993)	2,625	30% of total body surface area for ages 1–11	4,700	24% of total body surface area for males	4,700	24% of total body surface area
Environmental Risk Management Authority New Zealand	14,000	100% skin surface of 10-year-old child	4,500	25% of total body surface area averaged for females and males: assumes head, hands, forearms, lower legs and exposure under clothing	NA	
US EPA (2001a)	2,800	Head, hands, forearms, lower legs, feet	5,500	Forearms, head, hands, lower legs	3,300	Forearms, head, hands
Canada	2,600	Head, arms, hands, lower legs	4,300	Head, arms, hands	4,300	Head, arms, hands
UK1	466	23% hands, forearms, lower legs	293	Hands 5%	293	Hands 5%
Netherlands	2,800 50	Outdoor Indoor	1,700 90	Outdoor Indoor	NA	

Table 10:Area of exposed skin and body parts considered to be exposed for different
receptors, used in national and international protocols

1 Assumes that soil adheres to only one-third of the total exposed skin surface (Defra and EA, 2002a). NA = not applicable.

US EPA (2004c) assumes a child resident will be wearing a short-sleeved shirt and shorts, but no shoes. Therefore, the exposed skin is limited to face, hands, forearms, lower legs, and feet. This is probably a high-end estimate for year-round exposure, more clothes being worn in the winter. Arguably, it may be appropriate to reduce the average skin area by some factor (eg, two) to allow for more clothes in winter; however, some people may tend to wear fewer clothes all year round in the warmer parts of New Zealand. The dermal exposure route is generally minor and attempting to modify the skin areas for different times of the year is probably an unnecessary refinement.

Table 11 shows the body-part-specific skin areas for a child aged 1-2 and 2-3 years, the two age ranges closest to a 13 kg weigh, taken from US EPA (2004c). The mean of the two ranges will sufficiently approximate a 13-kg child. The total, 1900 cm², is proposed as the skin area for a child in both residential and recreational settings. This is smaller by about 30 per cent than all the existing New Zealand guidelines.

Table 11:	Body parts exposed for a child and associated skin areas (c	;m²)
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Age	Face	Forearms	Hands	Lower legs	Feet	Total
1–2	325	346	336	544	371	1,921
2–3	280	314	313	550	418	1,874
Mean	302	330	324	547	394	1,897

US EPA (2004c) assumes the adult resident wears a short-sleeved shirt, shorts and shoes; therefore, the exposed skin surface is limited to the head, hands, forearms and lower legs. The rationale for including the head area is not given. The average skin area for males and females for the face, hands, forearms and lower legs is given as 4850 cm^2 in US EPA (2004c), based on data from US EPA (1997). It is proposed to use this value for the adult resident.

The adult commercial / industrial worker was assumed to wear a short-sleeved shirt, long pants, and shoes. Therefore, the exposed skin surface was limited to the face, hands and forearms. It is probable that in New Zealand an outdoors worker would commonly wear shorts in summer. It is therefore proposed that the skin area be the total of face, hands, forearms and half the area of the lower legs, the latter to allow for the lower legs being exposed half the year. For the outdoor worker the area will therefore be 3670 cm². This skin area has also been adopted for adults playing outdoor sports throughout the year, on the assumption that wearing shorts is common while participating in active sports.

Soil adherence

The amount of soil that adheres to the skin will influence dermal exposure to soil contaminants. This factor differs for different activities and skin surfaces (eg, hands, forearms, lower legs). The soil adherence factor currently used in existing New Zealand guidelines, except the 'Sheep-dip Guide' for the residential scenarios (0.5 mg/cm²), is originally sourced from Hawley (1985) and is based on the amount of soil adhering to a child's hands. A greater amount of soil adheres to hands and feet compared to other parts of the body (eg, face, forearms, legs) so extrapolating the soil adherence factor based on hands to the rest of exposed skin overestimates soil adherence. For this reason, recent US EPA guidance for estimating dermal exposure has proposed revised soil adherence factors expressed as surface-area-weighted values.

US EPA (2004c) presents the data from a number of studies of children playing in soil and in childcare centres, and for adults carrying out various activities: mean and 95th percentile soil adherences have been calculated for various body parts. Body-part-weighted soil adherences for children and adults have then been calculated, using the body parts exposed as in the previous section. The central tendency and 95th percentile values are reproduced in table 12 for standard residential and industrial scenarios.

The 95th percentile values for adults and children from US EPA (2004c) were used for the existing 'Sheep-dip Guide' (MfE, 2006a). However, as a policy decision it had been decided to use central estimates for these factors in that protocol. The resulting skin adherences for adult resident and worker are 0.01 mg/cm^2 and 0.02 mg/cm^2 , respectively, and for a child 0.04 mg/cm^2 .

These values are not necessarily appropriate for all the proposed scenarios, particularly the recreational and outdoor worker scenarios: here soil adherence is expected to be significantly greater than the residential or the US EPA industrial scenario. US EPA (2004c) provides weighted-average values for various outdoor and sports activities. A selection of the activities is also shown in table 12.

Receptor	Central tendency	95th percentile
Residential adult	0.01	0.07
Industrial	0.02	0.2
Grounds keepers	0.02	0.1
Gardeners	0.1	0.5
Utility workers	0.2	0.1
Rugby player	0.1	0.6
Soccer player	0.01	0.08
Residential child	0.04	0.2

Table 12: Estimated soil adherence factors (mg/cm²)

Source: US EPA (2004c).

The residential value for a child is proposed for all scenarios involving a child, except the highdensity residential scenario – for which an arbitrary halving of the residential value is proposed. The high-density residential skin adherence value for adults is also arbitrarily half the standard residential value.

For the adult recreational scenario, an average between summer and winter is needed. Using a rugby player as the basis and assuming an adherence of 0.01 mg/cm^2 for summer, an average soil adherence of 0.06 mg/cm^2 has been adopted.

For most substances, dermal exposure is a minor pathway and the exact soil adherence values will make little difference to the SGVs for the residential and child recreational scenarios.

The proposed outdoor worker scenario is expected to cover a caretaker carrying out a variety of routine outdoor maintenance work, including gardening and occasional excavation; it will be conservative for a worker on an unpaved commercial / industrial site. A combination of soil adherence values for grounds keepers (most of the time), gardeners (once a week), and utility workers (five times per year) results in a weighted-average value of 0.04 mg/cm². The proposed values are summarised in table 13.

Table 13: Soil adherence values (mg/cm²)

Scenario	Adult	Child
Rural and residential	0.01	0.04
High-density residential	0.005	0.02
Parks / recreational	0.06	0.04
Outdoor worker	0.04	_

5.4.4 Inhalation of particulates and volatiles

As noted earlier, exposure from inhalation of particulates is so small relative to the soil ingestion pathway that it is not worth calculating for most generic SGVs_(health). Using the various parameters from the existing 'Timber Treatment Guidelines' (MfE and MoH, 1997) for illustrative purposes, the high dust environment of the commercial / industrial maintenance scenario contributes only 1.6 per cent to the combined soil ingestion and inhalation pathways. However, the exposure contribution from the inhalation pathway should be checked where the inhalation reference health standard is orders of magnitude lower than the oral reference health standard. In addition, there may be particularly dusty situations where inclusion of the inhalation pathway in site-specific assessment is appropriate. Inhalation rates are therefore summarised here for these situations.

Inhalation rates are also important for assessing the effects of inhalation of volatile compounds. While $SCSs_{(health)}$ for volatile compounds are not being calculated in the current document, the inhalation rates given here are appropriate for that purpose.

Inhalation rates

Inhalation rates for existing guidelines for various countries are summarised in table 14. A direct comparison between the guidelines is difficult because of the way the different models and methodologies use the values: as a combination of intake values, worst case for child and adult, combining indoor and outdoor, or treating indoor and outdoor separately for volatiles.

The Exposure Factors Handbook (US EPA, 1997) provides a review of inhalation studies. From that review a summary of inhalation rates for children of various ages and adults in different levels of activity was developed. The summary is reproduced in table 15. Daily rates are presented for long-term exposures, and hourly rates for short-term activities, as might be expected in an outdoor occupational setting when carrying out physical work.

The long-term average daily rates are applicable for residential setting, and are lower than the default values adopted in the Soil Screening Guidance (US EPA, 1996a, 1996b, 2002a) for adults as well as children (assuming the 15-kg child falls within the 1–2 year age range). The US EPA (1997) acknowledges this and recommends the summary values to be used in site-specific assessment; and also recommends that mean values, rather than high estimates, are employed. Taking the average of the male and female long-term rates as appropriate for a 70-kg adult, a rate of 13.3 m³/day is obtained, whereas 6.8 m³/day is appropriate for a 15-kg child in a residential setting.

The former value is lower than rates used in current New Zealand guidelines. The $20 \text{ m}^3/\text{day}$ used for the calculation of outdoor inhalation of volatiles in the 'Gasworks' and 'Oil Industry Guidelines' (MfE, 1997, 1999) is particularly unrealistic. Based on the US EPA figures, it suggests that on average, adults would spend a good part of their daylight hours outside carrying out moderate to heavy activities; whereas the reality is that an average adult is more likely to spend most time either inside the house or off-site at work.

It is notable that the Dutch CSOIL model assumes that an adult spends only an average of one hour outside each day at home, whereas the UK model does not consider adult inhalation exposure: a child is considered the critical receptor instead.

The US EPA young child rate of $6.8 \text{ m}^3/\text{day}$ is nearly twice that currently used in all the New Zealand guidelines except the 'Sheep-dip Guide', which uses a rate identical to the Dutch rate of 7.6 m³/day (although no source is cited for the value). The UK child inhalation rate falls between the New Zealand rates. It is proposed that a child rate of $6.8 \text{ m}^3/\text{day}$ be used for child inhalation in a residential setting.

When the methodology for calculating soil guidelines for exposure to volatiles is reviewed, the apportionment of the inhalation rate to indoor and outdoor residential activities will need to be considered.

For adult occupational exposure, the Exposure Factors Handbook (US EPA, 1997) recommends a 1.3 m^3 /hour average for outdoor workers. Using this rate, an occupational exposure of eight hours gives a daily inhalation rate of 10.4 m³/day. This is similar to the 9.6 m³/day values used in existing New Zealand industry-based guidelines. An indoor worker is likely to have an

inhalation rate of between 0.5 and 1 m^3 /day (table 15). Taking the higher rate as conservative, results in an indoor worker inhalation rate of 6 m^3 /day.

Inhalation of either particulates or volatiles is not likely to be a significant factor for the parks / recreation scenario and therefore no generic inhalation rate needs to be determined. Case-by-case determination of inhalation rates is recommended for the rare occasions when unusually high concentrations of particulates or volatiles are present in a parks / recreational setting, as part of site-specific assessment.

	New Zealand ^a				USA	Canada	Netherlands	UK
	TTG	GWG	OIG	SDG	SSL	SQG	CSOIL ^b	CLEA °
Residential								
Child indoor	3.8 ^d	3.8 ^d	3.8 ^d	7.6 ^d	10 ^d	_	6.7	5
Child outdoor							0.9	
Adult indoor	20 ^d	15	15	25 ^d	20 ^d	_	19	-
Adult outdoor		20	20			23	1	-
Parkland / recreation								
Child	-	1.1	-	7.6	-	5	0.32	-
Adult		2.4		25		23	0.83	
Commercial Industrial								
Adult indoor	9.6 ^d	10 ^e	10 ^e	10.4 ^d	20 ^e	23 ^e	5	4.7 [†]
Adult outdoor		10	10		20	23	0.8	0.95

 Table 14:
 Summary of daily inhalation rates in existing New Zealand and international guidelines (m³/day)

a TTG = 'Timber Treatment Guidelines' (MfE and MoH, 1997), GWG = 'Gasworks Guidelines' (MfE, 1997), OIG = 'Oil Industry Guidelines' (MfE, 1999), SDG = 'Sheep-dip Guide' (MfE, 2006a).

b CSOIL model calculates a combination of child and adult and indoor and outdoor exposure.

c Calculated using means values of probability distributions of inhalation rates and exposure times.

d No differentiation between indoor and outdoor.

e Soil guideline values for indoor and outdoor exposure calculated as separate scenarios.

f Calculated as combined indoor and outdoor exposure.
Short-term exposu	Long-term exposures			
Population	Mean rates (m ³ /hour)	Population		Mean rates (m³/day)
Adults		Children		
Rest	0.4	Infants <1 year		4.5
Sedentary activities	0.5	1.0		0.0
Light activities	1.0	1–2 years		6.8
Moderate activities	1.6	3–5 years	8.3	
Heavy activities	3.2	6–8 years		10
Children		9–11 years	Male	14
Rest	0.3	-	Female	13
Sedentary activities	0.4	2–14 years	Male	15
Light activities	1.0		Female	12
Moderate activities	1.2	15–18 years	Male	17
Heavy activities	1.9	2	Female	12
Outdoor workers		Adults		
Hourly average	1.3	19-65+ years	Female	11.3
Slow activities	1.1		Male	15.2
Moderate activities	1.5			
Heavy activities	2			

 Table 15:
 Inhalation rates recommended in Exposure Factors Handbook

Source: US EPA, 1997.

Particle emission factor

Existing New Zealand guidelines use particle emission factors based on US EPA studies and Australian measurement of dust concentrations (eg, see MfE and MoH, 1997 – chapter 5, Appendix C). The assumptions in MfE and MoH (1997) around what proportion of respirable dust comes from the contaminated site appear to have little basis. For example, it is assumed that 20 per cent of respirable dust is from soil sources, based on Australian measurements; this ignores the probability that for all but the largest sites, much of the soil in air-borne dust will be from uncontaminated off-site sources.

Consideration of the particulate inhalation pathway is only necessary for site-specific assessment of dusty sites. Rather than use generic particulate emission factors, it is therefore more appropriate to directly measure site-specific dust concentrations and the contaminant concentrations within the dust, and use these to estimate the intake of dust-borne contaminants.

5.5 Contaminant-specific factors

5.5.1 Plant uptake factors

Perhaps the greatest uncertainty in determining uptake of a contaminant in produce is selecting the plant uptake factors, otherwise known as bioconcentration factors (BCF).

Plants can accumulate contaminants via a number of pathways, the most important of which is typically absorption by roots (Kabata-Pendias and Pendias, 2000). Uptake of organic contaminants and metals occurs predominantly from the soil solution. Normally the concentration of a contaminant measured in the soil solution represents only a fraction of the total contaminant present in the soil. The ratio of the concentration in soil solution to the total in soil depends on a number of factors including soil pH, redox potential, soil organic matter, and soil texture (Kabata-Pendias and Pendias, 2000). In soils and sediments where the clay content is relatively low, the availability of organic contaminants is strongly related to the fraction of organic carbon present.

The uptake of contaminants from soil and transport within plant tissues also differs for different plant species, as well as for different contaminants and different soil types. Ideally, because of the complexity of soil-plant systems, the concentration of soil-derived contaminants in vegetables would be based on site-specific measured data. However, this is not practical for deriving generic assessment criteria, and there is a need to provide mechanisms for estimating plant uptake to enable the derivation of such criteria. A general review of the various BCF derivation methods is provided here, while specific discussion on the contaminant-specific BCFs used for each SCSs_(health) derivation is provided against each contaminant in section 6.

The simplest empirical relationship for estimating plant uptake of soil contaminants follows the form:

 $C_P = BCF \times C_s$

,

where C_P = concentration in produce (mg/kg)

 C_s = concentration in soil (mg/kg)

BCF = bioconcentration factor – typically expressed as the ratio of the contaminant content in produce (mg/kg dry weight) and soil (mg/kg dry weight); if either produce or soil concentration is other than dry weight, the BCF units will reflect this.

Typically the BCF for metals is based on field or laboratory experiments, while the BCF for organics is more often estimated from the octanol-water partition coefficient (K_{ow}) .⁹

For metals where field or laboratory data is available, point estimates of the BCF for a given metal have frequently been used by regulatory agencies. A number of different point estimates may be used including:

- geometric mean / median / mean of all available data
- geometric mean / median / mean of available data for selected vegetables of interest
- separate point (geometric mean) estimates for root and stem vegetables
- weighted-average BCF based on the consumption rates of different vegetables.

⁹ The octanol-water partition coefficient is a measure of the relative affinity an organic compound has to bind to organic material (eg, in soil) or dissolve in water.

Which point estimate is used depends on the available data and the methodological approach used to estimate contaminant intake via consumption of home-grown produce. For example, the UK bases estimates of contaminant intake from produce consumption on six vegetable and fruit groups that are commonly grown and consumed in the UK (EA, 2008a). In contrast, a consumption-weighted BCF that is also dependent on soil properties has been proposed for use in the Dutch CSOIL model (Otte et al, 2001).

For organics, there is a paucity of data on plant uptake, hence estimation of the plant uptake of organic contaminants is typically undertaken using models based on the octanol-water partition coefficient (K_{ow}) of the individual contaminants. These models tend to over-predict the bioconcentration factors compared to real data (Otte et al, 2001). The models include the relatively simple model of Travis and Arms (1988), which is an empirical relationship between the bioconcentration factor for organic contaminants in above-ground plants parts and the K_{ow} :

 $\log B_v = 1.588 - 5.78 \log K_{ow}$

where: B_v = bioconcentration factor for above-ground plant parts (dry weight basis) K_{ow} = octanol-water coefficient.

More complex models, such as that of Trapp and Matthies (1995), consider the sorption of contaminants to plant fats. See Rikken et al (2001) and EA (2006) for further discussion on plant uptake models for organic contaminants. While a considerable amount of research on plant uptake of organic contaminants has been undertaken, there is generally a lack of agreement with measured data (EA, 2006).

EA (2006) reviewed a number of models for the uptake of organic compounds in plants, comparing their predictions with actual measurements in the literature of uptake into common vegetable types from contaminated soil in both field and laboratory studies. Unfortunately, many of these studies were limited to persistent chemicals such as polycyclic aromatic hydrocarbons, polychlorinated biphenyls and dioxins / furans. Little data was available on common industrial chemicals, including petroleum hydrocarbons and low-molecular-weight chlorinated compounds. The authors were surprised to find they had to reject many studies because these did not report whether the results were on a fresh-weight or dry-weight basis, or whether the measurements were of roots, shoots, fruits or tubers. This indicates a general need for care if attempting to select representative uptake factors from the literature for a particular substance.

EA (2006) concluded that model performance was highly variable. All except one of the six models reviewed over-predicting root uptake by a least an order of magnitude. The exception, Travis and Arms (1988), only applies to above-ground parts.

Given this, the recommended approach is to simply use BCFs based on available data, and only resort to models when measured values are not available. As noted above, bioconcentration factors for the particular contaminants considered in this protocol, and justifications for the chosen values, are given for each $SCS_{(health)}$ derivation in section 6. Two elements excepted, all the $SCS_{(health)}$ derivations in this document use BCFs estimated from experimental data specific to the contaminant; BCFs were not used for the derivation of the copper and boron $SCSs_{(health)}$.

5.5.2 Bioavailability

Oral bioavailability is the fraction of a substance which, following ingestion, is actually absorbed and reaches systemic circulation. It is commonly expressed as the ratio (or percentage) of absorbed dose to ingested dose.

Bioaccessibility is the fraction of a substance that is soluble following gastrointestinal extraction and is therefore available for absorption. This term is specifically used when *in vitro* (literally, 'in glass', ie, in a test tube or petri dish) laboratory methods are used to estimate the relative bioavailability of a contaminant.

Until the science is better developed for New Zealand soils and conditions bioavailability considerations in site-specific assessments are not appropriate. Any adoption of site-specific assessment using reduced bioavailability in New Zealand should use a multiple lines of evidence approach. At present, the science does not support *in vitro* testing for other than lead and perhaps arsenic, despite wider use overseas. The present knowledge within the contaminated land community in New Zealand, both practitioners and regulators, is insufficient to give confidence that bioavailability test results would be applied correctly.

The recent review of the NEPM in Australia has included a review of the science on bioavailability and the available bioaccessibility tests. The draft policy resulting from the NEPM review is that oral bioavailability for lead and arsenic is assumed to be 50 per cent and 70 per cent respectively. This review has gone further than some other jurisdictions in accepting sufficient validation has been carried out for both lead and arsenic. For arsenic, this is based on parallel *in vitro* and *in vivo* (Latin for 'within the living', ie, using a whole, living organism) testing for a small number of Australian soil samples, and overseas research.

Non-statutory guidance associated with the NEPM provides specific equations for lead and arsenic to convert bioaccessibility results to relative bioavailability values. Unlike the UK and the US, a multiple lines of evidence approach is not being advocated in this guidance nor in the draft NEPM documents.

Much of the overseas research has been carried out on mining wastes and soils affected by metal smelting activities. Samples have contained relatively high contaminant concentrations. It is unknown whether such soils have similar behaviour to typical contaminated soils encountered in New Zealand, most of which are not associated with the primary metal production industry. This introduces a considerable uncertainty as to whether the *in vitro* – *in vivo* relationships developed in Australia and elsewhere are appropriate to New Zealand soils. Given expensive *in vivo* studies are unlikely to ever be justified for New Zealand sites, this uncertainty is unlikely to be resolved.

Overseas interlaboratory studies have shown significant differences between laboratories using the same *in vitro* test method. Small differences in the nominally identical methods may result in significantly different results. This introduces further uncertainty as to the applicability of the overseas-derived relationships, which may be specific to the particular laboratories in which the tests were carried out.

Reduced oral bioavailability only affects the soil ingestion pathway. It should not be applied to all pathways. Its influence on a site-specific SGV will depend on how significant the soil ingestion pathway is to the overall SGV. Soil ingestion is generally a large influence on most SGVs. The residential SGVs for cadmium are an exception.

5.5.3 Skin absorption factors

Dermal uptake of soil contaminants is largely dependent on the physico-chemical properties of individual contaminants, although it can be modified by soil loading on the skin and exposure duration (eg, McKone, 1990).

An absorption factor (expressed as a percentage or fraction) is generally used to estimate dermal uptake when deriving generic soil guideline values. Factors specific to the individual priority contaminants are given in MfE (2011), where more detailed discussion is provided for the individual priority contaminants. A brief general discussion is provided below.

Absorption rates need to be determined on the basis of available information, which is often limited. Further, dermal absorption studies are often only available for animals, and the dermal permeability of different animals can be different to that of humans. Pig and guinea pig are suggested to be most representative of absorption across human skin (McKone, 1990). Expert judgement is required to interpret available dermal studies. Adsorption on soil particles may reduce the dermal uptake of some contaminants: eg, Hawley (1985) suggested that only 15 per cent of contaminants adsorbed onto soil would be dermally absorbed. Yet for some contaminants this reduction in dermal uptake may be insignificant. For example, pentachlorophenol (PCP) from contaminated soil was absorbed at a similar rate to PCP in acetone, with 24 per cent of PCP absorbed over a 24-hour period (Wester et al, 1993).

Detailed models for estimating the dermal absorption of (primarily organic) contaminants (eg, McKone, 1990; US EPA, 2001a) are available. These models take into account factors such as skin thickness, chemical properties of the contaminants (eg, octanol-water partition coefficient, K_{ow}), and soil loadings. Generally, these models are unnecessarily complex in terms of developing a generic soil guideline value. Nevertheless in the absence of data for specific contaminants, the model results can provide a useful insight into potential dermal uptake of contaminants. For example, based on model estimates from McKone (1990) for compounds with a log K_{ow} of 6 and below and a K_h (Henry's Law Coefficient) below 0.001, it is reasonable to assume 100 per cent absorption in 12 hours. For compounds with a K_h of 0.01 and above, the uptake fraction is unlikely to exceed 40 per cent in 12 hours; for contaminants with a K_h of 0.1 and above, uptake of no more than 3 per cent in 12 hours is expected.

In the case of $SCSs_{(health)}$ derived in this document, dermal absorption factors were determined for seven of the 12 contaminants. Of these seven, the dermal pathway had no significant effect on the derived SCSs for two of the contaminants (arsenic and cadmium), a marginal effect on the SCSs for four of the contaminants (benzo(a)pyrene, DDT, dieldrin and dioxins), and a significant effect on the derived SCS for PCP. In all cases, dermal absorption factors were obtained from the literature, rather than resorting to models.

5.6 Summary of exposure factors

A summary of the recommended general and scenario-specific exposure parameters is presented in table 16.

Generic factors								
Body weight (child):	ody weight (child): 13 kg Averaging time (non-three							
Body weight (adult):	70 kg		Averaging time	e (threshold):	6 years			
Scenario-specific factors	Lifestyle block	Standard residential	High-density residential	Parks / recreational	Commercial / industrial indoor worker	Commercial / industrial outdoor worker	Unit	
Exposure frequency	350	350	350	200	230	230	day/year	
Exposure duration (child)	6	6	6	6			years	
Exposure duration (adult)	24	14	14	8	20	20	years	
Soil ingestion rate (child)	50	50	25	25			mg/day	
Soil ingestion rate (adult or older child)	25	25	15	25	0	50	mg/day	
Age-adjusted ingestion factor	31.6	28.1	14.5	14.4	0	14.3	mg year/kg day	
Inhalation rate (child)	6.8	6.8	6.8				m³/day	
Inhalation rate (adult)	13.3	13.3	13.3		8	10.4	m³/day	
Age-adjusted inhalation rate	7.7	5.8	5.8		2.3	3.0	m ³ year/kg day	
Particulate retention	0	0	0	0	0	0	dimensionless	
Particle emission factor	-	-	-	-	-	_	m³/kg	
Skin area (child)	1,900	1,900	1,900	1,900			cm ²	
Skin area (adult)	4,850	4,850	4,850	3,670	3,670	3,670	cm ²	
Soil adherence (child)	0.04	0.04	0.02	0.04			mg/cm ²	
Soil adherence (adult)	0.01	0.01	0.005	0.04	0	0.04	mg/cm ²	
Age-adjusted dermal exposure factor	51.7	44.8	22.4	51.9	0	41.9	dimensionless	
Produce ingestion (child)	0.0105	0.0105					kg/day (DW)	
Produce ingestion (adult)	0.0322	0.0322					kg/day (DW)	
Proportion of above- ground produce	0.3	0.3	0.0	0.0	0.0	0.0	dimensionless	
Proportion of root (not tuber) produce	0.1	0.1	0.0	0.0	0.0	0.0	dimensionless	
Proportion of tuber produce	0.6	0.6	0.0	0.0	0.0	0.0	dimensionless	
Age-adjusted produce ingestion	0.0159	0.0113	0.0	0.0	0.0	0.0	kg year/kg day	

Table 16: General and scenario-specific exposure parameters

DW = Dry weight.

6 Soil Contaminant Standards for Selected Priority Contaminants

Soil contaminant standards have been derived for a group of priority contaminants, specifically arsenic, boron, cadmium, chromium, copper, inorganic lead, inorganic mercury, benzo(a)pyrene, DDT, dieldrin, dioxin (as 2,3,7,8-tetrachlorodibenzo-*p*-dioxin) and dioxin-like polychlorinated biphenyls (PCBs), and pentachlorophenol. All calculations use the updated contaminant-specific toxicological, background intake and skin absorption factors presented in MfE (2011) and produce bioconcentration factors presented in each contaminant-specific section below.

The details of the derivations are provided in appendix 1 (and appendix 2 for additional calculations for cadmium); summary results for each contaminant are provided against each contaminant below and in an overall summary for all contaminants in section 7.

In the $SCS_{(health)}$ summaries, values have generally been rounded down to two significant digits, with some exceptions for pragmatic reasons. If the value was close to a round number, the value has been rounded up or down to that number, eg, 69 to 70. Values in the hundreds have generally been rounded to the nearest 10 unless close to a multiple of a hundred, eg, 410 becomes 400.

For values between an arbitrary 30 and 100 mg/kg, rounding has been to the nearest 5 mg/kg. Values below 0.1 mg/kg have been rounded to 1 significant digit. Dioxins are an exception, as these are in units of μ g/kg and have been rounded to two significant digits.

Derived values greater than 10,000 mg/kg have been shown as 'no limit' (NL). In practice, such high concentrations are unlikely to be found on most sites. The derived values may be found in Appendix 1.

A 'no limit' value is also shown for the commercial / industrial indoor worker scenario in the $SCS_{(health)}$ summary table for each contaminant. These are not derived values, but are based on the assumption that there is no exposure.

The soil contaminant standards are intended to be compared with the results of chemical analysis of soil sample results (dry weight basis). Guidance on how to compare laboratory results of soil analyses with SGVs is provided in MfE (2004) (and updates).

In summary, for metals and metalloids, the chemical analysis sample preparation should be the equivalent of 'total recoverable' (eg, equivalent to US EPA Method 200.2 (USEPA 1994)). More aggressive digestion techniques or x-ray fluorescence analysis techniques that give 'total' concentrations are also permissible, but the results from such techniques will be a more conservative comparison than results from total recoverable techniques. Comparison of $SCSs_{(health)}$ with the results of analyses using less aggressive extraction techniques, such as may be used for trace element analysis of agricultural soils (eg, extraction using ethylenediaminetetraacetic acid – EDTA), is not appropriate.

For analysis of organic compounds the objective is to determine the total amount in the soil. There are many methods available for both extraction from the matrix and detection following extraction. The analytical technique should be discussed with the laboratory to ensure that it is suitable for the target compounds and matrix. International Accreditation New Zealand (IANZ) (or equivalent) accreditation of the laboratory for the method should provide reassurance that the analysis will perform as intended.

6.1 Arsenic

6.1.1 Bioconcentration factor for arsenic

The US EPA (2007) reports a median bioconcentration factor (BCF) of 0.03752 for arsenic, determined from 122 data points with a range of 0.00006 to 9.0741 (Bechtel-Jacobs, 1998). The 'Timber Treatment Guidelines' (MfE and MoH, 1997) use a bioconcentration factor of 0.01 for root vegetables, and 0.002 based on the assumption that bioaccumulation in vegetative parts is one-fifth of that in the roots. In contrast, Baes et al (1984) found a higher BCF in vegetative parts (0.04) as compared to non-vegetative parts (0.006, tubers, seeds, fruits). Dutch agencies have used a BCF of 0.021 in the derivation of the current intervention value, although this has been revised and a BCF of 0.009 has now been proposed for use (Lijzen et al, 2001). This value is a consumption-weighted BCF, however, and the median value of the revised data set is 0.025. The Environment Agency (EA, 2009a) provides the most recent summary of plant BCFs. Their report reviews the existing literature and provides recommendations for BCFs (based on fresh weight) for the edible portion of six produce types, which are used in the CLEA model for the available data for each produce type.

The BCFs recommended in EA (2009a) are lower than those provided in Bechtel-Jacobs (1998) and used in Otte et al (2001). Close inspection of the Bechtel-Jacobs data indicates that most of it relates to non-vegetables, and thus has arguable application to the current work. The data in Otte et al (2001) appears to be based only on three data points. For the purpose of deriving soil guideline values protective of human health, the recent EA (2009a) review could be used in the derivation of New Zealand soil guideline values. However, there is some New Zealand-specific data (Gaw et al, 2008), which is not included in the EA (2009a) review which indicates a higher BCF to be appropriate. As such, the original references used in EA (2009a) were gathered where possible, and this data along with that of Gaw et al (2008) was used to derive BCFs. The derived BCFs are shown in table 17 along with BCFs determined from other sources (MfE, 2010b).

Table 17: BCF for arsenic from different sources

MfE and I	MoH, 1997	Du (Otte et	tch al, 2001)	US EPA (2007)	UK (EA, 2009) Fresh weight (dry weight) ²		٦	This study	y	
Root	Leafy	CW ¹	Median		Green (n=46)	Root (n=26)	Tuber (n=6)	Green (n=29)	Root (n=16)	Tuber (n=2)
0.01	0.002	0.009	0.025	0.0375	0.00043 (0.0043)	0.0004 (0.004)	0.00023 (0.001)	0.011	0.011	0.001

1 Consumption-weighted BCF.

2 Calculated using conversion factors of 0.096 kg DW/kg FW, 0.103 kg DW/kg FW and 0.21 kg DW/kg FW for green, root and tuber vegetables, respectively (EA, 2009b).

It should also be noted that Swartjes et al (2007) recently concluded that use of a fixed value for concentration of arsenic in vegetables, that is independent from arsenic soil concentrations and soil properties, is most appropriate for assessing the human health risks due to the consumption of produce from contaminated soil. This is because no significant relationship between vegetable concentration and soil concentration exists.

6.1.2 Calculations for arsenic standard

The calculations for the soil contaminant standard on arsenic use the contaminant-specific parameters in table 18 and the derived $SCSs_{(health)}$ are set out in table 19. Arsenic has been treated as a non-threshold substance and therefore a risk-specific dose has been employed. A single BCF value was used for both root and leafy parts of vegetables, with a separate factor for tubers; these are combined into a single consumption-weighted mean value, using weighting factors of 0.4 and 0.6 (dry weight), respectively.

 Table 18:
 Contaminant-specific parameters for the derivation of the arsenic soil contaminant standard

Risk-specific dose: oral		0.0086 μg/kg BW/day	
Background exposure		Not applicable (non-threshold substance)	
Dermal absorption factor		0.005	
Plant bioconcentration factor	Green	0.011	
	Root	0.011	
Tuber		0.001	
	Consumption-weighted mean	0.005	

 Table 19:
 Arsenic soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards			
	No produce	10% produce	25% produce	
Rural residential / lifestyle block	21	17 ^a	17 ^b	
Standard residential	24	20 ^a	17 ^b	
High-density residential	45			
Recreational	80			
Commercial / industrial indoor worker	NL			
Commercial / industrial outdoor worker / maintenance	70			

a Different rural residential and residential exposure durations result in different SCSs because non-threshold substance SCS derivation uses age-adjusted exposure rates.

b Derived values are less than 99th percentile of national dataset of background concentrations and therefore take the 99th percentile value. See Appendix 1 for values as derived.

NL = No limit.

The controlling pathway for arsenic is soil ingestion. Produce consumption has a significant influence for residential scenarios at high home-grown produce proportions but only a moderate influence at the standard residential proportion of 10 per cent.

The derived rural residential $SCS_{(health)}$ and site-specific SGVs derived for high home-grown produce percentages may be less than the local background concentration for arsenic. To avoid a SGV_(health) being below local background, the lowest value a SGV_(health) can take is, as a matter of policy, the 99th percentile concentration from a dataset collected from around the country from soil thought not to have been affected by anthropogenic activities. For arsenic this value is 17 mg/kg (see Appendix 6).

6.2 Boron

6.2.1 Bioconcentration factor for boron

The 'Timber Treatment Guidelines' (MfE and MoH, 1997) calculate SGVs using a BCF_{root} of 3 based on a measured range of 1–10 sourced from ECETOC (1990) and a BCF_{stem} of 0.6, the latter derived assuming stem concentrations are 20 per cent of root concentrations. No other regulatory agencies appear to consider boron and plant uptake of boron in the derivation of soil guideline values. The 'Timber Treatment Guidelines' also use a threshold for plant toxicity of boron of 3 mg/kg (as water-soluble boron) sourced from UK Interdepartmental Committee on the Redevelopment of Contaminated Land (ICRCL, exact reference not specified) as the basis for guideline values adopted for boron. These values are no longer considered appropriate.

It has not been possible to develop bioconcentration factors for boron. Reviewing the literature shows that boron uptake into plants is highly variable between species with no relationship with soil concentration or other soil parameters. Boron is an essential element for plant growth, but what may be optimal boron for one species may be toxic or insufficient for other species (Blevins and Lukaszewski, 1998; Nable, 1997). In addition, potential human health effects arising from ingestion of produce containing boron are generally considered to be protected by the soil-plant barrier, where toxicity to the plant will occur at concentrations far lower than what would affect human health (Chaney, 1980 in Langmuir et al, 2004). It is therefore appropriate that a maximal concentration of boron in produce is used in preference to a BCF.

Sensitive crop plants, considered to be cereals and cotton, may be affected by boron concentration in soil solution at 1 mg/L while 5 mg/L may be tolerated by various plant species, include most vegetables, and up to 15 mg/L by tolerant species (Blevins and Lukaszewski, 1998). Nable (1997) indicates that soil containing more than 5 to 8 mg/L hot water soluble boron may require special revegetation requirements.

Langmuir et al (2004) consider that 75 mg/kg represent phytotoxic levels in plants. However Nable (1997) indicates that in species that accumulate boron in their leaves, leaves can contain 250 mg/kg (dry weight) when boron in the soil approaches toxic levels and may exceed 1000 mg/kg (dry weight) in extreme conditions of boron toxicity. In species that do not accumulate boron in their leaves under conditions of toxicity, boron concentrations greater than 300 mg/kg (dry weight) may indicate that boron toxicity is present (Nable et al, 1997). In some species, concentrations of boron in plant tissue that hasn't resulted in toxicity may range up to 4800 mg/kg dry weight in corn (Kabata-Pendias and Pendias, 2000), although other studies using the same species indicate toxicity (yield reduction) at plant concentrations of 100 mg/kg.

Determining the significance of plant uptake of boron to human exposure is difficult, given the wide ranging and overlapping concentrations that determine boron essentiality and toxicity in various species. Nonetheless, it appears that 300 mg/kg is a reasonable upper limit of non-toxic plant boron concentrations and thus can be used as the reasonable maximum amount of boron likely to be taken up in home-grown vegetables. Beyond that point vegetables are unlikely to be harvestable. Alternatively, a hot water-soluble boron concentration of 8 mg/L could be considered as an upper limit of non-toxic concentrations for plants although the relationship between hot water-soluble boron and total boron concentrations is unclear, and likely to be highly variable.

6.2.2 Calculations for boron standard

The soil contaminant standard calculations for boron use the contaminant-specific parameters in table 20 and the derived $SCSs_{(health)}$ are set out in table 21. Boron has been treated as a threshold substance.

As noted above, BCF values could not be determined for boron and the $SCSs_{(health)}$ calculation has been carried out differently from the other guidelines with respect to the produce consumption pathway, because a produce pathway value cannot be determined as a function of soil concentration. This means that a soil guideline combining soil ingestion and the produce pathway cannot be calculated in the usual way. Instead, a soil ingestion guideline has been calculated for the residential-with-produce scenarios by subtracting a further notional background intake to take into account the amount of produce that could theoretically be consumed if the produce was at the phytotoxic limit of 300 mg/kg tissue concentration. The modified background is subtracted off the TDI in the usual fashion.

To obtain the additional background intake, a child's produce consumption (0.0105 kg DW/day) was multiplied by 300 mg/kg and divided by the child body weight of 13 kg to obtain the maximum additional background daily intake for 100 per cent of produce being home-grown. This was then multiplied by the home-grown produce percentage relevant to the particular SCS_(health).

As the dermal and inhalation intakes are insignificant there is no contribution from these pathways and the result with the modified background becomes the $SCS_{(health)}$ value.

This method breaks down for home-grown produce consumption percentages greater than about 49 per cent because the 'produce background' exceeds the TDI minus the normal background intake, resulting in nonsensical negative SCSs. For a site where greater home-grown produce consumption is a possibility, site-specific assessment will need to be carried out by measuring boron concentrations in relevant plants. This can be used to assess human exposure from plants in addition to soil.

Tolerable daily intake: oral	0.2 mg/kg BW/day		
Background exposure	Child		
	Adult	0.017 mg/kg BW/day	
Dermal absorption factor	0		
Plant bioconcentration factors	Root	Not applicable	
	Tuber	Not applicable	
	Leafy	Not applicable	
Additional child background for given produce	10%	0.024 mg/kg BW/day	
percentage at produce concentration of 300 mg/kg	25%	0.061 mg/kg BW/day	

Table 20: Contaminant-specific parameters for the derivation of the boron soil contaminant standard

Scenario	Combined soil contaminant standards				
	No produce	10% produce	25% produce		
Rural residential / lifestyle block	NL	NL	NL		
Standard residential	NL	NL	NL		
High-density residential	NL				
Recreational	NL				
Commercial / industrial indoor worker	NL				
Commercial / industrial outdoor worker / maintenance	NL				

Table 21: Boron soil contaminant standard (mg/kg)

Note: NL = No limit - the derived value exceeds 10,000 mg/kg, a concentration that is unlikely to be exceeded in practice. The derived values may be found in Appendix 1.

It should be noted that the derived SCSs values are all well above what would normally be encountered on contaminated sites in New Zealand, and have therefore been assigned a value of 'NL' for no limit. The derived values may be found in Appendix 1. Soil concentrations well below the derived concentrations may be above the phytotoxic threshold. If high boron concentrations are encountered on a site, the risk assessor will need to consider whether this could affect the use to which a site could be put. This is unlikely to be a consideration for most industrial or commercial uses.

6.3 Cadmium

6.3.1 Bioconcentration factors for cadmium

Cadmium uptake by plants is a function of cadmium concentration in the soil solution, although plant species and cultivars differ widely in their ability to absorb and accumulate cadmium (Kabata-Pendias and Pendias, 2000). Many researchers have found conflicting evidence for the relativity of uptake between different garden vegetables. Cadmium absorption has been shown to depend strongly on soil pH, and to a lesser degree on hydrous oxide and soil organic matter content (Alloway, 1995). Cadmium absorption (ie, plant uptake of cadmium) increases with decreasing pH. This means that in acid soils, produce consumption can represent a significant exposure route for cadmium (Swartjes et al, 2007).

Various regression equations have been developed for regulatory agencies to describe the plant uptake of cadmium in relation to various soil parameters. For example in the Netherlands, Otte et al (2001) and Swartjes et al (2007) used a regression equation describing plant concentrations in relation to four soil parameters: clay, organic carbon, total soil metal concentration and pH. Bechtel-Jacobs (1998) and McBride (2002) developed regression equations for plant uptake based on soil concentrations and pH; in the UK, Defra and EA (2002c) used regression equations for the behaviour of BCF in relation to soil pH for root and leafy vegetables.

Soil pH appears to be a dominant influence of plant uptake of cadmium, with greater uptake at lower pH. Use of the UK equation (Defra & EA, 2002c) in the derivation of soil guideline values in New Zealand is particularly attractive, as it uses only soil pH to describe the BCF. However, the original report describing this approach has now been withdrawn and a revised report on the derivation of soil guideline values for cadmium has recently been released which does not use the BCF-soil-pH relationship (EA, 2009b). Instead the geometric means of BCFs determined from the literature for the edible portion of six produce types are used. The applicability of the Dutch equations to New Zealand soils is debatable as it is based on data

from Dutch soils, and the Bechtel-Jacobs relationship was developed for all plant types, not just vegetables.

A review of the primary literature was undertaken and the relationships between plant uptake of cadmium and soil concentrations re-examined using collated data including those for New Zealand (MfE, 2010b). Specifically, only data from field studies was used, ie, studies where cadmium was already present in the soils (as opposed to experimental studies where solutions of cadmium salts are added to the soil: plant uptake of salts often overestimates uptake, particularly for cadmium – Effroysom et al, 2004). Further, only studies that reported soil pH, soil cadmium concentration and cadmium concentrations in the edible portions of plants were used. This resulted in 108 data points from 13 studies, with 51 data points for leafy vegetables and 33 for root and tuber vegetables. These data were used to examine the relationship between plant cadmium concentrations and soil pH. Based on the previous research described above, two relationships were examined using regression analyses:

$$Ln (BCF) = a + b (soil pH) Eqn 23$$

$$Ln (plant Cd) = a + b [ln (soil Cd)] + c (soil pH)$$
Eqn 24

Table 22 provides a summary of the equations developed and the percentage of the variability in the data explained (R^2).

Vegetable type	Ln (BCF) = a + b (soil pH)					
	а		b			R ²
All data	6.42		-0.991			23.7
Leafy	6.16		-0.844		23	
Root / tuber	7.04		-1.14		40.8	
Vegetable type	Ln (plant Cd) = a + b ln(soil Cd)+c pH)+c pH
	а	b		С		R ²
All data	5.462	0	.6981	-0.859		59.8
Leafy	5.66 0		.8783	-0.77	9	65.9
Root / tuber	4.61	0	.6215	-0.777		77.7

 Table 22:
 Coefficients determined for equations 23 and 24, and the percentage of the variability in the data

As a greater proportion of the data is explained using equation 24, it is considered this general form of equation is appropriate for determining vegetable uptake of cadmium. Further, the equations developed for the different vegetable types (leafy, root / tuber) were used as these equations have a better fit than those developed using all vegetable types.

Further analyses was undertaken using an extended data set and restricted to only leafy and root / tuber data (MfE, 2010b). The equations developed from this data are shown in table 23. The leafy and root / tuber values were used in the SCS derivation.

 Table 23:
 Coefficients determined for equation 24, and the percentage of the variability in the data explained using an extended data set

Vegetable type	Ln (plantCd) = a + b [ln(soilCd)] + c (soil pH)						
	а	R ²					
All data	4.98	0.728	-0.764	58.0			
Leafy	4.58	0.759	-0.626	68.1			
Root / tuber	4.73	0.600	-0.838	58.6			

The BCFs derived using the parameters shown in table 23 are plotted in figures 3 and 4 for leafy and root / tubers, respectively, for soil concentrations ranging from 0.1 to 4 mg/kg at soil pH 5, 5.5, 6 and 6.5. Studies on the properties of New Zealand soils indicate that soils under various land uses are acidic, with typical pH values for individual land types ranging from 5.2 to 6.2 (Sparling et al, 2000; Sparling and Schipper, 2002, 2004).

6.3.2 Calculations for cadmium standard

Cadmium has been treated as a threshold substance. The dependence of plant uptake on both soil pH and soil concentration requires an iterative derivation procedure for the calculation of the standard and rural residential SCSs. Different SGVs may be derived over a range of soil pH. However, as noted previously, New Zealand soils tend to be acidic and even where soil pH has been artificially raised (eg, by the addition of lime) the pH will revert to the natural value over time. The residential SCSs_(health) have therefore been calculated for a default pH of 5 and these values are to be applied in the first instance regardless of soil pH.

A SGV_(health) calculated for a higher pH may be used in a site-specific assessment if it can be demonstrated with sufficient certainty that some other soil pH is appropriate, with such an assessment including consideration of whether the pH is naturally high (eg, calcareous soils) or has merely been temporarily altered by artificial means (eg, liming). Further information is contained in appendix 2 on site-specific assessment for soil pH greater than 5.

The calculation procedure for a particular soil pH requires an estimated starting soil concentration to calculate the BCF. The resultant trial SGV was then compared with the starting concentration: if not within one per cent, a value midway between the starting value and the resultant value was fed back into the calculation (interval halving). This was carried out successively until convergence to a single value was achieved. This typically occurred in fewer than 10 iterations provided the starting value was chosen carefully.





Figure 4: Variation in plant bioconcentration factors (BCF, dry weight) for root and tuber vegetables with soil concentration and pH



The contaminant-specific parameters for the SCS calculations for cadmium are shown in table 24. The derived SCSs for pH 5 are set out in table 25. The detailed calculations from which the values in table 25 are derived are given in appendices 1 and 2, together with residential SGV calculations with pH values from pH 5 to pH 7, in 0.5 intervals. The BCF values are unique to each residential SGV and are not shown in table 24 but are given in appendix 2.

 Table 24:
 Contaminant-specific parameters for the derivation of the cadmium soil contaminant standard

Tolerable daily intake	: oral	0.000 83 mg/kg BW/day
Background exposure	Child Adult	0.000 41 mg/kg BW/day 0.000 26 mg/kg BW/day
Dermal absorption fac	ctor	0.001
Plant bioconcentratio	n factors	Depend on pH and soil concentration. See separate calculations, appendix 2

Table 25:	Cadmium	soil	contaminant	standard	(mg/kg)
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Scenario	Combined soil contaminant standards				
		No produce	10% produce	25% produce	
Rural residential / lifestyle block	pH 5	110	3	0.82	
Standard residential	pH 5	110	3	0.82	
High-density residential		230			
Recreational		400			
Commercial / industrial indoor worker		NL			
Commercial / industrial outdoor wo	orker	1,300			

Notes: NL = No limit.



Figure 5: Dependence of cadmium soil contaminant standard on pH

The soil ingestion rate has little or no influence on the SCSs derived for the standard and rural residential scenarios, with the produce consumption pathway being dominant. Where there is no produce consumption the derived SCSs are much larger. Dermal absorption is insignificant.

SGVs calculated for the site-specific situation of high home-grown produce consumption for acidic soils may be less than the background concentration of cadmium for the locality. The background concentration is typically in the range 0.02 to 0.5 mg/kg in New Zealand. To avoid a site-specific SGV_(health) being below local background, the lowest value a cadmium SGV_(health) can take is, as a matter of policy, the 99th percentile concentration from a dataset collected from around the country from soil thought not to have been affected by anthropogenic activities. For cadmium this value is 0.65 mg/kg (see appendix 6, section A6.3.2).

6.4 Chromium

6.4.1 Bioconcentration factor for chromium

Concentrations of chromium in plant-available form are extremely low in most soils, and this is reflected in low concentrations of the element in plants (McGrath, 1995). Anthropogenic sources of chromium are believed to be responsible for elevated concentrations of this metal in plants, although overall there have been few studies of its uptake and accumulation (Kabata-Pendias and Pendias, 2000).

Chromium typically occurs in two valence states, chromium III and chromium VI; the former is the only state normally found in aerobic soils. In theory, chromium VI is likely to be more available to plants for uptake (because of its greater mobility in soils), but there have been few studies to support this (Kabata-Pendias and Pendias, 2000).

The 'Timber Treatment Guidelines' (MfE and MoH, 1997) uses BCFs of 0.015 for root and 0.003 for leafy vegetables for chromium III (table 26) based on the assumption that bioaccumulation in vegetative parts is one-fifth of that in the roots. For chromium VI a BCF of 0.24 for root vegetables is used, with a BCF of 0.048 for leafy vegetables also being one-fifth of the root value. These values are nominally sourced from ECOTOC (1990).

The US EPA (2007) reports a median BCF of 0.041, determined from 28 data points with a range of 0.021 to 0.48 (Bechtel-Jacobs 1998). Defra and EA (2002d) use BCFs of 0.02 and 0.055 for root and leafy vegetables, respectively, based on a literature review of the uptake of total and chromium III.

A summary of international values is shown in table 26. Given that there is little evidence for preferential uptake of chromium VI, and the lack of information on the source of data provided in the 'Timber Treatment Guidelines' (MfE and MoH,1997), it is proposed that the BCF for chromium is based on a simple average¹⁰ of all values shown in bold in table 26. This gives a BCF of 0.0324, and is considered to be applicable to all vegetables.

Substance	MfE ar (1997, T	MfE and MoH (1997, Table A1)		et al, 2001)	US EPA (2007)	Defra a (200	and EA)2d)	Mean
	Root	Leafy ^a	CMp	Median		Root	Leafy	
Cr III	0.015	0.003						
Cr VI	0.24	0.048						
Cr			0.011	0.0135	0.041	0.02	0.055	0.0324

 Table 26:
 Summary of BCFs for chromium from different sources

a Assuming leafy five times less than root.

b CW = consumption weighted BCF, calculated using plant-soil relationships and produce consumption data.

6.4.2 Calculations for Chromium Standard

Soil contaminant standards have been calculated for both, chromium III and chromium VI as threshold substances. The SCS calculations for chromium use the contaminant-specific parameters in table 27 and the derived SCSs are set out in tables 28 and 29 for chromium III and VI, respectively.

A single BCF is used, applicable to both root/tuber and leafy parts of vegetables. The same background intake and BCF were used for both valance states. There is no data for the background intake of chromium VI. In accordance with MfE (2011) the background intake is taken as five per cent of the TDI.

¹⁰ Produce-type consumption-weighted mean BCFs have generally been used in this document, however, this refinement was not considered appropriate for the available data.

Table 27: Contaminant-specific parameters for the derivation of the chromium soil contaminant standard

		Chromium III	Chromium VI
Tolerable daily intake: oral		1.5 mg/kg BW/day	0.003 mg/kg BW/day
Background exposure	Child	0.0012 mg/kg BW/day	No data
	Adult	0.000 53 mg/kg BW/day	No data
Dermal absorption factor		0	
Plant bioconcentration factor	Leafy	0.0324	
	Root	0.0324	
	Tuber	0.0324	

Table 28: Chromium III soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards			
	No produce	10% produce	25% produce	
Rural residential / lifestyle block	NL	NL	NL	
Standard residential	NL	NL	NL	
High-density residential	NL			
Recreational	NL			
Commercial / industrial indoor worker	NL			
Commercial / industrial outdoor worker / maintenance	NL			

Note: NL = No limit - the derived value exceeds 10,000 mg/kg, a concentration that is unlikely to be exceeded in practice. The derived values may be found in appendix 1.

Table 29: Chromium VI soil contaminant standard (mg/kg)

Scenario	Combined	soil contaminant s	tandards
	No produce	10% produce	25% produce
Rural residential / lifestyle block	770	460	290
Standard residential	770	460	290
High-density residential	1,500		
Recreational	2,700		
Commercial / industrial indoor worker	NL		
Commercial / industrial outdoor worker / maintenance	6,300		

Note: NL = No limit.

It is notable that the calculated values for chromium III are all very high, some in excess of pure chromium, which is impossible. The results indicate no practical limit to allowable soil concentrations and have therefore been assigned 'NL' for no limit. However, acute effects for a pica child (consuming 5 to 10 grams of soil in a day) have not been checked. This should be assessed on a site-specific basis in the unlikely event that very high concentrations of chromium and a child with the pica habit are present. In practical terms, a concentration of chromium III of greater than 10,000 mg/kg is not likely.

Apart from possibly acute effects for children, phytotoxicity will occur well before the indicated soil concentrations are reached. This suggests that the calculated intake from produce consumption will not occur because produce will not reach a harvestable condition.

Chromium in soil is commonly analysed as total recoverable chromium, which will not differentiate between the trivalaent and hexavalent forms of chromium. For most situations where chromium is present, it will be in only the trivalent form, however, it is appropriate to conservatively compare the results of total recoverable analyses with the chromium VI SCS in the first instance. If the SCS is exceeded, then analysis for chromium VI should be carried out to determine whether chromium VI is present.

6.5 Copper

6.5.1 Bioconcentration factor for copper

The 'Timber Treatment Guidelines' (MfE and MoH, 1997) use a BCF for copper of 0.28 for roots, and assume that the BCF for leafy vegetables is 20 per cent of this. However, as discussed in Cavanagh (2004b), the equation used to estimate BCF is incorrectly stated to be root vegetables, as opposed to above-ground vegetables. This means that BCF_{leafy} should be 0.28, and BCF_{root} should be 1.4.

The US EPA (2007) reports a median bioconcentration factor of 0.124 from 180 data points and a range of 0.0011 to 7.4. Baes et al (1984) have shown that plant uptake of copper is dependent on the copper soil concentration. Copper is phytotoxic at relatively low concentrations, and plant uptake of copper is limited by its toxic effect on plants. A tissue copper concentration of 15–20 mg/kg (dry weight) is considered to be representative of excessive tissue concentration in agronomic species, while a 10 per cent growth yield decrease is most likely at 10–30 mg/kg (dry weight) tissue copper concentrations (Kabata-Pendias and Pendias, 2000).

Given the variable uptake of copper by plants from soil, and the known phytotoxic effects of copper, it is recommended that a maximal concentration of copper in produce is used in preference to a BCF. Based on the discussion above it is further recommended that a produce concentration of 30 mg/kg (dry weight) is used as the maximum amount of copper likely to be taken up in home-grown vegetables. Vegetables containing greater than this concentration would be so stunted and deformed that harvesting would be unlikely.

6.5.1 Calculations for copper standard

The soil contaminant standard calculations for copper use the contaminant-specific parameters in table 30 and the derived SCSs are set out in table 31. Copper has been treated as a threshold substance.

A BCF was not used to calculate the SCS for scenarios with produce consumption. As noted above, vegetables are unlikely to be harvested with tissue concentrations greater than 30 mg/kg dry weight. Any concentrations well in excess of this would theoretically accumulate in the plants at soil concentrations that are safe for the soil ingestion pathway, so a produce pathway soil guideline value cannot be determined as a function of soil concentration. Instead, a soil ingestion guideline has been calculated for the residential-with-produce scenarios by subtracting a further notional background intake to take into account the amount of produce that could theoretically be consumed if the produce was at the phytotoxic limit of 30 mg/kg tissue concentration. The modified background is subtracted from the TDI in the usual fashion.

To obtain the additional background intake, a child's produce consumption (0.0105 kg DW/day) was multiplied by 30 mg/kg and divided by the child's body weight of 13 kg to obtain the maximum additional background daily intake for 100 per cent of produce being home-grown. This was then multiplied by the homegrown produce percentage relevant to the particular SCS.

As the dermal and inhalation intakes are insignificant there is no contribution from these pathways and the result with the modified background becomes the SCS.

Table 30:	Contaminant-specific parameters for the derivation of the copper soil
	contaminant standard

Tolerable daily intake: oral	0.15 mg/kg BW/day	
Background exposure	Child	
	Adult	0.02 mg/kg BW/day
Dermal absorption factor	0	
Plant bioconcentration factors	Root	na
	Tuber	na
	Leafy	na
Additional child background for given produce	10%	0.0024 mg/kg BW/day
percentage at produce concentration of 30 mg/kg	25%	0.0061 mg/kg BW/day

na = not applicable.

Table 31: Copper soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards				
	No produce	10% produce	25% produce		
Rural residential / lifestyle block	NL	NL	NL		
Standard residential	NL	NL	NL		
High-density residential	NL				
Recreational	NL				
Commercial / industrial indoor worker	NL				
Commercial / industrial outdoor worker / maintenance	NL				

NL = No limit - the derived value exceeds 10,000 mg/kg, a concentration that is unlikely to be exceeded in practice. The derived values may be found in appendix 1.

It should be noted that the derived SCSs values are all well above what would normally be encountered on contaminated sites in New Zealand, and have therefore been assigned a value of 'NL' for no limit. The derived values may be found in appendix 1. Soil concentrations well below the derived concentrations may be above the phytotoxic threshold. If high copper concentrations are encountered on a site, the risk assessor will need to consider whether this could affect the use to which a site could be put. This is unlikely to be a consideration for most industrial or commercial uses.

6.6 **Inorganic lead**

Bioconcentration factor for lead 6.6.1

Inorganic lead is generally considered to be relatively immobile in soil and has limited plant uptake. Lead BCFs for most plants typically range from 0.001 to 0.03 (Jones and Johnston, 1991). Various regulatory agencies have examined plant uptake of lead for use in deriving soil guideline values. The US EPA (2007) used a relationship describing plant foliage lead concentrations as a function of soil lead concentrations derived by Bechtel Jacobs (1998) in the derivation of ecological soil screening levels (Eco-SSLs).

The Ontario Ministry for the Environment and Energy (OMEE) used a similar BCF of 0.039 (dry weight) determined from a study of the uptake of lead from Canadian soils by common backyard vegetables (OMEE, 1994). This is mentioned in CCME environmental guidelines on lead in the section on derivation of environmental guidelines, but does not appear to be used in the derivation of guidelines for the protection of human health. The Dutch use plant concentrations as a function of soil properties (Otte et al, 2001)

As there have been no recent summaries of the plant uptake of lead, a literature search was undertaken and data compiled to determine the geometric mean for leafy, root (carrot and radish) and tuber (potato) vegetables (MfE, 2010b). Seventy-eight data points from 10 studies were found. These derived geometric means are recommended for use in deriving soil guideline values in New Zealand and are shown in table 32, along with BCFs determined by other authors for a similar purpose.

Table 32: Recommended BCF (dry weight) and BCF (dry weight) for lead from different sources

Recomm		Dutch (Otte et al, 2001)		Bechtel-Jacobs (1998)	OMEE (1994)	
Leafy / above ground (n=41)	Root (n=13)	Tuber (n=4)	CW ^a	Median ^b	Geometric mean	
0.019	0.015	0.005	0.017	0.015	0.038	0.039

a CW = consumption weighted BCF, calculated using plant-soil relationships and produce consumption data.

Median of all measured data for vegetables.

6.6.2 Calculations for lead standard

Soil contaminant standards have been calculated for lead as a threshold substance. The calculations use the contaminant-specific parameters in table 33 and the derived SCSs are set out in table 34.

Three BCF values were used, one each for leafy, root and tuber vegetables, combined into a single consumption-weighted mean value, using weighting factors of 0.3, 0.1 and 0.6 (dry weight), respectively.

Table 33:	Contaminant-specific parameters for the derivation of the lead soil
	contaminant standard

Tolerable daily intake: oral	0.001 90 mg/kg BW/day		
Background exposure	Child	0.000 97 mg/kg BW/day 0.000 41 mg/kg BW/day	
	Adult		
Dermal absorption factor		0	
Plant bioconcentration	Leafy	0.019	
factors	Root	0.015	
	Tuber	0.005	
	Consumption weighted mean	0.0102	

Table 34: Lead soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards			
	No produce	10% produce	25% produce	
Rural residential / lifestyle block	250	210	160	
Standard residential	250	210	160	
High-density residential	500		L	
Recreational	880			
Commercial / industrial indoor worker	NL			
Commercial / industrial outdoor worker / maintenance	3,300			

NL = No limit.

The derived SCSs for lead are dominated by the ingestion pathway. The dermal pathway has no influence. For the residential with produce scenarios, the produce pathway has a significant influence.

The derived SCSs are not applicable to organic lead compounds.

6.7 Inorganic mercury

6.7.1 Mercury bioconcentration factor

There is limited data available on mercury uptake into home produce. Three sources have examined plant uptake of mercury for the purpose of developing regulatory guideline values. Bechtel-Jacobs (1998) provide a summary of plant uptake data from a range of plants including vegetables. Using regression analyses, this data was used to develop relationships between concentrations in plants and soil that are used in the derivation of US ecological soil screening levels (Eco-SSLs (US EPA, 2007 (in attachment 4.1). The geometric mean and median values provided in Bechtel-Jacobs (1998) are shown in table 35.

Versluijs and Otte (2001) developed a series of equations relating mercury (and other metal) concentrations in the edible parts of various vegetables to soil mercury concentrations and soil parameters using multiple regression. Using a standard soil, these equations are used to determine a consumption-weighted bioconcentration factor that is used to develop Dutch intervention values. This value is reported in Lijzen et al (2001) and Otte et al (2001); the latter also provides a summary of the actual plant BCFs from the collected data. The consumption-weighted BCF is shown in table 35, along with the BCF used for calculating the current Dutch mercury intervention value.

The Environment Agency (EA, 2009c) provides the most recent and relevant summary of plant BCFs. This report reviewed the existing literature and provides recommendations for BCFs (based on fresh weight) for the edible portion of six produce types, which are used in the CLEA model for the derivation of soil guideline values. The recommended BCFs are the geometric mean of the available data for each produce type.

The BCFs recommended by Environment Agency (EA, 2009c) are lower than provided by Bechtel-Jacobs (1998) and used in Otte et al, 2001. Close inspection of the Bechtel-Jacobs (1998) reveals a high proportion of non-vegetables that have significantly higher BCFs than vegetables provided in the same dataset. The vegetable BCFs in the Bechtel-Jacobs data set are similar to that provided by the Environment Agency (EA, 2009c), and also similar to that previously used in the derivation of Dutch intervention values. In contrast, the revised BCF determined from derived soil-plant relationships is higher than that determined from the literature. As the review undertaken by EA (2009c) is recent and for the purpose of deriving soil guideline values for the protection of human health, these values are recommended for the derivation of New Zealand soil guideline values. Conversion of the uptake factors to dry weight has been undertaken using the dry weight conversion factors specified in EA (2009a).

Dutch (Otte et al, 2001)		Bechtel-Jacobs (1998)	UK (EA 2009c) Fresh weight (dry weight) ^c		Recon	nmended	values		
CW ^a	GM⁵	Previous	GM	Green (n=52)	Root (n=52)	Tuber (n=13)	Green (n=52)	Root (n=52)	Tuber (n=13)
0.15	0.28	0.02	0.35	0.0038 (0.04)	0.0069 (0.07)	0.0043 (0.02)	0.04	0.07	0.02

 Table 35:
 BCF for mercury from different sources

a CW = consumption weighted BCF, calculated using plant-soil relationships and produce consumption data.

b Geometric mean of all measured data for vegetables.

c Dry weight (DW) calculated using conversion factors of 0.096 kg DW/g FW, 0.103 kg DW/kg fresh weight (FW) and 0.21 kg DW/kg FW for green, root and tuber vegetables respectively (EA, 2009a).

6.7.1 Calculations for mercury standard

Soil contaminant standards have been calculated for inorganic mercury as a threshold substance. The calculations use the contaminant-specific parameters in table 36 and the derived SCSs are set out in table 37.

Three BCF values were used: one each for leafy, root and tuber vegetables combined into a single consumption-weighted mean value, using weighting factors of 0.3, 0.1 and 0.6 (dry weight), respectively.

Table 36:	Contaminant-specific parameters for the derivation of the inorganic mercury
	soil contaminant standard

Tolerable daily intake: oral		0.002 mg/kg BW/day	
Background exposure Child		0.00005 mg/kg BW/day	
	Adult	0.000065 mg/kg BW/day	
Dermal absorption factor		0	
Plant bioconcentration	Leafy	0.04	
factors	Root	0.07	
	Tuber	0.02	
	Consumption weighted mean	0.031	

Scenario	Combined soil contaminant standards		
	No produce	10% produce	25% produce
Rural residential / lifestyle block	510	310	200
Standard residential	510	310	200
High-density residential	1,000		
Recreational	1,800		
Commercial / industrial indoor worker	NL		
Commercial / industrial outdoor worker / maintenance	4,200		

Table 37: Inorganic mercury soil contaminant standard (mg/kg)

Note: NL = No limit.

Produce consumption has a significant influence on the SCSs for inorganic mercury for the residential-with-produce scenarios. For scenarios without produce consumption, soil ingestion is the dominant pathway. Dermal absorption is insignificant.

The inorganic mercury SCS is not intended to be applied to a site contaminated with elemental mercury or organic mercury compounds (eg, methyl mercury).

6.8 Benzo(a)pyrene (BaP)

The SCS for benzo(a)pyrene (BaP) is intended to represent the several polycyclic aromatic hydrocarbons (PAHs) thought to be carcinogenic (MfE, 2011). PAHs typically occur as complex mixtures in which one or more carcinogenics may be present. The application of the SCS, using potency equivalency factors (PEFs) to calculate a BaP equivalence, is explained at the end of this subsection.

6.8.1 BaP bioconcentration factor

Limited data is available on plant uptake of BaP and a number of sources have used various models to determine plant uptake. For example, in New Zealand, the 'Oil Industry Guidelines' (MfE, 1999) use the method of Ryan et al (1988), whereas the 'Gasworks Guidelines' (MfE, 1997) use the method of Travis and Arms (1988). Both methods are based on the log K_{ow} of the organic contaminant, and slightly different values are used in the two guideline documents – the 'Oil Industry Guidelines' use a log $K_{ow} = 6.04$ whereas the 'Gasworks Guidelines' use $K_{ow} = 9.55 \times 10^5$, which gives a log $K_{ow} = 5.98$. The Travis and Arms (1988) relationship determines a plant bioaccumulation factor on a dry-weight basis in above-ground parts. In the *Gasworks Guidelines*, this is converted to a fresh-weight plant uptake factor (PUF) using the assumption that dry weight is 20 per cent of fresh weight for all vegetables. It should be noted that the US EPA (2003) have criticised the Travis and Arms relationship as being based on few data, some of which are at variance with the source documents cited by Travis and Arms.

The Ryan et al (1988) model determines fresh-weight PUF for roots and leaves directly from concentrations in the pore water. However, the 'Oil Industry Guidelines' do not indicate how the pore water concentrations relate to soil concentrations provided in the document.

Dutch authors use the model of Briggs et al (1982, 1983; in Lijzen et al, 2001) to estimate uptake of organic contaminants in leafy vegetables and the model of Trapp and Matthies (1995, in Lijzen et al, 2001) with modified parameters to estimate uptake of organic contaminants by root vegetables.

In contrast, the US EPA (2007) reports a median bioaccumulation factor (BAF) for plant foliage of 0.1 from 15 data points with a range of 0.039–0.2; it uses a relationship describing plant concentration as a function of soil concentration based on measured data to derive ecological soil screening levels (Eco-SSLs).

The preferred approach in the current work is to use BCFs determined from measured data. As such, literature data on plant uptake of BaP was compiled and used to determine the BCF for leafy, root and tuber vegetables (MfE, 2010b). Root and tuber vegetables were kept separate as the available data indicates a difference in uptake between the two vegetable types. It should be noted that uptake of vapour-phase BaP (or any PAH) is primarily from ambient air as opposed to volatilisation from soil. This is often implicated as being the most significant pathway of plant uptake leading to accumulation in all plant parts, even in plant roots (eg, Kipopoulou et al, 1999; Wild et al, 1992). This suggests that BCFs, particularly for leafy vegetables, are likely to be overestimated.

6.8.2 Calculations for BaP standard

The soil contaminant standard calculations for BaP use the contaminant-specific parameters in table 38 and the derived SCSs are set out in table 39. BaP has been treated as a non-threshold substance and therefore a risk-specific dose has been employed. Three BCF values were used: one each for leafy, root and tuber vegetables, combined into a single consumption-weighted mean value` using weighting factors of 0.3, 0.1 and 0.6 (dry weight), respectively.

MfE (2011) provides two skin absorption factors, a 'worst case' of 0.06 and 0.026 for 'aged' contamination. The worst-case value has been used for the SCS derivation, although as dermal absorption has little influence on the SCS, the difference between the two values is small.

Risk-specific dose: oral		0.0048 μg/kg BW/day	
Background exposure		Not applicable (non-threshold substance)	
Dermal absorption factor		0.06	
Plant bioconcentration	Leafy (n=10)	0.005	
factors	Root (n=17)	0.031	
	Tuber (n=3)	0.004	
	Consumption weighted mean	0.007	

 Table 38:
 Contaminant-specific parameters for the derivation of the BaP soil contaminant standard

Table 39: BaP soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards		
	No produce	10% produce	25% produce
Rural residential / lifestyle block	11	8 ^a	6
Standard residential	12	10 ^a	7
High-density residential	24		
Recreational	40		
Commercial / industrial indoor worker	NL		
Commercial / industrial outdoor worker / maintenance	35		

a Different rural residential and residential exposure durations result in different SCSs because non-threshold substance SSV derivation uses age-adjusted exposure rates.

NL = No limit.

The controlling pathway for BaP is soil ingestion but produce ingestion also has an influence on the derived SCS for residential scenarios, even at the default produce proportion of 10 per cent. Dermal absorption has minimal influence.

As noted earlier, BaP is used to represent the carcinogenic PAHs. To enable an estimate of the potential carcinogenicity of polycyclic aromatic hydrocarbon mixtures, potency equivalence factors have been used previously in New Zealand guidance (MfE, 1997, 1999), and are now employed for the SCS to calculate a BaP equivalence concentration (BaP_{eq}).¹¹ As recommended in MfE (2010b), the PEFs developed by Kalberlah et al (1995 cited in WHO, 1998) are to be used. The PEFs are given in table 40. The PEFs cover a wider range than used in current New Zealand guidance documents (MfE, 1997, 1999).¹²

0.01

0.1

Polycyclic aromatic hydrocarbon	Potency equivalency factors
Benz(a)anthracene	0.1
Benzo(b)fluoranthene	0.1
Benzo(j)fluoranthene	0.1
Benzo(k)fluoranthene	0.1
Benzo(a)pyrene	1.0
Chrysene	0.01
Dibenz(a,h)anthracene	1.0

Table 40: PEFs for use in assessing potential carcinogenicity of PAH mixtures

Source: WHO, 1998.

Indeno(1,2,3-c,d)pyrene

Fluoranthene

¹¹ The equivalent BaP concentration is calculated as the sum of each of the detected concentrations of nine carcinogenic PAHs (benz(a)anthracene, benzo(b)fluoranthene, benzo(j)fluoranthene, benzo(k)fluoranthene, benzo(a)pyrene, chrysene, dibenz(a,h)anthracene, fluoranthene and indeno(1,2,3-cd) pyrene), multiplied by their respective potency equivalency factors (table 40).

¹² FAO/WHO (2006) identified 13 PAHs as carcinogenic, of which eight, together with fluoranthene, are included in table 40. Consideration is being given to expanding table 40 to include PEFs for dibenzo(a,e)pyrene, dibenzo(a,h)pyrene, dibenzo(a,i)pyrene and dibenzo(a,l)pyrene.

6.9 DDT

The SCS for DDT has been derived to represent DDT and its metabolites DDD and DDE. The sum of the six isomers¹³ is commonly referred to as $\sum DDT$.

6.9.1 Bioconcentration factor for DDT

Limited data on the plant uptake of DDT are available, although two recent studies provide some data on the uptake of DDT and its primary degradation products, DDE, in vegetables from historically contaminated soils (Gaw et al, 2008; Mikes et al, 2009). Gaw et al (2008) examined uptake into lettuce and radish, whereas Mikes et al (2009) examined uptake into radishes. Using data from the edible portions of the vegetables (lettuce leaf, radish root) from these studies, the geometric mean for the BCFs for p,p'-DDT and p,p'-DDE in the two vegetable types are shown in table 41 (MfE, 2010b). These values compare with a median plant uptake factors for grasses of 0.136 (n=3) for DDE and 0.037 (n=6) for DDT in US EPA (2007).

A plant uptake factor for DDE as opposed to DDT is more appropriate, as DDE is the primary metabolite of DDT and is the compound most commonly found in highest concentrations in contaminated soils.

 Table 41:
 BCF (geometric mean) for *p,p*'-DDT and *p,p*'-DDE in root and leafy vegetables determined from the literature

Parameter	Root		Lea	afy
	BCF n		BCF	n
<i>p,p⁻</i> DDE	0.038	10	0.012	9
p,p'-DDT	0.022	9	0.003	7

6.9.2 Calculations for DDT standard

The soil contaminant standard calculations for DDT use the contaminant-specific parameters in table 42 and the derived SCSs are set out in table 43. DDT has been treated as a threshold substance. Two BCF values were used for root / tuber and leafy parts of vegetables, combined into a single consumption-weighted mean value, using weighting factors of 0.7 and 0.3 (dry weight), respectively.

 Table 42:
 Contaminant-specific parameters for the derivation of the DDT soil contaminant standard

Tolerable daily intake: oral		0.0005 mg/kg BW/day
Background exposure Child		0.000 051 1 mg/kg BW/day
	Adult	0.000 019 3 mg/kg BW/day
Dermal absorption factor		0.018
Plant bioconcentration	Leafy	0.012
factors	Root / tuber	0.038
	Consumption weighted mean	0.030

¹³ The six isomers are *p*,*p*'-DDT, *o*,*p*'-DDT, *p*,*p*'-DDE, *o*,*p*'-DDD, *o*,*p*'-DDD.

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Table 43: DDT soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards		
	No produce	10% produce	25% produce
Rural residential / lifestyle block	120	70	45
Standard residential	120	70	45
High-density residential	240		
Recreational	400		
Commercial / industrial indoor worker	NL		
Commercial / industrial outdoor worker / maintenance	1,000		

Note: NL = No limit.

Produce consumption has a significant influence on the SCS for DDT for the residential-withproduce scenarios. For scenarios without produce consumption, soil ingestion is the dominant pathway. Dermal absorption has only a minor influence on the combined value.

As noted early, the SCS is for the sum of DDT, DDD and DDE. The value is compared with the sum of the concentrations from the laboratory analysis of all six isomers of these compounds.

6.10 Dieldrin

6.10.1 Bioconcentration factor for dieldrin

Limited data on the plant uptake of dieldrin are available. A plant uptake factor of 0.41 was used for dieldrin, based on the median value for nine observations provided in US EPA (2007). These data were determined for the above-ground portions of three plants; in the absence of any further information, this value is considered to be applicable to all vegetable parts.

6.10.2 Calculations for dieldrin standard

The soil contaminant standard calculations for dieldrin assume it is a threshold substance. The contaminant-specific parameters used in the calculations are in table 44 and the derived SCSs are set out in table 45. A single BCF value was used for root, tuber and leafy parts of vegetables.

Table 44: Contaminant-specific parameters for the derivation of the dieldrin soil contaminant standard

Tolerable daily intake: oral		0.0001 mg/kg BW/day
Background exposure Child Adult		0.000 003 6 mg/kg BW/day 0.000 001 4 mg/kg BW/day
Dermal absorption factor		0.1
Plant bioconcentration factors	Leafy Root / tuber Consumption weighted mean	0.41 0.41 0.41

Scenario	Combined soil contaminant standards		
	No produce	10% produce	25% produce
Rural residential / lifestyle block	22	2.6	1.1
Standard residential	22	2.6	1.1
High-density residential	45		
Recreational	70		
Commercial / industrial indoor worker	NL		
Commercial / industrial outdoor worker / maintenance	160		

Table 45: Dieldrin soil contaminant standard (mg/kg)

Note: NL = No limit.

Produce consumption has a dominant influence in the SCSs for dieldrin for the residential-withproduce scenarios. For scenarios without produce consumption soil ingestion is the dominant pathway, although dermal absorption has some influence with dieldrin.

6.10.3 Applicability of the dieldrin SCS to aldrin

Aldrin was last used in New Zealand almost 50 years ago (MFE, 2006a). As aldrin degrades to dieldrin in the environment, with reported half lives in soil of 20–100 days (FAO, 2000), only small amounts of aldrin are expected to be detected, and most likely in conjunction with dieldrin, which would be at higher concentrations.

As the TDI for dieldrin is also applicable to aldrin (MfE, 2010b) the SCS is applicable to dieldrin or aldrin separately, or to the sum of aldrin and dieldrin if both are involved.

6.11 Dioxin and dioxin-like PCBs

The term 'dioxins' encompasses a group of 75 polychlorinated dibenzo-*p*-dioxin (PCDD) and 135 polychlorinated dibenzofuran (PCDF) congeners. PCDDs and PCDFs are formed during incomplete combustion processes. They also occur as contaminants during various industrial processes, eg, the chemical manufacture of some chlorinated compounds and chlorine bleaching of paper pulp.

The toxicity of individual dioxin congeners differs considerably. The congeners that are toxicologically important have chlorine atoms substituted in each of the 2-, 3-, 7- and 8-positions. Thus, from 210 theoretically possible congeners, only 17 are of toxicological concern. These compounds have a similar toxicological profile to that of the most toxic congener, 2,3,7,8-tetrachlorodibenzo-*p*-dioxin (2,3,7,8-TCDD).

Twelve of the 209 possible polychlorinated biphenyl (PCB) congeners also exhibit 'dioxin-like' toxicity. This document only considers dioxin-like PCB congeners and does not consider other PCB congeners. The toxicity of other PCB congeners will need to be considered separately in any assessment where PCBs are of concern.

Dioxins and PCBs do not occur as pure compounds but as complex mixtures of many congeners. When considering the toxicity of mixtures, the total toxicity is assessed as a toxic equivalency (TEQ) to 2,3,7,8-TCDD using toxic equivalency factors (TEF). The TEQ is defined as the sum of the products of the concentration of each compound multiplied by the value of its TEF; it is an estimate of the total 2,3,7,8-TCDD-like activity of the mixture.

A number of TEF schemes have been developed, the most recent from the 2005 World Health Organization re-evaluation by Van den Berg et al (2006). These TEFs are given in table 46 and are the values recommended for calculating the TEQ to be compared with dioxin and dioxin-like PCB SCSs.

Compound	Abbreviation	WHO (2005)
Polychlorinated dibenzodioxins		
2,3,7,8-Tetrachlorodibenzodioxin	TCDD	1
1,2,3,7,8-Pentachlorodibenzodioxin	1,2,3,7,8-PeCDD	1
1,2,3,4,7,8-Hexachlorodibenzodioxin	1,2,3,4,7,8-HxCDD	0.1
1,2,3,6,7,8-Hexachlorodibenzodioxin	1,2,3,6,7,8-HxCDD	0.1
1,2,3,6,7,9-Hexachlorodibenzodioxin	1,2,3,6,7,9-HxCDD	0.1
1,2,3,4,6,7,8-Heptachlorodibenzodioxin	1,2,3,4,6,7,8-HpCDD	0.01
Octachlorodibenzodioxin	OCDD	0.0003
Polychlorinated dibenzofurans		
2,3,7,8-Tetrachlorodibenzofuran	2,3,7,8-TCDF	0.1
1,2,3,7,8-Pentachlorodibenzofuran	1,2,3,7,8-PeCDF	0.03
2,3,4,7,8-Pentachlorodibenzofuran	2,3,4,7,8-PeCDF	0.3
1,2,3,4,7,8-Hexachlorodibenzofuran	1,2,3,4,7,8-HxCDF	0.1
1,2,3,6,7,8-Hexachlorodibenzofuran	1,2,3,6,7,8-HxCDF	0.1
1,2,3,7,8,9-Hexachlorodibenzofuran	1,2,3,7,8,9-HxCDF	0.1
2,3,4,6,7,8-Hexachlorodibenzofuran	2,3,4,6,7,8-HxCDF	0.1
1,2,3,4,6,7,8-Heptachlorodibenzofuran	1,2,3,4,6,7,8-HpCDF	0.01
1,2,3,4,7,8,9-Heptachlorodibenzofuran	1,2,3,4,7,8,9-HpCDF	0.01
Octochlorodibenzofuran	OCDF	0.0003
'Non-ortho' polychlorinated biphenyls		
3´,4,4´-Tetrachlorobiphenyl (PCB 77)	3,3´,4,4´-TCB	0.0001
3,4,4´,5,-Tetrachlorobiphenyl (PCB 81)	3,4,4´,5-TCB	0.0003
3,3´,4,4´,5-Pentachlorobiphenyl (PCB 126)	3,3',4,4',5-PeCB	0.1
3,3´,4,4´,5,5´-Hexachlorobiphenyl (PCB 169)	3,3´,4,4´,5,5´-HxCB	0.03
'Mono-ortho' polychlorinated biphenyls		
2,3,3´,4,4´-Pentachlorobiphenyl (PCB 105)	2,3,3´,4,4´-PeCB	0.00003
2,3,4,4´,5-Pentachlorobiphenyl (PCB 114)	2,3,4,4´,5-PeCB	0.00003
2,3´,4,4´,5-Pentachlorobiphenyl (PCB 118)	2,3',4,4',5-PeCB	0.00003
2,3´,4,4´,5'-Pentachlorobiphenyl (PCB 123)	2,3´,4,4´,5´-PeCB	0.00003
2,3,3',4,4',5-Hexachlorobiphenyl (PCB 156)	2,3,3´,4,4´,5-HxCB	0.00003
2,3,3´,4,4´,5´-Hexachlorobiphenyl (PCB 157)	2,3,3´,4,4´,5´-HxCB	0.00003
2,3´,4,4´,5,5´-Hexachlorobiphenyl (PCB 167)	2,3´,4,4´,5,5´-HxCB	0.00003
2,3,3´,4,4´,5,5´-Heptachlorobiphenyl (PCB 189)	2,3,3´,4,4´,5,5´-HpCB	0.00003

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6.11.1 Bioconcentration factors for dioxin and dioxin-like PCB

Plant uptake of dioxins and dioxin-like PCBs is suggested to primarily occur through atmospheric deposition, eg, uptake of a gaseous fraction or particle-bound contaminants via the leaves (Hulster and Marschner, 1993; McLachlan, 1997; Meneses et al, 2002; Collins et al, 2006). Root-uptake / translocation of PCDD/PCDFs from the soil, or volatilisation of PCDD/ CDFs from the soil and their subsequent adsorption onto the vegetation is negligible (McLachlan, 1997 and references contained therein; Jones and Duarte-Davidson, 1997).

Plant uptake via soil for dioxins and dioxin-like PCBs is only considered for the cucurbits, particularly those in the genus *Cucurbita* (zucchini and pumpkin): these are the only plants for which uptake via soil has been conclusively demonstrated (Hulster et al, 1994; Inui et al, 2008). This is suggested to be due to the production of root exudates in these species, although differences in uptake also exist between different cultivars (Inui et al, 2008). For other vegetables, uptake from soil is not considered a relevant exposure pathway.

Transfer of dioxin-like PCBs is indicated to be greater than that of PCDD/PCDFs, with pentaand hexa-chloro-biphenyls showing the highest BCFs (fresh weight): up to about 0.13 in highaccumulator cultivars. Average BCF for PCBs was 0.045, and for PCDDs and PCDFs, 0.01. BCFs in low-accumulator cultivars were typically less than 0.001, although up to 0.003 for tetrachloro-biphenyl and 0.002 for the penta- and hexachloro-biphenyls.

Hulster et al (1994) provide BCFs for different PCDD/PCDF-homologue groups for zucchini fruit. The values from Inui et al (2008) and Hulster et al, 1994 are shown in table 47. The recommended values for cucurbits are the geometric mean values determined from Inui et al (2008) and Hulster et al (1994), and are also show in table 47.

Polychlorinated	Inui et al (2008)		Hulster et al (1994)		Recommended BCF	
dibenzodioxins	Fresh weight	Dry weight ^a	(dry weight)		value (dry weight)"	
TCDD	0.065	0.65	0.25	0.08	0.24	
1,2,3,7,8-PeCDD	0.035	0.35	0.2	0.09	0.18	
1,2,3,4,7,8-HxCDD	0.01	0.1	0.17	0.04	0.09	
1,2,3,6,7,8-HxCDD	0.01	0.1				
1,2,3,6,7,9-HxCDD	0.01	0.1				
1,2,3,4,6,7,8-HpCDD	<0.001	<0.01	0.03	0.01	0.017	
OCDD	<0.001	<0.01	<0.005	<0.005	<0.005	
2,3,7,8-TCDF	0.045	0.45	0.14	0.09	0.18	
1,2,3,7,8-PeCDF	0.022	0.22	0.14	0.011	0.07	
2,3,4,7,8-PeCDF	0.02	0.2				
1,2,3,4,7,8-HxCDF	0.005	0.05	0.1	0.04	0.06	
1,2,3,6,7,8-HxCDF	0.005	0.05				
1,2,3,7,8,9-HxCDF	<0.001	<0.01				
2,3,4,6,7,8-HxCDF	0.05	0.5				
1,2,3,4,6,7,8-HpCDF	<0.001	<0.01	0.03	0.01	0.017	
1,2,3,4,7,8,9-HpCDF	<0.001	<0.01				
OCDF	<0.001	<0.01	0.01	0.05	0.02	
Total PCBs	0.045	0.45			0.45	

Table 47:BCF for dioxins and dioxin-like PCBs from Inui et al (2008), Hulster et al
(1994) and recommended BCFs

a Calculated using a conversion factor of 0.1 kg DW/kg FW.

b Geometric mean of data provided in Inui et al (2008) and Hulster et al (1994).

6.11.2 Calculation for the dioxin and dioxin-like PCB standards

The most common occurrence of dioxins in New Zealand contaminated land is that associated with the use of pentachlorophenol (PCP) as a wood preservative. The dioxin contamination in PCP is dominated by the octa- and hepta-chlorinated congeners (in TEQ terms). Dioxin contamination associated with the manufacture of trichlorophenol and 2,4,5-T herbicide is dominated by 2,3,7,8-TCDD. Combustion-derived dioxin mixtures will fall between these extremes.

Because the lower chlorinated congeners are taken up in cucurbits considerably more than the octa- and hepta-congeners, it can be expected that the produce pathway will be more significant for dioxin mixtures dominated by the lower-chlorinated congeners. Similarly, the produce pathway is significant for dioxin-like PCB mixtures. Because of this, SCSs have been separately calculated for OCDD- and TCDD-dominated mixtures and dioxin-like PCBs for residential scenarios with produce consumption (appendix 4). As noted above, the only vegetable type with significant dioxin uptake is the cucurbit family. The proportion of cucurbits relative to total vegetable consumption has been taken to be 0.04 for calculating the consumption-weighted BCFs.

It is necessary to choose representative BCFs from table 47 to calculate the residential-withproduce consumption SCSs. For TCDD-dominant mixtures, the BCF value of TCDD was chosen as conservative. For OCDD- and HpCDD-dominated mixtures, the BCF for HpCDD was chosen rather than the lower value for OCDD, again to be conservative.

It is also necessary to choose representative skin absorption factors. PCDFs contribute little to the TEQ in most PCDD/PCDF mixtures encountered, therefore the PCDD skin absorption factor 0.02 recommended in MfE (2011) has been used for all dioxin SCS derivations, but the larger factor 0.07 that is specific to dioxin-like PCBs has been used for that derivation.

The soil contaminant standard calculations for dioxin and dioxin-like PCBs use the contaminant-specific parameters in table 48 and the derived SCSs are set out in table 49. Dioxins and dioxin-like PCBs have been assumed to be threshold substances in the derivation.

Only the SCSs for TCDD and dioxin-like PCBs for residential scenarios are presented in table 49. The calculated SCSs with produce consumption for OCDD and TCDD are similar for 10 per cent homegrown produce, as the uptake into cucurbits has only a small influence on the final value (appendix 1). The TCDD values have therefore been chosen as the default value to cover all dioxin mixtures. However, the SCSs for derived dioxin-like PCBs are sufficiently different from the TCDD values to warrant separate default values for PCB mixtures.

It may be advantageous to derive site-specific SGVs for OCDD-dominated or other mixtures, rather than using the TCDD defaults, particularly if the occupants of the particular site grow a high proportion of their vegetables. Use of other than the defaults would have to be demonstrated on a case-by-case basis.

Table 48:	Contaminant-specific parameters for derivation of the dioxin and dioxin-like
	soil contaminant standards

Tolerable daily intake: oral		0.000 001 µg/kg BW/day				
Background	Child	0.000 000 33 µg/kg BW/day				
exposure	Adult	0.000 000 33 μg/kg BW/day				
		TCDD OCDD PCBs		PCBs		
Dermal absorption factor		0.02	0.02	0.07		
Plant	Leafy	0	0	0		
bioconcentration factors	Root	0	0	0		
	Tuber	0	0	0		
	Cucurbits	0.24	0.017	0.45		
	Consumption weighted mean	0.0096	0.000 68	0.018		

Table 49:	Dioxin and dioxin-like PCB soil contaminant standards (µg TEQ/kg)
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Scenario	Combined soil contaminant standards					
	No produce		10% produce		25% produce	
	Dioxins	PCBs	Dioxins	PCBs	Dioxins	PCBs
Rural residential / lifestyle block	0.18	0.16	0.15	0.12	0.12	0.09
Standard residential	0.18	0.16	0.15	0.12	0.12	0.09
High-density residential	0.35	0.33				
Recreational	0.6	0.52				
Commercial / industrial indoor worker	NL	NL				
Commercial / industrial outdoor worker / maintenance	1.4	1.2				

Note: NL = No limit.

For scenarios without produce consumption, soil ingestion is the dominant pathway for dioxins and dioxin-like PCBs, but dermal absorption affects the final SCS by up to about 10 per cent, depending on the congener mix / compound type. Skin absorption is greater for PCDFs than for PCDDs and absorption for dioxin-like PCBs is greater again (MfE, 2010b). The produce consumption pathway is a significant pathway for those residential scenarios with home-grown produce consumption, and becomes dominant for high home-grown produce percentages. The produce consumption pathway has the greatest significance for the dioxin-like PCBs.

Note: The SCS values for dioxins have units of µg TEQ/kg, that is, 1000 times lower than the other SCSs presented in this document. The guidelines are applicable to dioxin or PCB concentrations after converting to TEQ (toxic equivalents) and should not be applied to individual congener concentrations or a simple sum of dioxin or PCB concentrations. The PCB guideline is not applicable to the many non-dioxin-like PCBs. The guidelines should also not be applied to dioxin 'screens' for which all the 2,3,7,8 congeners have not been analysed, without considering whether the analytical screen provides a sufficiently good estimate for the TEQ. For example, the commonly used OCDD screen¹⁴ may underestimate the TEQ by a factor of two to three for a PCP-contaminated site. The PCB screens currently available from New Zealand laboratories include all the dioxin-like PCBs.

¹⁴

The OCDD screen provides concentrations of OCDD and 1,2,3,4,6,7,8-HpCDD.

6.11.3 Dioxin and dioxin-like PCB uptake into home-produced eggs

Eggs may accumulate lipophilic organic compounds, if the hens producing the eggs forage in an area contaminated with such compounds. Dioxins and dioxin-like PCBs are lipophilic. Several studies have noted that free-range eggs have higher concentrations of dioxins than commercial eggs (eg, Van Overmeire et al, 2006 and references contained in Van Overmeire et al, 2009). The contact with the outdoor environment, in particular soil and soil organisms, is believed to contribute to egg contamination of free-range hens (Stephens et al, 1995; Schuler et al, 1997; Harnley et al, 2000; Van Overmeire et al, 2009).

Keeping hens for egg production is a reasonably common activity on rural properties and some urban-dwellers also keep hens. This may be a more common activity for households who already grow a significant proportion of their vegetable consumption. It is therefore appropriate to consider a SGV incorporating both consumption of home-grown produce and home-produced eggs. Accordingly, SGVs have been derived for illustrative purposes – but these should not be used for site assessment without consideration of the site-specific circumstances. Three produce scenarios and two egg-consumption scenarios have been considered: 0, 10 and 50 per cent of vegetables consumed being home-grown for each of average egg consumption and twice the average egg consumption (the latter arbitrarily chosen as being a more likely estimate for those households that run hens).

According to Statistics New Zealand (see www.stats.govt.nz), New Zealanders consume an average of about 200 eggs per year, or a little over half an egg per day. This is somewhat higher than the average 20 g/day¹⁵ assumed for a 25+ adult (averaged across males and females) in the simulated diets of the New Zealand Total Diet Survey. The survey also assumes a 1- to 3-year-old toddler consumes an average of about 8 g/day (Brinsdon, 2004).

Dioxins are taken up into the egg lipids, therefore it is the egg-lipid content of eggs that is important in determining exposure. Egg lipids are 10–11 per cent of the total egg weight (Van Overmeire et al, 2009), or, using the consumption values from Brinsdon (2004), the equivalent of about 2.2 grams lipid per day for an adult and 0.9 grams lipid per day for a toddler (taking the higher lipid percentage and rounding to one decimal place). When normalised by standard body weights, this converts to 0.03 and 0.06 g lipid/kg BW/day. The child is therefore the critical receptor with respect to eggs, as it is for soil ingestion and produce consumption.

There is no statistical information on egg consumption of New Zealand toddlers, so the egg lipid consumption value derived from Brinsdon (2004) has been used for the SGV calculation.

¹⁵ 20 g/day equates to about 0.4 egg/day if it is assumed the edible portion of a 'typical' egg is about 50 g. The most common egg sizes available for purchase are sizes 6 and 7, which must be a minimum of 53 and 62 g, respectively. If it is assumed a typical egg lies between these sizes, then such an egg would weigh approximately 60 g. The egg shell makes up 11–13 per cent of the egg, suggesting the edible part is around 50 g.

To calculate dioxin intake from eggs and hence a SGV, a relationship between soil concentration and egg lipid concentration is necessary. Some studies have found a correlation between egg and soil concentrations (Harnley et al, 2000; Van Overmeire et al, 2009), others have not (Schuler et al, 1997; Pirard and de Pauw, 2004). This lack of a relationship is likely a reflection of the amount of soil and/or soil organisms ingested by the chickens (Schuler et al, 1997; Van Overmeire et al, 2009). This in turn will be influenced by foraging activity, the number of chickens in a given area and the extent of bare soil exposed, and therefore the amount of soil and/or soil organisms available to be ingested per chicken.

Schuler et al (1997) provides an estimate of egg lipid/soil concentration factors based on soil concentrations in a hen yard for different PCDD/PCDF congeners. Additionally, crude egg lipid/soil ratios can be derived from the median egg lipid and soil concentrations provided in Van Overmeire et al (2009), who assessed PCDD/PCDF concentrations in eggs and soil from 10 private hen owners. The egg lipid/soil ratios from Schuler et al (1997) and Van Overmeire et al (2009) are similar in the sense that both are based on egg lipid with an unknown / variable amount of soil ingestion, and yield similar ratios (table 50).

Few studies have examined the uptake of dioxin-like PCBs into eggs (Van Overmeire et al, 2009 and references contained therein), although PCBs in eggs may contribute significantly to the total toxic equivalent concentrations. For example, Van Overmeire et al (2009) found dioxin-like PCBs contributed, on average, 47 per cent to the total TEQs in eggs compared with 14 per cent in soil.

The egg lipid / soil ratios for dioxin-like PCBs determined from median egg lipid and soil concentration data in Van Overmeire et al (2009) are shown in table 50. These ratios may be used to provide crude estimates of potential contamination of eggs as a result of running hens on soils with elevated PCDD/PCDF and dioxin-like PCB concentrations. A better approach, if some information about a given site is known, would be to use a model similar to that developed by Waegeneers et al (2009), which accounts for different sources of intake, bioavailability of dioxins and variable intake of soil depending on hen-run conditions.

Table 50 shows greater uptake into eggs for the less chlorinated congeners and for dioxin-like PCBs. This is similar to the plant uptake behaviour.

For the purposes of illustrative calculation, egg-lipid concentration factors of 1.9 for TCDD dominated mixtures, 0.7 for OCDD/HpCDD-dominated mixtures (average of the OCDD and HpCDD ratios given in table 50) and 17 for dioxin-like PCBs¹⁶ were used.

The detailed calculations (appendix 4) show egg consumption dominates all other pathways. The same combined SGV is arrived at regardless of home-grown produce consumption percentage. The results are shown in table 51.

¹⁶ Dioxin-like PCBs occur in small quantities within commercial PCB mixtures such as the Aroclor brand manufactured by General Electric. Aroclor 1254 and Aroclor 1260 were two mixtures commonly used in capacitors. Frame et al (1996) provide typical compositions of Aroclor mixtures. These have been used to calculate weighted average egg lipid/soil ratios, which for both Aroclor 1254 and 1260 was about 17 and for Aroclor 1242 about 14. Actual concentrations should be used to calculate a weighted average egg lipid/soil ratio for a site-specific analysis.

Dioxin	Schuler, 1997	Derived from Van Overmeire et al, 2009	Average
TCDD	1.2	2.7	1.9
1,2,3,7,8-PeCDD	2.4	4.1	3.2
1,2,3,4,7,8-HxCDD	1.5	2.5	2.0
1,2,3,6,7,8-HxCDD	1.6	2.7	2.2
1,2,3,6,7,9-HxCDD	0.8	1.4	1.1
1,2,3,4,6,7,8-HpCDD	0.4	0.9	0.6
OCDD	0.1	0.5	0.3
2,3,7,8-TCDF	3.3	2.4	2.9
1,2,3,7,8-PeCDF	4.4	1.8	3.1
2,3,4,7,8-PeCDF	0.8	1.7	1.2
1,2,3,4,7,8-HxCDF	0.9	0.9	0.9
1,2,3,6,7,8-HxCDF	1	0.8	0.9
1,2,3,7,8,9-HxCDF	0.1	1.0	0.6
2,3,4,6,7,8-HxCDF	0.6	0.7	0.6
1,2,3,4,6,7,8-HpCDF	0.2	0.2	0.2
1,2,3,4,7,8,9-HpCDF	0.1	1.0	0.5
OCDF	0.1	0.2	0.15
PCB77		6.4	
PCB81		3.1	
PCB126		7.9	
PCB169		6.8	
PCB105		15.8	
PCB114		18.5	
PCB118		17.2	
PCB123		13.3	
PCB156		17.2	
PCB157		16.1	
PCB167		18.4	
PCB189		17.8	

Table 50: Egg lipid / soil ratios for PCDD/PCDFs and dioxin-like PCBs

Table 51:Dioxin and dioxin-like PCB soil guideline values for egg consumption
pathway (µg TEQ/kg)

Scenario	Combined soil guideline values					
	Average egg consumption			Twice average egg consumption		
	TCDD	OCDD/ HpCDD	Dioxin-like PCBs	TCDD	OCDD/ HpCDD	Dioxin-like PCBs
Rural and standard residential	0.005	0.013	0.000 6	0.003	0.007	0.000 3

The calculations show that running hens on a dioxin-contaminated property is generally not advisable without precautions against hens foraging in contaminated areas. However, a number of assumptions have been made in these calculations. Assessment of such a site should be carried out using site-specific information, in particular the actual amount of eggs consumed, the actual dioxin or dioxin-like PCB concentrations in the soil or, better still, the concentrations within the home-produced eggs.
6.12 Pentachlorophenol

6.12.1 Bioconcentration factor for PCP

A number of factors influence plant uptake of PCP. For example, if PCP does not persist in the soil for a sufficiently long period (eg, over the growth period of the plant), a significant quantity is unlikely to be taken up into a plant (Anon, 1987). The form in which PCP exists in the soil is also a critical factor in determining plant uptake. For example, in a non-ionised form a limited amount of PCP will be taken up into plants through dissolution in soil pore water and passive diffusion into the roots (Anon, 1987).

A number of models describing the accumulation of *non-ionised* organic compounds in plants exist (eg, Travis and Arms, 1988; Ryan et al, 1988). However, at pH 6.7, as much as 99 per cent of PCP is ionised and exists as pentachlorophenate anion – compared to 1 per cent ionisation at pH 2.7 (Crosby, 1981). Pentachlorophenate is highly soluble in water and leaches readily. However, while ionisation potentially increases the uptake of PCP into a plant (due to increased solubility in water), no model is available to adequately describe plant uptake of *ionised* organic compounds. Furthermore, PCP can be metabolised by plants, which will also reduce the concentrations present in the plant (Weiss et al, 1982 in Anon., 1987; Casterline et al, 1985; Haque et al, 1988; Bellin and O'Connor 1990). In fact, based on the weight of evidence of uptake, metabolism and elimination of PCP, the Canadian Council of Ministers of the Environment concluded that bioaccumulation of PCP in plants would not be significant (CCME, 1997).

In contrast, US EPA (2007) determined a BCF of 5.93 for PCP in plant foliage, which was the median of 10 data points from four studies, of which only one was based on field data. The field study (Bellin and O'Connor 1990) reported BCFs in fescue of less than 0.000 72 and 0.000 1 (dry weight), in contrast to the laboratory studies which resulted in calculated BCFs ranging from 2.3 to 46. Only one of these studies included verification of intact PCP in plant tissue (Casterline et al, 1985). Calculated BCFs for PCP (in spinach and soybean) from this study ranged from 2.3 to 7.8. Bellin and O'Connor (1990) suggested the higher BCFs determined by Casterline et al (1985) are due to the persistence of PCP in the soil. Other authors (Scheunert et al, 1986) suggest that radio-labelled PCP metabolites, including CO_2 , arising from the degradation of radiolabelled PCP used in the experiments were responsible for the observed radioactivity in plant tissue – giving rise to 'erroneously' high BCF values for PCP based on that radioactivity.

Dutch authors use models for estimating plant uptake of PCP, and suggest it forms a significant pathway of exposure.

The 'Timber Treatment Guidelines' (MfE and MoH, 1997) consider plant uptake of PCP and nominally use a BCF(stem) of 0.09 (wet weight, 0.4 dry weight) for PCP: this is based on the Travis and Arms (1988) relationship and a log K_{ow} for PCP at pH 7 of 3.3 (appendix A1 in MfE and MoH, 1997). However, in the tables for calculating the guideline values (tables 5.10 and 5.12 in MfE and MoH, 1997) a soil / stem concentration of 63.5 or a BCF (stem) of 0.016 and BCF (root) of 0.078 is apparently used (the latter derived by assuming stem concentrations are 20 per cent of root concentrations). The basis for these BCFs is unclear.

However, it is considered that plant uptake of PCP is not a significant pathway of exposure – given that:

- PCP is known to be metabolised by plants (and hence there is over-prediction of plant uptake by models predicting plant uptake of organic contaminants)
- BCFs reported in a field-based studies are low
- recent papers on plants and PCP in soil focus on phytoremediation (through enhanced microbial activity associated with plant roots: eg, He et al, 2005; Lin et al, 2006) as opposed to plant uptake.

6.12.2 Calculation for PCP standard

The soil contaminant standard calculations for PCP use the contaminant-specific parameters in table 52 and the derived SCSs are set out in table 53. PCP has been assumed to be a threshold substance. The BCF for all produce types has been set to zero, indicating no plant uptake. This results in SCSs for residential scenarios with produce uptake being the same as scenarios without produce uptake.

 Table 52:
 Contaminant-specific parameters for the derivation of the PCP soil contaminant standard

Tolerable daily intake: oral	0.000 3 mg/kg BW/day		
Background exposure	0.000 02 mg/kg BW/day 0.000 02 mg/kg BW/day		
Dermal absorption factor	0.24		
Plant bioconcentration fac	tor	0	

 Table 53:
 PCP soil contaminant standard (mg/kg)

Scenario	Combined soil contaminant standards							
	No produce	10% produce	25% produce					
Rural residential / lifestyle block	55	55	55					
Standard residential	55	55	55					
High-density residential	110							
Recreational	150							
Commercial / industrial indoor worker	NL							
Commercial / industrial outdoor worker / maintenance	360							

NL = No limit.

The SCS for PCP is dominated by the soil ingestion pathway, however, dermal absorption is also significant.

It should be noted that, as technical grade PCP was contaminated with dioxins, consideration should be given to investigating dioxins even if SCSs for PCP are complied with. Data from the investigation of New Zealand sawmill sites (T&T and SPHERE, 2008) shows that at PCP concentrations between about 0.3 and 55 mg/kg (the latter being the residential SCS for PCP), roughly 50 per cent of samples will also exceed the residential SCS for dioxins of 0.15 μ g TEQ/kg.

7

Summary of Soil Contaminant Standards and Guideline Values

	Arsenic	Arsenic Boron ¹ Cadmium		Chron	nium ¹	Copper ¹	Inorganic	Inorganic
			(pH 5) ²	111	VI		lead	mercury compounds ³
Rural residential / lifestyle block no produce	21	NL^4	110	NL	770	NL	250	510
Rural residential / lifestyle block 10% produce	17	NL	3	NL	460	NL	210	310
Rural residential / lifestyle block 25% produce	17 ⁵	NL	0.8	NL	290	NL	160	200
Residential no produce	24	NL	110	NL	770	NL	250	510
Residential 10% produce	20	NL	3	NL	460	NL	210	310
Residential 25% produce	17 ⁵	NL	0.8	NL	290	NL	160	200
High-density residential	45	NL	230	NL	1,500	NL	500	1,000
Recreational	80	NL	400	NL	2,700	NL	880	1,800
Commercial / industrial outdoor worker / maintenance	70	NL	1,300	NL	6,300	NL	3,300	4,200

Table 54:	Summary of soil contaminant standards (shaded) and guideline values for
	inorganic substances <i>(unshaded)</i> (mg/kg)

1 SCSs for boron, chromium III and copper are much greater than the soil concentration at which plant health will be affected. Plant and other environmental effects may need to be considered separately.

2 Default value is for pH 5. See appendix 1 for SGVs at other soil pH values.

3 The inorganic mercury SCS does not apply to elemental (pure) mercury.

4 NL = No limit. Derived value exceeds 10,000 mg/kg.

5 Derived value replaced with 99th percentile of national dataset of background concentrations (appendix 6). Note: Shading indicates SCS used for the purpose of the NES.

Table 55: Summary of soil contaminant standards (shaded) and guideline values for organic compounds (unshaded)

Scenario	BaP ¹	DDT (mar/lan)	Dieldrin ²		Dioxin (µg/kg TEQ)⁴		
	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	TCDD	Dioxin-like PCBs⁵	
Rural residential / lifestyle block no produce	11	120	22	55	0.18	0.16	
Rural residential / lifestyle block 10% produce	8	70	2.6	55	0.15	0.12	
Rural residential / lifestyle block 25% produce	6	45	1.1	55	0.12	0.09	
Residential no produce	12	120	22	55	0.18	0.16	
Residential 10% produce	10	70	2.6	55	0.15	0.12	
Residential 25% produce	7	45	1.1	55	0.12	0.09	
High-density residential	24	240	45	110	0.35	0.33	
Recreational	40	400	70	150	0.60	0.52	
Commercial / industrial outdoor worker / maintenance	35	1,000	160	360	1.4	1.2	

- 1 SCS to be compared with the equivalent BaP concentration calculated as the sum of each of the detected concentrations of the nine PAHs listed in table 40 multiplied by the respective PEF.
- 2 SCS for dieldrin also applies to aldrin separately, or to the sum of aldrin and dieldrin where both are present.
- 3 Consideration should be given to investigating dioxins for PCP concentrations in excess of 0.3 mg/kg, see last paragraph of section 6.
- 4 TCDD (WHO, 2005) TEQ calculated as the sum of each of the 17 PCDDs and PDDFs, or 12 PCBs listed in table 46, multiplied by the respective 2005 WHO TEF (table 46).
- 5 The SCS applies to only the 12 dioxin-like PCBs. The 'ordinary' toxicity of the simple sum of the concentrations of these and all other detected PCBs must be considered separately.

Note: Shading indicates SCS used for the purpose of the NES.

8

Limitations of Soil Contaminant Standards

The use to which the soil contaminant standards are put is inevitably limited by the assumptions behind the derivations, in particular the limitations of the chronic toxicity values (MfE, 2011), the generic exposure scenarios (Section 3) and the specific exposure parameters (Section 5). There will be actual scenarios for which the standards are not applicable and should not be used. Scenarios that are outside the intended use include:

1. Protection against allergic reactions. The toxicity values have not taken allergic reactions into account. Some of the contaminants are known to cause allergic reactions, however, in general such allergies occur in only a small proportion of the population. Consideration of effects on small sub-populations (ie, outside 95 per cent of the population) is outside the scope of the SCSs.

A case in point is the allergic reaction some people suffer to chromium. The derived chromium III SCS is very high. In the unlikely event that actual soil concentrations are very high, the possibility that a site occupant could suffer from an allergy will require separate consideration.

2. Acute toxicity and the pica child. The toxicity values used to derive the SCSs are for chronic effects and the SCSs are therefore not appropriate for being applied to acute exposure situations. Checks to ensure that acute toxicity thresholds are not exceeded by a single exposure event as might occur for a pica child (who may consume gram amounts of soil for each exposure event) have not been carried out on the SCSs. This is because it is considered that behaviour modification is appropriate for the relatively small number of children who suffer from the condition, rather than having a modified SCS applied to the complete population.

It is open, however, for an assessor to carry out a site-specific check for a particular situation, and modify the SCSs accordingly. It is noted that there is a paucity of reliable acute toxicity thresholds that are required to carry out such an evaluation for many of the contaminants.

- 3. Consumption of eggs, milk and meat from animals raised on-site is excluded. Produce consumption is limited to home-grown vegetables. Sites for which consumption of home-grown eggs, milk or meat is important will need to be evaluated on a site-specific basis. Such assessment may be required where the contaminants of concern are lipophilic organic compounds (eg, organochlorine pesticides), but some heavy metal contaminants may be of concern for sites where the offal of home-raised animals is routinely consumed.
- 4. The SCSs are restricted to human health. This means that separate consideration will need to be given to other environmental receptors where the conceptual site model shows this to be relevant. This could include:
 - phytotoxity
 - soil microbial health
 - surface water and the aquatic life it contains
 - groundwater, whether for stock or human consumption, irrigation, or possible discharge to nearby surface water and its aquatic environment
 - stock or wildlife.

5. The SCSs do not apply to productive agricultural and horticultural land. The safety of commercially grown food crops is administered by the New Zealand Food Safety Authority. The SCSs have not been derived to ensure that food maximum residue limits (MRLs) are not exceeded in home-grown vegetables or commercial crops. While in most cases MRLs will not be exceeded, there may be some specific situations for some contaminants (eg, cadmium, lead and DDT) where MRLs are exceeded.

The policy basis for MRLs is quite different from the policy basis of the SCSs, in that MRLs are not health-toxicity-based numbers. Instead, the ALARA (as low as reasonably achievable) principle is used in setting MRLs. MRLs therefore have limited relevance to the application of SCSs for contaminated soil. In addition, the SCS derivation method should ensure human health is not compromised even if MRLs are exceeded because the total intake of a contaminant is taken into account in the derivation, including that from vegetables. The derivation method ensures that total intake does not exceed the allowable intake provided the actual soil concentration of the particular contaminant does not exceed the SCS for that contaminant.

It should also be noted that MRLs are not legally applicable to home grown produce but apply only when produce is sold on domestic or international markets. However, if homegrown vegetables were to be sold at a produce market, it is the legal responsibility of the grower/seller to ensure the produce complies with food safety standards.

9 Site-specific Risk Assessment

9.1 Purpose

The methods and guidance provided in this section have been prepared to support practitioners deriving, or contemplating deriving, site-specific soil guideline values as provided for in the National Environmental Standard for Assessing and Managing Contaminants in Soil to Protect Human Health. If accepted by the relevant territorial authority, these become legally binding for the particular piece of land.

9.2 Introduction

Site-specific or 'Tier 2' assessment in contaminated site practice is using site-specific information to modify the generic assumptions used for the SCS derivation; this will more accurately estimate a person's exposure and therefore the risk to human health for the particular situation. Site-specific soil guideline values (SSGV) are then derived using the same basic methodology used to derive the generic standards as described in section 4 and 5 in this document.

Site-specific assessment considers each element in the hazard (source)-pathway-receptor model of risk assessment, and theoretically could involve modifying one or more of:

- the toxicity of the chemical of concern, particularly in relation to chemical speciation (overlaps with bioavailability issues)
- the default assumptions about the receptors considered to be at risk in the generic derivations, including the physical characteristics of those receptors (eg, weight, skin areas) and their behaviour (frequency and duration on the site, lifestyle)
- the exposure (intake) estimates, such as whether all the exposure pathways assumed to exist actually exist, or exist to the degree assumed; and whether the generic exposure rates (eg, soil ingestion, soil adherence, produce consumption) and other exposure factors are realistic for the particular situation.

As a fundamental starting point, any site-specific assessment needs a well-developed conceptual site model with all contaminants of concern, exposure pathways (and any barriers) and receptors identified and quantified.¹⁷ A good understanding of the soil concentrations and distribution of concentrations is required. Only then can consideration be given to modifying the generic scenarios and associated factors.

Site-specific assessment is a task for an appropriately qualified and experienced contaminatedland professional. Expert professional judgement and an intimate knowledge of the derivation methodology is required to vary factors used in the derivation of the guideline values. In considering site-specific assessments, local authorities may need to obtain independent expert review.

¹⁷ Refer to *Contaminated Land Management Guidelines No. 5* (MfE, 2004b) for a discussion on the conceptual site model and sampling requirements.

When a site-specific assessment should not 9.3 be carried out

One could argue that a site-specific assessment should be carried out on any site because not one will exactly fit the generic exposure assumptions. Also, the SCS are deliberately conservative so that many sites fitting a generic scenario would still be safe at concentrations in excess of the SCS. However, the intention is that, for sites that generally fit within a particular generic exposure scenarios set out in Section 3.1, and there is no resource consent or other mechanism (eg, encumbrance on land titles) providing for site-specific management of exposure, then site-specific assessment is not allowed. This is because without the enforcement mechanism provided by the resource consent, there is no guarantee that any current site-specific circumstances creating lower exposure will remain into the future.

For example, all urban housing with a typical section will fit within the standard residential scenario, even if there is no vegetable garden; they should be assessed as such unless a resource consent that allows variation of the standard exposure is applied for and approved.

Site-specific principle 1:

Site-specific assessment is not permitted for sites that fit within the generic scenarios of appendix 1 above unless a resource consent is granted that ensures the exposure assumed in the site-specific assessment will continue into the future.

It may not be economic to go to the expense of a site-specific assessment for some sites, particularly small or low-value sites, where a small amount of remedial work would cost less than the cost of the site-specific assessment and applying for a resource consent to manage the site. In that case, site-specific assessment should not be carried out.

Site-specific assessment should also not be carried out if the owners decide that remediation best suits their intentions for the site. Some owners prefer to know their site is fit for any purpose.

When a site-specific assessment must be 9.4 carried out

Site-specific assessment must be carried out if it is clear the current site use, or intended site use, does not fit within any of the generic exposure scenarios such that the selection of the most relevant generic SCS would under-estimate actual human exposure. In this situation the derivation of a site-specific SGV would be indicated and would become mandatory via a resource consent. Note, however, that the assessor is free to use a more conservative generic scenario so long as it is sufficiently protective. For example, if the actual use of a park was more intensive than the generic parkland scenarios envisaged, then use of the residential SCS would be sufficiently protective without the need to go to a site-specific assessment.

Site-specific assessment is not triggered or required, however, for reassessing the proportion of home-grown produce consumed. The two generic land-use scenarios included within the scope of the national environmental standard are based on an assumption that up to 10 per cent (residential) or 25 per cent (rural residential / life style block) of fruit and vegetables consumed are grown on the property.

Any site-specific assessment and adjustment of the SCS to allow for greater produce consumption is a voluntary step and cannot be required. If the adjusted SCS (now referred to as a SSGV) is exceeded, non-regulatory advice is appropriate. In this situation it would be good practice to record the circumstances on the property file, and inform the owner of the exposure risk and the range of measures that could be adopted to mitigate this risk. Although not a NES requirement, a site-specific assessment is strongly advised if the owners consume home-produced eggs, poultry or dairy products **and** the contaminants of concern are highly toxic lipophilic compounds (eg, dioxins).

Site-specific principle 2:

Site-specific assessment must be carried out if the current site use, or intended site use, results in greater human exposure than for any of the generic exposure scenarios.

9.5 When a site-specific assessment may be carried out

Site-specific assessment and derivation of site-specific SGV is appropriate if both of the following apply:

- a site has been sampled and the results exceed one or more relevant SCS
- the generic land-use scenarios for which SCS are available do not fit the actual site use or configuration with sufficient accuracy. In this case the generic SCS for the contaminants of concern are too protective, resulting in an unjustified restriction on site use or unnecessary remediation with associated financial burden.

As noted in *Contaminated Land Management Guidelines No. 5* (MfE, 2004), it is not envisaged that occasional SCS exceedances would necessarily trigger a site-specific assessment (or management action or remediation), rather the site should be assessed on the basis of average exposure over appropriate exposure (averaging) areas, taking into account any hotspot contamination, as necessary.

There is no compulsion to carry out site-specific assessment, unless it is to support a resource consent application. An owner may be prepared to tolerate, or require, a more conservative assessment than the site use would suggest, and would then carry out remediation as appropriate to any SCS exceedances.

Particular situations where site-specific assessment **could** be carried out fall into two situations:

- SCS are exceeded for a site that fits squarely with one of the generic definitions, but the site 1. has current or proposed circumstances that limit exposure, and the owner is to apply for a resource consent to permit limited remediation or management.
- The site falls outside the generic exposure scenarios, or between two of the generic 2. scenarios, and is of a type that the use is not likely to change for the foreseeable future, eg, here is a long history of the particular use, or the land is designated for particular purposes (eg, education), or there are district plan restrictions on the type of use for the particular site. A resource consent would not be required in these cases, as a change to some more sensitive use is likely to come to the attention of the territorial local authority and a reassessment would be required.

Examples of the former situation include:

- A conventional residential property where there is no vegetable garden and no likelihood of a garden (eg, the backyard is too small or is paved) and this situation will be preserved by restrictions imposed by a resource consent.
- A conventional residential property where there is, or intended to be, a vegetable garden, but the owner has installed (or will install) a raised-bed garden with clean soil. The owner would need a resource consent that restricted vegetable gardens to raised-bed gardens. In this case the site-specific assessment may be to simply recalculate the SCS without the produce pathway.

Examples of the second situation include:

- A childcare centre a residential scenario could be used as a conservative screening but site-specific assessment will probably result in higher (less conservative) SGV. This would only be appropriate where the site has been, or is likely to be, a childcare centre for a long time. A childcare centre in a converted house that may revert to residential at any time should be assessed as residential.
- Primary or secondary schools a residential scenario is too conservative. A parkland scenario might be appropriate for parts of the site, but a site-specific assessment taking into account different sub-uses of the school grounds is more appropriate.
- Rural land not used for residential accommodation, or reserve land used for occasional or passive recreation where the generic human-health scenarios provided are probably too conservative.

9.6 What factors may be changed?

Any factors may be changed **except**:

- the contaminant toxicity values
- the dermal absorption factors
- the averaging time for non-threshold substances (MfE, 2011)
- the use of 100 per cent contaminant oral bioavailability.

The *toxicity values* and *dermal absorption factors* have been considered and approved by a panel of government experts and should not be changed without going through the same process. Where toxicity values do not exist, then a similarly rigorous process to that described in *Toxicological Intake Values for Priority Contaminants in Soil* (MfE, 2011) in developing values for the current priority contaminants for which SCS have been derived, should be followed. However, if it is clear that people on a site have a higher background intake than has been assumed in the SCS calculation, then the residual tolerable daily intake assigned to soil must be reduced.

The *averaging time for non-threshold contaminants* is, by definition, a lifetime. Again, the duration of a lifetime has been approved by a panel of government experts on the basis of population statistics. Until such time as average life expectancy changes the value must remain fixed.

Contaminant bioavailability has been subject to much debate internationally. The consensus is that, currently, the test methods available in New Zealand for estimating site-specific bioavailability are not yet good enough and the use of generic bioavailability values from the literature is not appropriate.

Some factors are less likely (or less appropriate) to be changed than others. Factors that are less likely to be changed, or require greater justification to change, include: body weights, inhalation rates, and skin areas for given body parts for the standard receptors (ie, adults and young children). As skin areas are proportional to body weight, fixing body weights will fix total skin areas (but not skin area exposed if fewer or more body parts are likely to be exposed than the generic scenario).

Any varying of exposure factors should be fully justified in the assessment report, either on the basis of professional judgement or by citing scientific studies. The calculations should be presented.

Typical situations that would enable the generic factors to be changed are described in table 56.

Factor	Situations	Examples ¹
Background intake (increase only)	Where a non-soil exposure results in greater than the assumed background intake used for the generic SGV derivation	Groundwater used on-site has natural or anthropogenic contamination
Exposure duration	Non-residential situations where duration of occupancy is likely to be at variance from the standard situations	Childcare centre Secondary school Construction site
Exposure frequency	Occupancy for a typical person is discontinuous throughout the year, or for less than five or seven days per week	Some childcare facilities Schools Short-tem construction Some parks, public gardens and reserve land
Body weight	Situations where the critical occupant is at variance from the standard child (toddler) or adult weights	Primary school – choose body weight for 5-year- old female Secondary school – 12-year-old
Skin area	As for body weight	As for body weight
Soil ingestion rate	Where enforceable management controls create permanent or semi-permanent barriers to soil Partial remediation reduces area of soil above SGV ² Where the typical activities increase or reduce likelihood of soil ingestion	Installation of paving, decking, soil cap with marker layer, gravel with geotextile (reduced exposure) Parks and gardens for passive recreation (reduced exposure) Construction sites where excavation is carried out (increased exposure) ³
Dermal adherence factor	Activities that result in increased or decreased likelihood of getting dirty	Parks and gardens intended for passive recreation (little soil adherence) Excavation activities (increased soil adherence) ³ Lakes or beaches with contaminated sediments where children play
Percentage home- grown produce (in the extreme, eliminating this pathway)	Where site-circumstances or enforceable management controls eliminate or reduce risk of produce uptake Lifestyle of owners indicates substantial home-grown produce	Raised-bed vegetable garden allows dispensing with produce pathway Remediation of backyard allows dispensing with produce pathway Rural property with substantial vegetable garden and favourable climate for year-round growing
Produce uptake factors	Applicable only to residential gardens	Deriving site-specific bio-concentration factors - using soil and produce concentrations from the particular site ^{4, 5}
Additional pathways	Situations where the soil ingestion, dermal absorption and produce consumption pathways do not account for a significant part of the contaminant exposure	Extremely dusty sites such as mines and construction sites ³ Consumption of home-grown eggs, poultry or dairy where the contaminant is highly toxic and lipophilic On-site abstraction and use of groundwater impacted by contaminants

Table 56: Modifiable exposure factors, typical situations and examples

1 This is an indicative list. There are many other situations where adjustment of factors may be appropriate. The services of an experienced contaminated-site professional should be used to decide whether site-specific assessment is appropriate and, if so, the modified factors to be used.

2 Arguably, this does not need site-specific adjustment of the ingestion rate as the partial remediation enables the average site concentration to be redefined.

- 3 It is probable that such a situation would be controlled by a requirement for appropriate personal protective equipment and site occupational health and safety controls (a health and safety plan), rather than modifying allowable SCS.
- 4 Deriving site-specific bio-concentration factors (BCFs) will require measurement of soil concentrations and concentrations within appropriate plant species grown in the site soil (with sufficient testing to be statistically significant). If suitable plants species are not available, growing trials would be necessary. Field-scale studies are likely to be more realistic of the home gardener, but are difficult to control. Pot experiments are often seen as a compromise between control and realism. However, pot experiments also have known problems resulting in overprediction of plant uptake (EA, 2006).

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5 In theory, soil properties could be modified to reduce produce uptake, eg, liming to reduce pH or cause reduction of metal solubility. However, there is a question of the long-term effectiveness of such treatment where the intention is to meet regulatory requirements (eg, make a residential site suitable for growing up to 10 per cent of produce consumption). It is unlikely that such an approach would be acceptable to regulators unless it is conclusively demonstrated that the treatment would be effective, long-lived and/or irreversible for the particular site. Evidence required might include bench trials, field trials or full-scale remediation, with appropriate and sufficient (statistically significant) chemical testing. Testing could include such things as soil pH, soil mineralogy, soil organic carbon, soil clay content, and sequential extraction tests. However, there may be non-regulatory circumstances where such treatment may be appropriate and require a lower standard of evidence. For example, an owner is seeking to grow more than 10 per cent of vegetable consumption and the site complies with SCS for 10 per cent produce but would fail a non-regulatory SGV for a greater percentage.

9.7 The site-specific assessment process

The need for a site-specific assessment will tend to be an exception; before embarking on this path its relevance should be determined. For most sites it will not be economic or useful to carry out a site-specific assessment, since remediation or site management can be achieved simply or more readily.

The site-specific assessment process is set out in diagrammatic form in figure 6: Site-specific assessment steps. Carrying out a site-specific assessment presupposes the site is well understood and sufficient work has been carried out to properly characterise the soil conditions. If the site has not been properly characterised, or the site conceptual model is poorly developed, it is important that these be remedied before embarking on site-specific assessment. Further site characterisation may reveal, for example, that average site concentrations are actually below SCS values; and/or, the contaminant-pathway-receptor relationships are not as first thought and require better defining.

The steps shown in figure 6 should be carried out at a level of detail consistent with the size and complexity of the site. A small or simple site would warrant no more than a brief consideration of the various steps, while a complex site undergoing extensive investigation or a site undergoing a high-cost redevelopment could warrant detailed analysis at each step.

Step 1 involves reconsideration of the receptors at risk, the mechanisms by which those receptors might be exposed by contaminants, and whether the conditions exist on the site for that exposure to occur. Not all contaminants will behave in the same way, with the result that the exposure mechanism can be different for different contaminants, eg, the critical exposure pathway for a volatile organic compound may be different from a heavy metal. Setting out a matrix of contaminants (and associated characteristics), receptors and exposure pathways is a good way of assessing the linkages.

Step 2, in determining how actual exposure differs from the generic exposure, should consider for each receptor in turn (and for each exposure area relevant to that receptor):

- whether the standard parameters adequately describe the receptors' physical characteristics and exposure
- whether all of the standard pathways are relevant and, if not, which can be eliminated
- whether additional pathways are relevant.

If the answer is 'no' to either of the first two questions or 'yes' to the last, then it is necessary at **Step 3** to assemble sufficient information to decide whether it is practical and economic to proceed to a site-specific assessment. This could be very simply exercising professional judgement for a small site or a formal assembling of options and carrying out a cost-benefit analysis for a large complex site. This will tend to be interactive with Step 4, for complex sites.

Step 4 involves deciding on what factors can be changed (as set out in table 56), and what values these factors should take. This may involve carrying out soil and plant testing if produce bio-concentration factors are to be changed, researching receptor body weights and skin areas, researching occupancy, and carrying out literature searches to justify modifying soil ingestion rates or dermal adherence factors. Inevitably, professional judgement will be involved. At this point, a site-specific assessment would be abandoned if it is obvious that the site-specific SGV will not be sufficiently different from the SCS to warrant going further. Otherwise, the next step is to proceed to derive the site-specific SGV at **Step 5**, developing additional exposure equations if necessary for any additional exposure pathways that need to be considered. Fate and transport modelling may be appropriate for the inhalation and groundwater pathways.



Figure 6: Site-specific assessment steps

Site-specific assessment examples

Example 1: Step 1 – Confirming the site conceptual model

The site (about 5000 m²) is currently occupied by an engineering workshop, with underground storage tanks holding degreasing solvents and old machinery stored in one corner of the site. It is being considered for conventional residential redevelopment. The proposed residential lots will be of a sufficient size to have gardens, in keeping with the surrounding residential use. The site is generally level. The site geology is mixed fill overlying sandy gravels, with the water table at 2 m. A stream is on one boundary of the site.

Contaminant	Pathway(s)	Receptor			
Metals A, B, C	Ingestion, direct contact	Future residents, site workers			
	Consumption of contaminated vegetables	Future residents			
Semi-volatile, non-	Ingestion, inhalation, direct contact	Future residents, site workers			
halogenated	Consumption of contaminated vegetables	Future residents			
	Dermal contact	Future residents			
	Migration through fill	Groundwater in gravel			
	Migration through gravels	River			
Volatile halogenated	Inhalation through migration into buildings	Future residents, neighbours (possibly)			
hydrocarbons X, Y, Z	Ingestion, direct contact	Future residents			
	Consumption of contaminated vegetables	Future residents			
	Migration through fill	Groundwater in gravel			
	Migration through gravels	River			

Possible contaminant-pathway-receptor linkages

Example 2: Step 2 – Determining differences from the generic model

A secondary school has been built on an old landfill. The site has been investigated and found to contain elevated concentrations of lead, zinc, copper and arsenic. Landfill gas was not being produced. As an initial conservative screening, the results were compared with residential guidelines. It was found that zinc and copper were well below the residential SCS but that lead in surface soil was up to three times the residential SCS of 750 mg/kg and arsenic up to four times the SCS of 24 mg/kg.

The assessor considered the conceptual model and decided that the site should be broken into two areas, based on likely exposure – the area around the buildings which was entirely grassed or paved, and the playing fields where an individual pupil might practise and play contact sports on up to four occasions a week.

The assessor also decided receptors that needed considering were the pupils and the school caretaker. Teaching staff were considered less at risk than the caretaker and therefore not the critical adult. Twelve-year-olds were considered to be the youngest (and lightest) likely group at school and were therefore used as the critical child receptor. An average 12-year-old weighs about 40 kg. This weight is also about the 25th percentile for a 13-year-old.

For the building area, it was assumed the 12-year-old pupil was at school five days a week for 38 weeks of the year and the school caretaker was carrying out maintenance and gardening activities five days a week for 48 weeks of the year.

For the playing field area, it was conservatively assumed a 12-year-old pupil would be practising and playing contact sport on four occasions a week during term time (38 weeks per year) and the school caretaker would be mowing the fields and carrying out miscellaneous activities two days per week on average.

It was decided that the residential guideline for lead was too conservative as it was based on a 15 kg two-year-old child with 350 day exposure to bare soil. Instead, soil guideline values were calculated for each of the two areas using the 40 kg body weight typical of a 12-year-old child, and soil ingestion rates for high-density residential and recreational, as being an approximation for the activities for the two areas. As contaminant intake for the two areas is additive, part of the lead tolerable daily intake was assigned to each area in proportion to expected intake. The reduced exposure frequencies for the two areas were used. Produce consumption was dispensed with as not a valid exposure pathway.

After further examination of the exposure parameters, it was decided for the caretaker that the standard commercial / industrial outdoor worker scenario was sufficiently accurate, with calculation of a site-specific guideline not warranted. The measured concentrations did not exceed these guidelines. As such, the caretaker was not considered further.

For arsenic, a similar approach was taken for the 12-year-old child, but as arsenic is nonthreshold and exposure is averaged over a lifetime, the exposure duration becomes important. The exposure duration for the child was reduced from the standard 14 years residential exposure for an adult to the five years a typical child would spend at high school.

The calculated site-specific SGVs were in excess of the measured concentrations and no remediation was required. The school implemented a management plan to control off-site disposal of soil in the event of redevelopment.

Example 3: Steps 3–5 – Revision of site conceptual model and site-specific assessment

A rural subdivision was proposed on a former timber treatment site. The subdivision was being promoted as a sustainable development for families who wanted to get away from the city. The show-home was to be of packed-earth construction, double glazed and with solar heating. The advertising brochures showed houses with large vegetable gardens. There was to be an on-site communal water supply using groundwater.

The site had been used for chromated copper arsenate (CCA) and boron treatment, but the site had been tested and remediated to residential guidelines. The site had also been tested for pentachlorophenol (PCP) as a precaution, although there was no known history of pentachlorophenol PCP use on the site. PCP was below the detection limit in the tested locations.

During the development, a bore was sunk and tested and found to only marginally comply with the arsenic drinking-water guideline. In addition, a former site worker informed the developer that PCP had definitely been used in the boron dip for a few years in the early 1980s. The developer consulted an environmental consultant who advised the following:

- (a) The additional exposure to arsenic through drinking water meant that residents would be subject to a greater risk of cancer. The consultant further advised that it was not appropriate to calculate a site-specific soil guideline value as arsenic was a nonthreshold substance and background intake did not figure in the calculation. Instead, the consultant calculated the increased risk of cancer using the measured soil concentrations and the additional exposure from water.
- (b) Despite PCP being below the detection limit, it was possible that dioxin was present. Dioxin was a known contaminant of PCP and very resistant to degradation. Testing of dioxin was recommended at the boron dip, the diffusion shed location, and locations in between.

The site-specific calculation of the risk from arsenic found that the risk from soil and drinking water was less than 1 in 100,000, however the developer decided to include arsenic removal in the water treatment process.

Dioxin at concentrations in excess of the residential guideline was found at the former boron dip location, but below the residential guideline elsewhere. However, the consultant, knowing how the development was being promoted, advised the developer that the concentrations measured in some locations would exceed an site-specific soil guideline value calculated for 30 per cent home-grown produce and was well in excess of a guideline that took home-produced eggs into account.

The proposed lot containing the former dip location became a community tennis court. Encumbrances were placed on property titles preventing the keeping of chickens and farm animals (also consistent with the desirability of a quiet neighbourhood). Topsoil in future backyards was replaced with imported topsoil.

Example 4: Steps 4 and 5 – Eliminate pathways and calculate SCS

A residential site was found to have arsenic contamination at twice the SCS over the complete site as a result of past orchard use. Associated lead contamination, while above the SCS in places, was not critical compared with arsenic. Elevated copper was not critical as it was well below the SCS.

The site was considered to fall squarely within the standard residential scenario. As such, 10 per cent produce home-grown produce was applicable.

The assessor determined that if the produce consumption pathway was eliminated then the modified SCS (now SSGV) would be in excess of the measured concentrations. The assessor proposed to the owners that the site would comply if:

- the owner undertook to pave the backyard or replace the site soil with at least a half metre cap of clean soil (this could be in a raised-bed garden separated from the contaminated soil by a geotextile marker layer)
- applied to the council for a resource consent which would have as a condition restrictions on changing the new site configuration without obtaining permission from the council.

After consulting their architect, the owners decided to install a raised-bed garden over a third of the backyard, with the remainder landscaped with paving and a half metre of contoured soil capping. They applied for and were granted a resource consent.

The remainder of the site was left un-remediated.

Appendix 1: Detailed Calculations

Note: The combined soil guideline values (SGV) shown in the following calculations are not rounded in accordance with the SCS rounding rules (see beginning of section 6). For rounded soil contaminant standards ($SCS_{(health)}$) and guideline values see tables 54 and 55.

Generic factors												
Body weight (child): 13 k	kg		Averaging time (non-threshold): 75 years									
Body weight (adult): 70 k	g		Averaging time (threshold): 6 years									
Scenario-specific factors Lifestyle Stan block reside		Standard residential	High-density residential	Parks/ recreational	Commercial/ industrial indoor worker	Commercial/ industrial outdoor worker	Unit					
Exposure frequency	350	350	350	200	230	230	day/year					
Exposure duration (child)	6	6	6	6			years					
Exposure duration (adult)	24	14	14	8	20	20	years					
Soil ingestion rate (child)	50	50	25	25			mg/day					
Soil ingestion rate (adult or older child)	25	25	15	25	0	50	mg/day					
Age-adjusted ingestion factor	31.6	28.1	14.5	14.4	0	14.3	mg year/kg day					
Inhalation rate (child)	6.8	6.8	6.8				m³/day					
Inhalation rate (adult)	13.3	13.3	13.3	3.3 8		10.4	m³/day					
Age-adjusted inhalation rate	7.7	5.8	5.8		2.3	3.0	m ³ year/kg day					
Particulate retention	0	0	0	0 0		0	dimensionless					
Particle emission factor	-	-	-	-	-	-	m³/kg					
Skin area (child)	1,900	1,900	1,900	1,900			cm ²					
Skin area (adult)	4,850	4,850	4,850	3,670	3,670	3,670	cm ²					
Soil adherence (child)	0.04	0.04	0.02	0.04			mg/cm ²					
Soil adherence (adult)	0.01	0.01	0.005	0.04	0	0.04	mg/cm ²					
Age-adjusted dermal exposure factor	51.7	44.8	22.4	51.9	0	41.9	dimensionless					
Produce ingestion (child)	0.0105	0.0105					kg/day (DW)					
Produce ingestion (adult)	0.0322	0.0322					kg/day (DW)					
Proportion of above- ground produce	0.3	0.3	0.0	0.0	0.0	0.0	dimensionless					
Proportion of root (not tuber) produce	0.1	0.1	0.0	0.0	0.0	0.0	dimensionless					
Proportion of tuber produce	0.6	0.6	0.0	0.0	0.0	0.0	dimensionless					
Age-adjusted produce ingestion	0.0159	0.0113	0.0	0.0	0.0	0.0	kg year/kg day					

NL = No limit.

Health-based arsenic soil guideline values (mg/kg)			Threshold									
Oral RHS (mg/kg BW/day)		0.0000086	6 Skin absorption factor 0.005									
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		0.011						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		0.011						
Background intake child (mg/kg BW/day)		na	Bioconcentratio	on factor tuber	r	0.001						
Background intake adult (mg/kg BW/day)		na	Max conc. in p	roduce (mg/kg	g DW)	0.005						
Scenario	Soil ingestion	Dermal	Inhalation		Produce in	ngestion	1	Combined SGV				
RHS-child background (mg/kg BW/day)	0.000086	0.0000086	na	No	10%	2	25%	50%	No	10%	25%	50%
RHS-adult background (mg/kg BW/day)	0.000086	0.0000086	na	produce	produce	e pro	bauce	produce	produce	produce	produce	produce
Rural residential / lifestyle block	21	2,602	na	na	85		34	17	21	17	13	9
Standard residential	24	3,004	na	na	119		48	24	24	20	16	12
High-density residential	46	6,009	na	na					46			
Parks / recreation	82	4,540	na						80			
Commercial / industrial indoor worker	NL	NL	na						NL			
Commercial / industrial outdoor worker	72	4,881	na						71			

Health-based boron soil guideline values		Threshold							
Oral RHS (mg/kg BW/day)	0.2	Skin absorption	n factor						
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		na			
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		na			
Background intake child (mg/kg BW/day)		0.08	Bioconcentratio	on factor tuber		na			
Background intake adult (mg/kg BW/day)		0.017	Max conc. in p	roduce (mg/kg D	W)	300			
10% produce additional background intake		0.024							
25% produce additional background intake		0.061							
50% produce additional background intake		0.121							
Scenario	Soil ingestion	Dermal	Inhalation			Combin	ed SGV		
RHS-child background (mg/kg BW/day)	0.120	na	na	No produce	10% p	oroduce	25% produce	50% produce	
RHS-adult background (mg/kg BW/day)	0.183	na	na						
Rural residential / lifestyle block	32,537	na	na	32,537	25	,967	16,112	-	
Standard residential	32,537	na	na	32,537	25	,967	16,112	-	
High-density residential	65,074	na	na	65,074	65,074				
Parks / recreation	113,880	na	na	113,880					
Commercial / industrial indoor worker	NL	NL	na	NL					
Commercial / industrial outdoor worker	406,578	na	na	406,578					

Note: BCF not used for produce pathway. Additional background intake subtracted from TDI on assumption that maximum plant tissue boron concentration = 30 mg/kg dry weight. Method breaks down for produce percentage greater than 49%.

NL = No limit.

na = Not applicable.

Health-based cadmium soil guideline values (mg/kg)			Threshold									
Oral RHS (mg/k Dermal RHS (m Inhalation RHS Background inta Background inta	ig BW/day) ig/kg BW/day) (mg/kg BW/day) ake child (mg/kg BW/day) ake adult (mg/kg BW/day)		0.000 833 na na 0.00041 0.00026	Skin absorption Bioconcentratio Bioconcentratio Bioconcentratio Mean bioconce	n factor on factor leaf on factor root on factor tuber entration facto	r r	0.001 Depend on pH a soil conc. See separate calcula	nd tions				
Scenario		Soil ingestion	Dermal	Inhalation		Produ	ce ingestion			Combir	ned SGV	
					No produce	10% produce	25% produce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child back	ground (mg/kg BW/day)	0.00042	0.00042	na		0	.00042					
RHS-adult back	ground (mg/kg BW/day)	0.00057	0.00057	na		0	.00057					
Rural	pH 5	115	75,456	na	na	3.12	0.83	0.30	110	3	0.8	0.3
residential /	pH 5.5	115	75,456	na	na	5.19	1.40	0.51	110	5	1.4	0.5
mostyle block	pH 6	115	75,456	na	na	8.50	2.34	0.87	110	8	2.3	0.9
	pH 6.5	115	75,456	na	na	13.7	3.86	1.45	110	12	3.7	1.4
	рН 7	115	75,456	na	na	21.7	6.28	2.38	110	18	6	2.3
Standard	pH 5	115	75,456	na	na	3.12	0.83	0.30	110	3	0.8	0.3
residential	pH 5.5	115	75,456	na	na	5.19	1.40	0.51	110	5	1.4	0.5
	pH 6	115	75,456	na	na	8.50	2.34	0.87	110	8	2.3	0.9
	pH 6.5	115	75,456	na	na	13.7	3.86	1.45	110	12	3.7	1.4
	pH 7	115	75,456	na	na	21.7	6.28	2.38	110	18	6	2.3
High-density res	sidential	229	150,912	na	na				230			
Parks / recreation 401		132,048	na					400				
Commercial / in	dustrial indoor worker	NL	NL	na					NL			
Commercial / in	dustrial outdoor worker	1,273	433,602	na					1,300			

Note: The default soil pH for initial assessment is pH 5. The residential SGVs provided here for pH other than pH 5 are for site-specific assessment purposes where it can be demonstrated that some other pH is appropriate. Information for demonstrating that a pH greater than 5 is appropriate is provided in appendix 2.

NL = No limit.

Health-based chromium III soil guideline	values (mg/kg)		Threshold									
Oral RHS (mg/kg BW/day)		1.5	Skin absorptior	n factor		0						
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		0.0324						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		0.0324						
Background intake child (mg/kg BW/day)		0.0012	Bioconcentratio	on factor tube	r	0.0324						
Background intake adult (mg/kg BW/day)		0.00053	Mean bioconce	entration facto	r	0.0324						
Scenario	Soil ingestion	Dermal	Inhalation		Produ	ce ingest	ion			Combir	ed SGV	
				No 10% 25 produce produce prod			5% duce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	1.425	1.425	na			1.425						
RHS-adult background (mg/kg BW/day)	1.425	1.425	na			1.425						
Rural residential / lifestyle block	386,379	na	na	na	567,870) 227	,148	113,574	386,379	229,932	143,050	87,773
Standard residential	386,379	na	na	na	567,870) 227	,148	113,574	386,379	229,932	143,050	87,773
High-density residential	772,757	na	na	na					772,757			
Parks / recreation	1,352,325 ^ª	na	na						1,352,325 ^ª			
Commercial / industrial indoor worker	NL	NL	na						NL			
Commercial / industrial outdoor worker	3,165,978 ^ª	na	na						3,165,978 ^ª			

Note: Background intake less than 5% of the TDI, therefore background taken as 5% of the TDI.

NL = No limit.

na = Not applicable.

a calculated values greater than 1,000,000 are impossible in reality and indicate no limit.

Health-based chromium VI soil guideline	values (mg/kg)		Threshold									
Oral RHS (mg/kg BW/day)		0.003	Skin absorption	n factor		0						
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		0.0324						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		0.0324						
Background intake child (mg/kg BW/day)		No data	Bioconcentratio	on factor tube	r	0.0324						
Background intake adult (mg/kg BW/day)		No data	Mean bioconce	entration facto	r	0.0324						
Scenario	Soil ingestion	Dermal	Inhalation		Produ	ce ingesti	ion			Combin	ed SGV	
				No 10% 25 produce produce proc			5% duce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.00285	na	na		0	.00285						
RHS-adult background (mg/kg BW/day)	0.00285	na	na		0	.00285						
Rural residential / lifestyle block	773	na	na	na	1,136	4	54	227	773	460	286	176
Standard residential	773	na	na	na	1,136	4	54	227	773	460	286	176
High-density residential	1,546	na	na	na					1,546			
Parks / recreation	2,705	na	na						2,705			
Commercial / industrial indoor worker	NL	na	na						NL			
Commercial / industrial outdoor worker	6,332	na	na						6,332			

Note: There is no data on the background intake of CrVI. In accordance with MfE (2010b) the background intake is taken to be 5 per cent of the tolerable daily intake.

NL = No limit.

Health-based copper soil guideline value	s (mg/kg)		Threshold						
Oral RHS (mg/kg BW/day)		0.15	Skin absorption	n factor	0				
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf	na				
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root	na				
Background intake child (mg/kg BW/day)		0.056	Bioconcentratio	on factor tuber	na				
Background intake adult (mg/kg BW/day)		0.02	Max conc. in p	roduce (mg/kg DW)	30				
10% produce additional background intake		0.0024							
25% produce additional background intake		0.0061							
50% produce additional background intake		0.0121							
Scenario	Soil ingestion	Dermal	Inhalation			Combined SGV			
RHS-child background (mg/kg BW/day)	0.0940	na	na	No produce	10% pro	duce	25% produce	50% produce	
RHS-adult background (mg/kg BW/day)	0.1300	na	na						
Rural residential / lifestyle block	25,487	na	na	25,487	24,83	0	23,845	22,202	
Standard residential	25,487	na	na	25,487	24,83	0	23,845	22,202	
High-density residential	50,975	na	na	50,975					
Parks / recreation	89,206	na	na	89,206					
Commercial / industrial indoor worker	NL	NL	na	NL					
Commercial / industrial outdoor worker	288,826	na	na	288,826					

Note: BCF not used for produce pathway. Additional background intake subtracted from TDI on assumption that maximum plant tissue boron concentration = 300 mg/kg dry weight. NL = No limit.

Health-based inorganic lead soil guidelin	e values (mg/kg)		Threshold									
Oral RHS (mg/kg BW/day)		0.001 90	Skin absorptior	n factor		0						
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		0.019						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		0.015						
Background intake child (mg/kg BW/day)		0.00097	Bioconcentratio	on factor tube	r	0.005						
Background intake adult (mg/kg BW/day)		0.00041	Mean bioconce	entration facto	r	0.0102						
Scenario	Soil ingestion	Dermal	Inhalation		Produ	ce ingest	ion			Combin	ed SGV	
				No produce	10% produce	e proc	5% duce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.00093	na	na		0	.00093						
RHS-adult background (mg/kg BW/day)	0.00149	na	na		0	.00149						
Rural residential / lifestyle block	252	na	na	na	1,177	4	71	235	252	208	164	122
Standard residential	252	na	na	na	1,177	4	71	235	252	208	164	122
High-density residential	504	na	na	na					504			
Parks / recreation	883	na	na						883			
Commercial / industrial indoor worker	NL	na	na						NL			
Commercial / industrial outdoor worker	3,310	na	na						3,310			

Health-based inorganic mercury soil guid	deline values (mg/	′kg)	Threshold									
Oral RHS (mg/kg BW/day)		0.002	Skin absorption	n factor		0						
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		0.04						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		0.07						
Background intake child (mg/kg BW/day)		0.00005	Bioconcentratio	on factor tuber	r	0.02						
Background intake adult (mg/kg BW/day)		0.000065	Mean bioconce	entration facto	r	0.031						
Scenario	Soil ingestion	Dermal	Inhalation		Produ	ce ingest	ion			Combin	ed SGV	
				No 10% 25 produce produce proc			5% duce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.00190	na	na		C).00190						
RHS-adult background (mg/kg BW/day)	0.00190	na	na		C).00190						
Rural residential / lifestyle block	515	na	na	na	791	3	17	158	515	312	196	121
Standard residential	515	na	na	na	791	3	17	158	515	312	196	121
High-density residential	1,030	na	na	na					1,030			
Parks / recreation	1,803	na	na						1,803			
Commercial / industrial indoor worker	NL	na	na						NL			
Commercial / industrial outdoor worker	4,221	na	na						4,221			

Note: Background intake less than 5 per cent of the TDI, therefore background taken as 5 per cent of the TDI.

NL = No limit.

Health-based BaP soil guideline values (r	mg/kg)		Non-threshold	k								
Oral RHS (mg/kg BW/day)		0.000 004 8	Skin absorptior	n factor		0.06						
Dermal RHS (mg/kg BW/day)		na	Bioconcentratio	on factor leaf		0.005						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentratio	on factor root		0.031						
Background intake child (mg/kg BW/day)		na	Bioconcentratio	on factor tuber		0.004						
Background intake adult (mg/kg BW/day)		na	Mean bioconce	entration factor	r	0.007						
Scenario	Soil ingestion	Dermal	Inhalation		Produ	ce ingesti	on			Combin	ed SGV	
				No produce	10% produce	25 e proc	i% Juce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.0000048	0.0000048	na		0.	000048						
RHS-adult background (mg/kg BW/day)	0.0000048	0.0000048	na		0.	000048						
Rural residential / lifestyle block	12	121	na	na	34	1	4	7	11	8	6	4
Standard residential	13	140	na	na	48	1	9	10	12	10	7	5
High-density residential	26	279	na	na					24			
Parks / recreation	46	211	na						38			
Commercial / industrial indoor worker	NL	NL	na						NL			
Commercial / industrial outdoor worker	40	227	na						34			

Health-based \sum DDT soil guideline values	(mg/kg)		Threshold									
Oral RHS (mg/kg BW/day)		0.0005	Skin absorptio	n factor		0.018						
Dermal RHS (mg/kg BW/day)		na	Bioconcentrati	on factor leaf		0.012						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentrati	on factor root		0.038						
Background intake child (mg/kg BW/day)		0.0000511	Bioconcentrati	on factor tuber	r	0.038						
Background intake adult (mg/kg BW/day)		0.0000193	Mean bioconc	entration facto	r	0.0302						
Scenario	Soil ingestion	Dermal	Inhalation		Produ	ce ingest	tion			Combin	ed SGV	
				No produce	10% produce	e pro	5% duce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.0004489	0.0004489	na		0	.000449						
RHS-adult background (mg/kg BW/day)	0.0004750	0.0004750	na		0	.000475						
Rural residential / lifestyle block	122	4,449	na	na	192	7	77	38	118	73	47	29
Standard residential	122	4,449	na	na	192	7	77	38	118	73	47	29
High-density residential	243	8,897	na	na					237			
Parks / recreation	426	7,785	na						404			
Commercial / industrial indoor worker	NL	NL	na						NL			
Commercial / industrial outdoor worker	1,055	19,969	na						1,002			

Health-based dieldrin soil guideline value	es (mg/kg)		Threshold									
Oral RHS (mg/kg BW/day)		0.0001	Skin absorpti	on factor		0.1						
Dermal RHS (mg/kg BW/day)		na	Bioconcentra	tion factor leaf		0.41						
Inhalation RHS (mg/kg BW/day)		na	Bioconcentra	tion factor root		0.41						
Background intake child (mg/kg BW/day)		0.0000036	Bioconcentra	tion factor tube	er	0.41						
Background intake adult (mg/kg BW/day)		0.0000014	Mean biocon	centration facto	or	0.41						
Scenario	Soil ingestion	Dermal	Inhalation		Produce	e ingestior	n			Combin	ed SGV	
				No produce	10% produce	25% produ	ice	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.000095	0.000095	na		0.0	00095						
RHS-adult background (mg/kg BW/day)	0.000095	0.000095	na		0.0	00095						
Rural residential / lifestyle block	26	169	na	na	2.99	1.20)	0.60	22	2.6	1.1	0.58
Standard residential	26	169	na	na	2.99	1.20)	0.60	22	2.6	1.1	0.58
High-density residential	52	339	na	na					45			
Parks / recreation	90	297	na						69			
Commercial / industrial indoor worker	NL	NL	na						NL			
Commercial / industrial outdoor worker	211	719	na						163			

Note: Background intake less than 5 percent of the TDI, therefore background taken as 5 per cent of the TDI.

NL = No limit.

Health-based dioxin (TCDD/PeCDD domin (µg TEQ/kg)	nant) soil guidelir	e values	Threshold									
Oral RHS (μg/kg BW/day)		0.000 001	Skin absorp	tion factor		0.02						
Dermal RHS (µg/kg BW/day)		na	Bioconcentr	ation factor leat	f	0						
Inhalation RHS (µg/kg BW/day)		na	Bioconcentr	ation factor roo	t	0						
Background intake child (µg/kg BW/day)		0.000 000 33	Bioconcentr	ation factor tub	er	0						
Background intake adult (µg/kg BW/day)		0.000 000 33	Bioaccumula	ation factor cuc	urbits	0.24						
			Mean bioco	ncentration fact	or	0.0096						
Scenario	Soil ingestion	Dermal	Inhalation		Produc	e ingestio	n			Combir	ed SGV	
				No produce	25% produ	% uce	50% produce	No produce	10% produce	25% produce	50% produce	
RHS-child background (µg/kg BW/day)	0.00000067	0.00000067	na		0.00	0000067						
RHS-adult background (µg/kg BW/day)	0.0000067	0.0000067	na		0.00	0000067						
Rural residential / lifestyle block	0.182	6.0	na	na	0.90	0.3	6	0.18	0.18	0.15	0.12	0.09
Standard residential	0.182	6.0	na	na	0.90	0.3	6	0.18	0.18	0.15	0.12	0.09
High-density residential	0.363	12	na	na					0.35			
Parks / recreation	0.64	10	na						0.60			
Commercial / industrial indoor worker	NL	NL	na						NL			
Commercial / industrial outdoor worker	1.49	25	na						1.4			

Health-based dioxin (OCDD/HpCDD domi (µg TEQ/kg)	nant) soil guideli	ne values	Threshold								
Oral RHS (μg/kg BW/day)		0.000 001	Skin absorpt	tion factor		0.02					
Dermal RHS (µg/kg BW/day)		na	Bioconcentra	ation factor lea	af	0					
Inhalation RHS (µg/kg BW/day)		na	Bioconcentra	ation factor ro	ot	0					
Background intake child (µg/kg BW/day)		0.000 000 33	Bioconcentra	ation factor tul	ber	0					
Background intake adult (µg/kg BW/day)		0.000 000 33	Bioaccumula	ation factor cu	curbits	0.017					
			Mean biocor	ncentration fac	ctor	0.00068					
Scenario	Soil ingestion	Dermal	Inhalation	nhalation Produce ingesti					Combir	ned SGV	
				No 10% 25% produce produce produce			50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (µg/kg BW/day)	0.0000067	0.00000067	na		0.000	000067					
RHS-adult background (µg/kg BW/day)	0.0000067	0.0000067	na		0.000	000067					
Rural residential / lifestyle block	0.182	6.0	na	na	12.7	5.09	2.54	0.18	0.17	0.17	0.16
Standard residential	0.182	6.0	na	na	12.7	5.09	2.54	0.18	0.17	0.17	0.16
High-density residential	0.363	12	na	na				0.35			
Parks / recreation	0.64	10	na					0.60			
Commercial / industrial indoor worker	NL	NL	na					NL			
Commercial / industrial outdoor worker	1.49	25	na					1.4			

Health-based dioxin-like PCB soil guidel	ine values (µg TE	Q/kg)	Threshold						
Oral RHS (μg/kg BW/day)		0.000 001	Skin absorpt	ion factor		0.07			
Dermal RHS (µg/kg BW/day)		na	Bioconcentration factor leaf			0			
Inhalation RHS (µg/kg BW/day)		na	Bioconcentra	ation factor ro	ot	0			
Background intake child (µg/kg BW/day)		0.000 000 33	Bioconcentra	ation factor tul	ber	0			
Background intake adult (µg/kg BW/day)		0.000 000 33	Bioaccumula	ation factor cu	curbits	0.45 0.018			
			Mean biocor	ncentration fac	ctor	0.018			
Scenario	Soil ingestion	Dermal	Inhalation		Produce	ingestion			
				No produce	10% produce	25% produce	50% produce		
RHS-child background (μg/kg BW/day)	0.0000067	0.0000067	na		0.000	000067			
RHS-adult background (µg/kg BW/day)	0.00000067	0.00000067	na		0.000	0000067			
	1					1	1		

Scenario	Soil ingestion	Dermal	Inhalation		Produce	ingestion		Combined SGV			
				No produce	10% produce	25% produce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (µg/kg BW/day)	0.0000067	0.0000067	na		0.000	00067					
RHS-adult background (µg/kg BW/day)	0.0000067	0.0000067	na		0.000	00067					
Rural residential / lifestyle block	0.182	1.7	na	na	0.481	0.192	0.096	0.16	0.12	0.09	0.06
Standard residential	0.182	1.7	na	na	0.481	0.192	0.096	0.16	0.12	0.09	0.06
High-density residential	0.363	3.4	na	na				0.33			
Parks / recreation	0.64	3.0	na					0.52			
Commercial / industrial indoor worker	NL	NL	na					NL			
Commercial / industrial outdoor worker	1.49	7.2	na					1.2			

Health-based pentachlorophenol soil guideline values (mg/kg)			Threshold								
Oral RHS (mg/kg BW/day) 0.0003		Skin absorption factor			0.24						
Dermal RHS (mg/kg BW/day) na		Bioconcentration factor leaf			0						
Inhalation RHS (mg/kg BW/day) na		na	Bioconcentration factor root			0					
Background intake child (mg/kg BW/day)		0.00002	Bioconcentration factor tuber 0								
Background intake adult (mg/kg BW/day)		0.00002	Mean bioconcentration factor na			na					
Scenario	Soil ingestion	Dermal	Inhalation	Produce ingestion				Combined SGV			
				No produce	10% produce	25% produce	50% produce	No produce	10% produce	25% produce	50% produce
RHS-child background (mg/kg BW/day)	0.00028	0.00028	na		0.0	0028					
RHS-adult background (mg/kg BW/day)	0.00028	0.00028	na		0.0	0028					
Rural residential / lifestyle block	76	208	na	na na n			na	56	56	56	56
Standard residential	76	208	na	na	na	na	na	56	56	56	56
High-density residential	152	416	na	na				111			
Parks / recreation	266	364	na					154			
Commercial / industrial indoor worker	NL	NL	na					NL			
Commercial / industrial outdoor worker	622	883	na					365			

Appendix 2: Detailed SGV_(health) Calculations for Cadmium

The detailed calculations presented here are for residential SGVs for cadmium over the soil pH range 5 to 7, in increments of 0.5. A plot of the variation of SGV with soil pH is shown in figure A2.1.

SGVs for soil pH above 7 or below 5 have not been calculated because of the unknown validity of the pH - BCF relationship above pH 7 or below pH 5. A lack of plant uptake data prevented the relationship being extended beyond the range pH 5 to 7.

The combined $SGVs_{(health)}$ shown in the following calculations are not rounded in accordance with the SCS rounding rules (see beginning of section 6). Rounded values over the pH range are given in the table for cadmium given in appendix A.

Residential soil SGVs for other than the default soil pH of 5 should not be used except for sitespecific assessments, and then only where the soil pH can be reliably determined. Determination of soil pH must include:

- samples taken of each soil type, based on field observations of such things as texture, organic content, moisture content and mineralogy different SGVs may apply for different soil types depending on the pH of each soil type
- a large enough number of samples of each soil type for the determination to be statistically valid
- avoidance of areas where the pH may have been artificially modified, including avoiding gardens and agricultural land that may have been fertilised and/or limed, as the soil pH of such land is likely to revert back to the natural pH with time. This may mean sampling nearby undeveloped land of the same soil type(s), bush land, road verges or other areas that are less likely to have had the soil pH artificially altered
- samples are to be analysed using an IANZ accredited laboratory.

In applying these SGVs, the nearest pH below the measured pH should be selected. Interpolating between SGVs for other pH is not warranted, given the various approximations involved in the derivation of the SGVs.

Health-based cadmium	Threshold										
Oral RHS (mg/kg BW/day	0.000 833	Homegrown produce %			10%	25	% 50%				
Dermal RHS (mg/kg BW/	na	Bioconcentr	r leaf	3.26	4.4	7 5.71					
Inhalation RHS (mg/kg B	W/day)			na	Bioconcentr	ation facto	r root	1.10	1.8	6 2.79	
Background exposure ch	0.000 41	Bioconcentr	ation facto	r tuber	1.10	1.8	6 2.79				
Background exposure ad	lult (mg/kg BV	V/day)		0.000 26	Mean biocor	ncentratior	n factor	1.75	2.6	4 3.66	
Skin absorption factor				0.001							
Scenario	Soil	Dermal	Pro	duce inges	stion		Combi	ined SGV			
ingestion			10% produce	25% produce	50% produce	No produce	10% produce	25% produce		50% produce	
RHS-child background (mg/kg BW/day)	0.00042	0.00042		0.00042							
RHS-adult background (mg/kg BW/day)	0.00057	0.00057		0.00057							
Rural residential / lifestyle block	115	75,456	3.12	0.83	0.30	115	3.0	0.82		0.30	
Standard residential	115	75,456	3.12	0.83	0.30	115	3.0	0.82		0.30	
High-density residential	229	150,912				229					
Parks / recreation	401	132,048				400					
Commercial / industrial indoor worker	NL	NL				NL					
Commercial / industrial outdoor worker	1,273	433,602				1,269					

na = Not applicable.

Health-based cadmium	5.5	Threshold									
Oral RHS (mg/kg BW/day	0.000 833	Homegrown	Homegrown produce %			25%	6 50%				
Dermal RHS (mg/kg BW/	na	Bioconcentr	ation facto	2.12	2.8	8 3.67					
Inhalation RHS (mg/kg B	na	Bioconcentr	ation facto	r root	0.59	0.9	9 1.48				
Background exposure ch	0.00041	Bioconcentr	ation facto	r tuber	0.59	0.9	9 1.48				
Background exposure ad	0.00026	Mean biocor	n factor	1.05	1.5	6 2.14					
Skin absorption factor				0.001							
Scenario	Soil	Dermal	Pro	duce inges	stion		ined SGV				
ingestion		10% produce	25% produce	50% produce	No produce	10% produce	259 prod	% uce	50% produce		
RHS-child background (mg/kg BW/day)	0.00042	0.00042		0.00042							
RHS-adult background (mg/kg BW/day)	0.00057	0.00057		0.00057							
Rural residential / lifestyle block	115	75,456	5.19	1.40	0.51	115	5.0	1.4		0.51	
Standard residential	115	75,456	5.19	1.40	0.51	115	5.0	1.4	4	0.51	
High-density residential	229	150,912				229					
Parks / recreation	401	132,048				400					
Commercial / industrial indoor worker	NL	NL				NL					
Commercial / industrial outdoor worker	1,273	433,602				1,269					

NL = No limit.
Health-based cadmium	soil guidelir	/kg) at pH (6	Threshold							
Oral RHS (mg/kg BW/da	y)			0.000 833	Homegrown	produce 9	%	10%	25	% 50%	
Dermal RHS (mg/kg BW/	/day)			na	Bioconcentra	ation facto	r leaf	1.38	1.8	2.36	
Inhalation RHS (mg/kg B	W/day)			na	Bioconcentr	ation facto	r root	0.32	0.5	0.79	
Background exposure ch	ild (mg/kg BV	V/day)		0.00041	Bioconcentr	ation facto	r tuber	0.32	0.5	0.79	
Background exposure ac	lult (mg/kg BV	0.00026	Mean biocor	ncentratior	n factor	0.64	0.9	3 1.26			
Skin absorption factor				0.001							
Scenario	Soil Dermal Product					duce ingestion					
	ingestion		10% produce	25% produce	50% produce	No produce	10% produce	259 prod	% uce	50% produce	
RHS-child background (mg/kg BW/day)	0.00042	0.00042		0.00042							
RHS-adult background (mg/kg BW/day)	0.00057	0.00057		0.00057							
Rural residential / lifestyle block	115	75,456	8.50	2.34	0.87 115 7.9		115 7.9 2.3		2.3 0.8		
Standard residential	115	75,456	8.50	2.34	0.87	115	7.9	2.3	3	0.86	
High-density residential	229	150,912				229					
Parks / recreation	401	132,048				400					
Commercial / industrial indoor worker	NL	NL				NL					
Commercial / industrial outdoor worker	1,273	433,602				1,269					

NL = No limit.

na = Not applicable.

Health-based cadmium	g/kg) at pH	6.5	Threshold								
Oral RHS (mg/kg BW/da	y)			0.000 833	Homegrown	produce 9	%	10%	25	% 50%	%
Dermal RHS (mg/kg BW/	/day)			na	Bioconcentra	ation facto	r leaf	0.91	1.2	1 1.5	53
Inhalation RHS (mg/kg B	W/day)			na	Bioconcentr	ation facto	r root	0.18	0.2	9 0.4	2
Background exposure ch	ild (mg/kg BV	V/day)		0.00041	Bioconcentra	ation facto	r tuber	0.18	0.2	9 0.4	2
Background exposure ac	lult (mg/kg BV	V/day)		0.00026	Mean biocor	ncentratior	n factor	0.40	0.5	7 0.7	'5
Skin absorption factor		0.001									
Scenario	Soil	Dermal	Pro	duce inges	stion		Combi	bined SGV			
	ingestion		10% produce	25% produce	50% produce	No produce	10% produce	25% prod	% uce	50% produc	ce
RHS-child background (mg/kg BW/day)	0.00042	0.00042		0.00042							
RHS-adult background (mg/kg BW/day)	0.00057	0.00057		0.00057							
Rural residential / lifestyle block	115	75,456	13.7	13.7 3.86 1.45 115 12 3.		1.45 115 12		3.7	7	1.4	
Standard residential	115	75,456	13.7	3.86	1.45	115	12	3.7	7	1.4	
High-density residential	229	150,912				229					
Parks / recreation	401	132,048				400					
Commercial / industrial indoor worker	NL	NL				NL					
Commercial / industrial outdoor worker	1,273	433,602	1,269								

NL = No limit.

Health-based cadmium	soil guidelir	/kg) at pH 7	7	Threshold							
Oral RHS (mg/kg BW/day	y)			0.000 833	Homegrown	produce 9	%	10%	25	% 50%	6
Dermal RHS (mg/kg BW/	/day)			na	Bioconcentr	ation facto	r leaf	0.61	0.7	9 0.99	Э
Inhalation RHS (mg/kg B	W/day)			na	Bioconcentr	ation facto	r root	0.10	0.1	6 0.23	3
Background exposure ch	ild (mg/kg BV	V/day)		0.00041	Bioconcentr	ation facto	r tuber	0.10	0.1	6 0.23	3
Background exposure ad	lult (mg/kg BV	V/day)		0.00026	Mean biocor	ncentration	n factor	0.25	0.3	85 0.46	3
Skin absorption factor				0.001							
Scenario	Soil	Dermal	Pro	duce inges	stion		Combi	bined SGV			
	ingestion		10% produce	25% produce	50% produce	No produce	10% produce	25° prod	% uce	50% produce	e
RHS-child background (mg/kg BW/day)	0.00042	0.00042		0.00042							
RHS-adult background (mg/kg BW/day)	0.00057	0.00057		0.00057							
Rural residential / lifestyle block	115	75,456	21.7	6.28	2.38	115	18	6.	C	2.3	
Standard residential	115	75,456	21.7	6.28	2.38	115	18	6.	C	2.3	
High-density residential	229	150,912				229					
Parks / recreation	401	132,048				400					
Commercial / industrial indoor worker	NL	NL				NL					
Commercial / industrial outdoor worker	1,273	433,602				1,269					

NL = No limit.

na = Not applicable.

Figure A2.1: Dependence of cadmium soil guideline value on pH



Appendix 3: Determination of the Amount of Produce Consumed and its Dry Weight

Estimating the intake of soil contaminants via the home-grown produce pathway requires as a basic parameter an estimate of the average child and adult's produce consumption. The estimate presented here is slightly modified from appendix 2 of Cavanagh (2005b), which in turn was a modification from Cavanagh and Proffitt (2005). The major modification from Cavanagh (2005b) is dividing vegetable types into root, tuber and above-ground (including cucurbits), rather than just below-ground and above-ground vegetables.

Typically, only the consumption of vegetables is taken into account in the derivation of soil guideline values (eg, the Netherlands, Canada), although the UK also takes fruit into account. In New Zealand, consideration will be given only to the consumption of vegetables. This approach is valid for two reasons. Firstly, fruit is not widely grown in residential gardens. Secondly, contaminant uptake and translocation to fruit (in fruit trees) is considered to be negligible (MfE and MoH, 1997; MfE, 1999).

For calculation purposes, table A3.1 shows the amount of produce consumed by different age and gender groups based on simulated diets (from Brinsdon, 2004). Quantities of fruit, vegetables and totals have been calculated for the average adult and the average child.

Table A3.1: Amount of fruit and vegetables consumed (grams per day) by differe	nt age-
gender groups based on simulated diets for the New Zealand 2003/0	4 Total
Diet Surveys	
-	

Produce	Young male 19–24 years	Male 25+ years	Female 25+ years	Average adult ¹	Child 5–6 years	Child 1–3 years	Child 6–12 months	Average child ²
Vegetables	224	294	232	254	115	63	42.5	77
Fruit	110	138	141	136	140	77	62.5	95
Total	336	432	373	390	256	140	105	173
Body weight ³	70	80	65	70	20	13	10	15

Source: Adapted from Brinsdon (2004).

1 Adjusted for difference in body weight before averaging and converted to the amount consumed by a 70-kg adult.

2 Adjusted for difference in body weight before averaging and converted to the amount consumed by a 15-kg child.

3 Vannoort et al, 2000.

Vegetable types

The extent of uptake of contaminants by different vegetables is dependent on the individual vegetable. In the development of soil guideline values, vegetables are most frequently separated into root (eg, carrots, potato) and above-ground (eg, lettuce, peas) vegetables. The vegetables considered in the development of the simulated diets were grouped into root vegetables, tubers, 'above-ground' vegetables (including the cucurbits), and vegetables unlikely to be grown at home, as shown in table A3.2.

'Above-ground' vegetables	Root vegetables	Unlikely to be grown at home
Bean	Carrot	Avocado
Broccoli / cauliflower	Kumara (tuber)	Celery
Cabbage	Potato (tuber)	Mushrooms
Capsicum		
Courgette (cucurbit)		
Cucumber (cucurbit)		
Lettuce		
Onion		
Peas		
Pumpkin (cucurbit)		
Silverbeet		
Tomato		

Table A3.2: Vegetables considered in simulated diets, and their grouping

Dry weight contents of individual vegetables were determined from the average of the values provided in US EPA (1996b; 1997) and Alloway et al (1988). The exception was avocado, whose dry weight content was derived from the average dry weight of the edible portion of avocado over a growing season (Hofshi et al, 2000).

Table A3.4 provides a summary of the amount of vegetables consumed by an average adult and child, using the above grouping. Plant uptake of contaminants is typically expressed on a dry-weight basis, and therefore requires conversion from the wet weights typically used to express the amount of produce consumed. However, the dry-weight content of different vegetables is also variable. The dry weight proportion of individual vegetables (table A3.3) and the amount of each vegetable consumed was used to determine the amount of produce consumed on a dry-weight basis (table A3.4). This also enables derivation of consumption-weighted dry weights for the different types of vegetables, and for different receptors. The consumption-weighted dry-weight content does not vary much for the different receptors (table A3.4), with root vegetables having an average dry-weight content of about 0.18, and above-ground vegetables of 0.09.

Vegetable	Conversion factor
Avocado	0.25
Beans	0.094
Broccoli / cauliflower	0.089
Cabbage	0.083
Capsicum	0.15
Carrot	0.110
Celery	0.116
Courgette	0.035
Cucumber	0.079
Kumara	0.21
Lettuce	0.046
Mushrooms	0.082
Onion	0.124
Peas	0.145
Potatoes	0.210
Potatoes, with skin	0.167
Pumpkin	0.084
Silverbeet (spinach)	0.073
Tomato	0.058

Table A3.3: Fresh to dry conversion factors for vegetables

Produce type	Wet weight perce	(g/day) and ntage	Dry weight perce	(g/day) and ntage	Consumption- weighted dry-weight content (g DW/g FW)			
	Average adult	Average child	Average adult	Average child	Adult	Child		
Tuber vegetable	92 (36%)	33 (43%)	18.9 (56%)	6.6 (63%)	0.21	0.20		
Root vegetables	18 (7%)	9 (12%)	1.9 (6%)	1.0 (10%)	0.11	0.11		
Above-ground vegetables (not including cucurbits)	119 (47%)	25 (33%)	10 (30%)	2.4 (23%)	0.08	0.10		
Cucurbits (courgette, pumpkin)	14 (6%)	6 (8%)	1.4 (4%)	0.46 (4%)	0.10	0.08		
Subtotal	243	73	32.2	10.5	0.13	0.14		
Unlikely to be grown at home	10 (4%)	3.6 (5%)	1.4 (4%)	0.57 (5%)	-	-		
Total	253	76.6	33.6	11.03	0.13	0.14		

Table A3.4: Amounts of different vegetable groups consumed by an average adult and average child, with percentage of total vegetables given in parenthesis

Source: Adapted from Brinsdon (2004).

Appendix 4: Dioxin SGV_(health) Calculations with Egg Consumption

Health-based dioxin (TCDD/PeCDD-dominant) soil guideline value (µg TEQ/kg) Threshold																		
Oral RHS (µg/kg BW/day)			0.000 001		Sk	in absorptio	n factor		0.02									
Dermal RHS (µg/kg BW/d	ay)		na		Bio	oconcentrati	on factor leaf		0									
Inhalation RHS (µg/kg BW	//day)		na		Bio	oconcentrati	on factor root		0									
Background exposure chil	d (µg/kg BW/da	iy)	0.000 000 3	33	Bio	oconcentrati	on factor tuber		0									
Background exposure adu	ılt (µg/kg BW/da	ay)	0.000 000 3	33	Bio	oconcentrati	on factor cucu	bits	0.24									
Egg lipid BCF			1.9		Me	an bioconce	entration factor		0.0096									
Scenario	Soil ingestion	Dermal	Inhalation	Pro	duce inges	gestion Average 2 x a egg ingestion ing			average egg	Com	bined SGV n	o eggs	Combined SGV average eggs			Combined SGV with 2 x average eggs		
				No produce	10% produce	25% produce	ingestion	ing	estion	No produce	10% produce	25% produce	No produce	10% produce	25% produce	No produce	10% produce	25% produce
RHS-child background (µg/kg BW/day)	0.00000067	0.00000067	na		0.0000006	7	0.00000067	0.00	000067									
RHS-adult background (µg/kg BW/day)	0.00000067	0.00000067	na		0.0000006	7	0.00000067	0.00	000067									
Rural residential / lifestyle block	0.182	6.0	na	na	0.901	0.360	0.005	0	0.003	0.176	0.147	0.118	0.005	0.005	0.005	0.003	0.003	0.003
Standard residential	0.182	6.0	na	na	0.901	0.360	0.005	0	0.003	0.176	0.147	0.118	0.005	0.005	0.005	0.003	0.003	0.003
High-density residential	0.363	12	na	na						0.353			0.353			0.353		
Parks / recreation	0.636	10	na							0.599			0.599			0.599		
Commercial / industrial indoor worker	NL	NL	na							NL			NL			NL		
Commercial / industrial outdoor worker	1.49	25	na							1.41			1.41			1.41		

NL = No limit.

Health-based dioxin (OCI	DD/HpCDD-doi	minant) soil g	uideline val	ue (µg TEC	ג(kg) Th	reshold													
Oral RHS (µg/kg BW/day)			0.000 001		Sk	in absorptio	n factor	0.02											
Dermal RHS (µg/kg BW/da	iy)		na		Bio	oconcentrati	on factor leaf	0											
Inhalation RHS (µg/kg BW/	/day)		na		Bio	oconcentrati	on factor root	0											
Background exposure child	d (µg/kg BW/da	y)	0.000 000 3	33	Bio	oconcentrati	on factor tuber	0											
Background exposure adul	t (µg/kg BW/da	y)	0.000 000 3	33	Bio	oconcentrati	on factor cucur	bits 0.017											
Egg lipid BCF			0.7		Me	an bioconc	entration factor	0.00068	3										
Scenario	Soil ingestion	Dermal	Inhalation	Pro	duce inge	stion	Average egg ingestion	2 x average egg	Com	bined SGV n	Combine	d SGV ave	rage eggs	S Combined SGV with 2 x average eggs					
				No produce	10% produce	25% produce		ingestion	No produc	10% produce	25% produce	No produce	10% produce	25% produce	No produce	10% produce	25% produce		
RHS-child background (µg/kg BW/day)	0.00000067	0.0000067	na		0.0000006	7	0.00000067	0.0000067											
RHS-adult background (µg/kg BW/day)	0.00000067	0.0000067	na		0.0000006	7	0.00000067	0.00000067											
Rural residential / lifestyle block	0.182	6.0	na	na	12.7	5.1	0.014	0.007	0.176	0.174	0.170	0.013	0.013	0.013	0.007	0.007	0.007		
Standard residential	0.182	6.0	na	na	12.7	5.1	0.014	0.007	0.176	0.174	0.170	0.013	0.013	0.013	0.007	0.007	0.007		
High-density residential	0.363	12	na	na					0.353			0.353			0.353				
Parks / recreation	0.636	10	na						0.599			0.599			0.599				
Commercial / industrial indoor worker	NL	NL	na						NL			NL			NL				
Commercial / industrial outdoor worker	1.49	25	na						1.41			1.41			1.41				

NL = No limit.

Health-based dioxin-like	PCBs soil gui	deline value (µ	ıg TEQ/kg)		Tł	reshold												
Oral RHS (µg/kg BW/day) Dermal RHS (µg/kg BW/da Inhalation RHS (µg/kg BW Background exposure chile Background exposure adu Egg lipid BCF	ay) /day) d (µg/kg BW/da It (µg/kg BW/da	y) ay)	0.000 001 na na 0.000 000 0.000 000 17	33 33	Sk Bi Bi Bi Bi	in absorptic oconcentrat oconcentrat oconcentrat oconcentrat oconcentrat	on factor ion factor leaf ion factor root ion factor tuber ion factor cucu centration facto	r rbits r	0.07 0 0 0.45 0.018									
Scenario	Soil ingestion	Dermal	Inhalatio n	Pro	duce inge	stion	Average egg	2 x a	average egg	Comb	oined SGV n	o eggs	Combine	d SGV ave	rage eggs	Comb	ined SGV v iverage egg	vith 2 x js
				No produce	10% produce	25% produce	ingestion	ing	jestion	No produce	10% produce	25% produce	No produce	10% produce	25% produce	No produce	10% produce	25% produce
RHS-child background (µg/kg BW/day)	0.0000067	0.0000067	na		0.0000006	7	0.0000067	0.00	0000067									
RHS-adult background (µg/kg BW/day)	0.0000067	0.0000067	na		0.0000006	7	0.0000067	0.00	0000067									
Rural residential / lifestyle block	0.182	1.7	na	na	0.48	0.19	0.0006	0	.0003	0.164	0.122	0.089	0.0006	0.0006	0.0006	0.0003	0.0003	0.0003
Standard residential	0.182	1.7	na	na	0.48	0.19	0.0006	0	.0003	0.164	0.122	0.089	0.0006	0.0006	0.0006	0.0003	0.0003	0.0003
High-density residential	0.363	3.4	na	na						0.328			0.328			0.328		
Parks / recreation	0.636	3.0	na							0.524			0.524			0.524		
Commercial / industrial indoor worker	NL	NL	na							NL			NL			NL		
Commercial / industrial outdoor worker	1.49	7.2	na							1.23			1.23			1.23		

NL = No limit.

Appendix 5: International Comparison of Soil Guideline Value Derivation

A5.1 Introduction

New Zealand has four industry-based guideline documents providing soil guideline values for a variety of contaminants (MfE and MoH, 1997; MfE, 1997, 1999, 2006a). The United States risk assessment procedures set out in US EPA (1989a) provided the basis for the derivation of the soil guideline values in these documents. In addition, New Zealand has typically looked to the United States and a number of other countries, in particular Australia, Canada, the Netherlands and the United Kingdom, to provide soil guideline values where New Zealand guidelines do not exist. A compilation of guidelines from these countries formed Contaminated Land Management Guideline No. 2 (MfE, 2003). Each of these overseas countries has well-developed contaminated sites risk assessment frameworks, with guideline documents readily available, including Dutch documents available in English.

This section explores the derivation methods and differences of the New Zealand documents and documents from the five overseas jurisdictions as background to proposing a consistent methodology for New Zealand, drawing on and updating earlier work for the Ministry for the Environment by Cavanagh (2003) and Cavanagh and O'Halloran (2003).

Given that the US has provided the basis for New Zealand's guideline, the US EPA guidance is considered first, then the other overseas jurisdictions, and finally the existing New Zealand guidelines.

A5.2 Regulatory context

The derivation of human-health numeric values largely focuses on the exposure of a given human receptor in designated scenarios that typically relate to different land uses. But the various choices in soil guideline derivation, including the selection of designated scenarios and exposure parameters, are influenced by the particular policy and legislation of each country. The various legislative and policy environments overseas may not (and frequently do not) accord with the New Zealand situation and therefore many overseas choices do not suit New Zealand.

Carlon (2007) recently compared the reasons for the differences in SGVss for 12 European countries. While only two of the countries studied are considered here (the Netherlands and the UK), his findings are generally applicable. Carlon identified five, often overlapping, categories of reasons for differences in SGVs:

- 1. **Geographical and biological**: associated with the environmental variability between countries (across Europe in his case).
- 2. **Socio-cultural**: associated with the variability of social behaviours and land use between countries.
- 3. **Regulatory**: associated with regulatory requirements, such as constitutional aspects or commonalities with existing laws.

- 4. **Political**: associated with the prioritisation of environmental and economic values, as made by policy makers and regulators, or forced by societal views.
- 5. Scientific: associated with arguments of different scientific views.

This document largely focuses on the last of these, with some input from socio-cultural aspects such as the New Zealand lifestyle (eg, ownership of rural 'lifestyle' properties and vegetable growing habits). However, the lack of New Zealand-specific data does not always allow proper accounting of how New Zealand's lifestyle and work habits differ from other countries.

Geographical and biological differences have a limited role in New Zealand. Its climate is sufficiently consistent across the country that the large differences in summer and winter lifestyles forced by heavy winter snows in some countries do not have to be taken into account in New Zealand guidelines. On the other hand, the lack of good New Zealand data and/or the ready accessibility of overseas data (particularly from the US) means that New Zealand typically draws on overseas data to describe common activities and the physical characteristics of the 'average' person.

This document has attempted to avoid 'importing' regulatory or policy decisions from other countries in making recommendations, other than the inevitable 'scientific' policy embedded in the relatively generic risk assessment frameworks common to most countries. An example of country-specific policies is Canada choosing to assign only one-fifth of the acceptable daily intake of a substance to the soil compartment, dividing the remainder amongst air, water, consumer products and food. There is no scientific justification for this – it is purely a policy decision.

The differences in approach can make large difference in final soil guideline values. In his analysis of values across the European Union, Carlon (2007) found a difference of one to two orders of magnitude between values for particular metals and metalloids; and an even larger range of two to three orders of magnitude for some organic compounds commonly found on contaminated sites.

Two important policy issues require mentioning at this point: the purpose to which soil guideline values are to be put, and whether the values are to protect human health only, or human and ecological receptors.

Soil guideline values are generally derived for one of the following purposes:

- investigation if the stated value is exceeded, further site investigation is required
- intervention if the stated value is exceeded, remediation (which may include management actions or imposition of a barrier to contact) or assessment of the urgency of remediation, is required. Typically an intervention value is higher than an investigation value
- remediation goals soil concentrations to which contaminated soil should be remediated or some other management action taken to minimise exposure. Such values may be higher than the equivalent intervention value.

Carlon (2007) set this down in diagrammatic form, as shown in figure A5.1. This shows that for there to be a negligible risk to all receptors, very low soil values have to be set (called target values in the figure). These tend to be aspirational, rather than realistic goals, and are not very useful for assessing whether a contaminated site is safe for its present use (with or without management) or whether further assessment is necessary. By their very nature, many sites would fail on the basis of screening using target values; using such values as remediation goals would result in many sites requiring expensive clean-ups with limited, if any, overall humanhealth benefit.



Figure A5.1: Derivation of screening values based on different risk levels and applications

Source: after Carlon, 2007.

In the middle of the continuum are values that provide a warning of potential risk – the orange light to use the traffic light analogy. Measured values below the designated screening value require no further action while concentrations above the value require further investigation. Such values are often called 'trigger', 'screening' or 'investigation' values. Exceeding such a value does not necessarily indicate a risk – particularly as most sites will not perfectly match the conservative generic assumptions behind the guideline value – but there is sufficient potential for risk that further enquiries should be made.

Screening values can move up or down the concentration axis depending on how conservative the policy and regulatory authorities wish to be (influenced by the various socio-cultural and other contexts). Lower values are more protective but potentially a greater unnecessary economic burden on a society, while higher values will be less protective, with the potential for some sites that actually present a risk slipping through the net.

The highest values are remediation or 'intervention' values. Such values indicate a risk is a distinct possibility and some form of intervention to moderate that risk is required. Site-specific assessment could result in even higher values being set, based on the actual contaminant exposure assessed for the site.

Most countries use more or less conservative values in the middle as screening values, as shown in table A5.1 (modified from Cavanagh, 2003). The art in setting such values is having them high enough to not trigger too much unnecessary further investigation, but not so high as to result in a risk for some sites. The latter is generally unlikely, because the toxicological values generally have large factors of safety given the uncertainty of the science. Not setting them too low will also result in less remediation and greater ongoing use of marginally contaminated sites: screening values are inevitably used as clean-up goals, either because a small site is not worth investigating further or because of regulatory expediency. The burden of unnecessarily low screening levels will typically fall on the small property owner; owners of larger, more valuable sites will more often have the resources to investigate further and possibly avoid remediation.

For this report, the purposes of soil guideline values, as enunciated in the MfE position paper (MfE, 2007), are to:

- serve as Tier 1 or screening criteria to assess whether there is a potential risk to human health
- when the criteria are exceeded, serve as conservative clean-up targets for many situations, ie, where further investigation or site-specific risk assessment is not warranted or economic
- inform onsite management actions to reduce the potential for adverse effects
- trigger further investigation to determine site-specific criteria.

The soil guidelines values represent 'clean down to levels' at contaminated sites and not 'pollute up to levels' for less contaminated sites, that is, the guideline values must not be considered as permission to contaminate up to a certain level. They are also not intended to be used to manage pristine sites (CCME, 2006).

Table A5.1: Summary of the name / terminology used, purpose, and basis for derivation of soil guideline values in different countries

Country		Terminology used in country / by agency	Purpose	Derivation basis		
New	Timber treatment	Soil acceptance criteria	Investigation	Human health / phytotoxicity		
Zealand	Gasworks	Soil acceptance criteria	Investigation	Human health		
	Oil industry	Soil acceptance criteria	Investigation	Human health		
	Sheep-dip sites	Soil guideline values	Investigation	Human health		
Australia		Investigation levels	Investigation	Ecological Human health		
United States	Federal Soil Screening Guidance	Soil screening level (SSL)	Investigation	Ecological Human health		
	US EPA Region 6	Media-specific human health screening levels	Investigation	Human health		
	US EPA Region 9	Preliminary remediation goals	Investigation	Human health		
Canada		Soil quality guidelines	Remediation goal	Integrated		
United Kin	gdom	Soil guideline values	Investigation	Human health		
The Nethe	erlands	Intervention value	Intervention	Integrated		
		Remediation values	Remediation goal			
		Target value	Long-term remediation goal	Ecological		

A largely philosophical decision also needs to be made as to whether soil numeric values should be protective of ecological or human receptors. In some countries, one or the other is considered; in other countries, both types of receptors are considered together to create 'integrated guidelines', eg, in Canada and the Netherlands. Some New Zealand guidelines, eg, some of the values in the 'Timber Treatment Guidelines' (MfE and MoH, 1997) have mixed consideration of human health and phytotoxicity (plant health), while other values are purely for human health. The underlying premise in existing New Zealand industry-based guidelines is that protection of on-site ecosystems is only required to the extent necessary to facilitate the use of the land.

This report only considers human health, and therefore the ecological component of the guidelines reviewed in the following section has not been considered.

A5.3 United States

A5.3.1 Legislative framework

The United States has a considerable body of legislation and guidance, both federal and state, that controls contaminated land assessment, management and remediation. The legislation is too voluminous to review in any detail here. As New Zealand has adapted parts of the US EPA's 'Superfund' guidance in developing its existing guidelines, the legislation controlling this programme will be reviewed.

The Federal Superfund programme was established under the Comprehensive Environmental Response Compensation and Liability Act (CERCLA) in 1980 and is administered by the US EPA. This law created a tax on the chemical and petroleum industries and provided broad federal authority to respond directly to releases or threatened releases of hazardous substances that may endanger public health or the environment. CERCLA and subsequent amendments established prohibitions and requirements concerning closed and abandoned hazardous waste sites. They provided for the liability of persons responsible for release of hazardous waste, and established a multi-billion dollar trust fund to provide for the clean-up when no responsible party could be identified. This is funded by taxes on the chemical and petroleum industries. The law authorises two kinds of response actions (US EPA, 2006b):

- short-term removals, where actions may be taken to address releases or threatened releases requiring prompt response
- long-term remedial response actions, that permanently and significantly reduce the dangers associated with releases or threats of releases of hazardous substances that are serious, but not immediately life threatening. These actions can be conducted only at sites listed on EPA's National Priorities List (NPL). There are about 1500 NPL sites.

CERCLA also enabled the revision of the National Contingency Plan (NCP). The NCP provided the guidelines and procedures needed to respond to releases and threatened releases of hazardous substances, pollutants or contaminants. The NCP also established the NPL.

The Hazard Ranking System (HRS) is the principal mechanism the US EPA uses to place uncontrolled waste sites on the NPL. It is a numerically based screening system that uses information from initial, limited investigations – the preliminary assessment and the site inspection – to assess the relative potential of sites to pose a threat to human health or the environment. Any person or organisation can petition the EPA to conduct a preliminary assessment.

The federal government plays a strong role in site-specific clean-up decisions and action. There are national guidelines for assessment of risk and decision making (see below), with the risk targets common to all clean ups.

Operating waste facilities fall within the Resource Conservation and Recovery Act (RCRA Corrective Action). The RCRA primarily provides that hazardous waste is properly managed so it does not contribute to future contamination, and under its corrective action programme, the clean up of existing contamination at operating industrial facilities is addressed. About 3800 sites were undergoing corrective action as of 2006, many more than on the Superfund NPL (US EPA, 2006b). Clean-up expectations are similar to that for Superfund. Thirty-eight states are authorised to run their own programmes. Federal oversight is limited.

A further federal programme is the clean up of underground storage tanks (UST), established under RCRA in 1984. US EPA's Federal UST regulations require that contaminated UST sites must be cleaned up to restore and protect groundwater resources and create a safe environment for those who live or work around these sites. The programme is intended to oversee clean-ups by responsible parties and to pay for clean ups at sites where the owner or operator is unknown, unwilling, or unable to respond, or which require emergency action. A trust fund, which receives US\$70 million per year from a fuel tax, has been established to enable this. There are 125,000 contaminated underground storage tank facilities to address under the programme, with thousands of new petroleum releases each year. The federal programme provides only general guidance, with states defining their own programmes. Risk-based decision making is encouraged but there is no specific protocol or expectations for clean up.

A5.3.2 Superfund risk assessment process

New Zealand has based much of its past soil guideline derivation on US guidance, and in particular the multi-volume Risk Assessment Guidance for Superfund (RAGS). This has been developed by the US EPA as a requirement of both CERCLA and RCRA¹⁸. Key guidance is RAGS Volume I (and its various parts in separate documents) covering human health risk assessment and Volume II covering ecological risk assessment. More recently, RAGS Volume III, Part A (US EPA, 2001b), on probabilistic risk assessment has been developed. Only Volume I is considered here.

RAGS Volume I has multiple parts, including supplementary guidance, developed over several years (US EPA, 1989a, 1991a, 1991b, 1991c, 1996a, 1996b, 2001a, 2001c, 2002a, 2004b), the most important of which for the New Zealand context are:

¹⁸ Comprehensive Environmental Response, Compensation, and Liability Act – otherwise known as CERCLA or Superfund; RCRA, Resource Conservation and Recovery Act.

- Risk Assessment Guidance for Superfund (RAGS): Volume I. Human Health Evaluation Manual (HHEM) (Part A, Baseline Risk Assessment) (US EPA, 1989a)
- RAGS Volume I, Part E: Supplemental Guidance for Dermal Risk Assessment (US EPA, 2004c)
- Supplemental Guidance for Developing Soil Screening Levels for Superfund Sites (US EPA, 2002a).

The risk assessment guidance is supported by the multi-volume *Exposure Factors Handbook*, first published in 1989 (US EPA, 1989b) as a three-volume set covering general factors, food intake factors and activity factors. The latest version was published in 1997 (US EPA, 1997) as a single 1200-page volume and available online at http://www.epa.gov/ncea/efh/. The handbook is intended to serve as a support document to *Guidelines for Exposure Assessment* (US EPA, 1992), the original version of which was published in 1986, and developed to promote consistency among the various exposure assessment activities carried out by the US EPA.

Chapter 6 of RAGS Volume 1, Part A (US EPA, 1989a) provides the basic scientific basis of exposure assessment and presents the basic equations to be used. Chapter 7 of this document covers the toxicological basis of risk assessment. RAGS Volume 1, Part B (US EPA, 1991a) originally provided additional information and equations on determining preliminary remediation goals (PRGs), but has since been superseded by the supplementary guidance of US EPA (2002a) for non-residential exposure and for the dermal exposure pathway by US EPA (2004c).

A PRG is an initially developed chemical concentration for an environmental medium that is expected to be protective of human health and ecosystems. Risk-based PRGs, either at scoping or later on, are initial guidelines. They do not establish that clean up to meet these goals is warranted, that is, they are not clean-up values (US EPA, 1989a, 1991a). The risk assessment process described in RAGS Volume 1, part A and B is, in effect, a site-specific risk assessment process and involves consideration of many exposure pathways.

The process described in the various RAGS documents is quite complex, time-consuming and expensive, and is only warranted for the typically large and complex sites that are named as Superfund sites. Soil screening levels (SSLs) have been developed to standardise and accelerate the evaluation and clean up of sites, initially in 1996 (US EPA, 1996a; US EPA, 1996b) with an updated procedure for non-residential exposure in 2002 (US EPA, 2002a).

It is the generic set of SSLs that are of interest for New Zealand, certainly with respect to the derivation methods. However, the generic values are potentially more conservative than would be derived in New Zealand. In the US context, SSLs are not national clean-up standards; instead, they are used as a first screening tool to identify areas, chemicals and pathways of concern at federally listed sites that need further investigation. They are but one step in a long and public statutory process of investigation, initial screening, site-specific risk assessment, establishment of clean-up criteria and selection of remediation.

In the US context, soil screening levels can be used as preliminary remediation goals (but not necessarily the final clean-up levels) provided conditions found during subsequent investigations at a specific site are the same as the conditions assumed in developing the SSLs. However, the conservative assumptions built into the generic SSLs, while appropriate for a screening analysis, may be overly conservative for setting PRGs and, ultimately, site clean-up levels (US EPA, 2002a). It is important to understand this context, as New Zealand practitioners frequently refer to SSLs when New Zealand guidelines are not available. The obvious conclusion from the US framework is that SSLs are rather more conservative than soil guideline values envisaged in the proposed framework for New Zealand (MfE, 2007).

Generic human-health SSLs based for residential land use were first developed in the Soil Screening Guidance (SSG) (US EPA, 1996a) with the technical background given in US EPA (1996b). The more recent supplemental guidance (US EPA, 2002a) widened the application of generic SSLs to commercial / industrial land use, for both indoor and outdoor workers, and to the construction scenario, with the latter having site-specific exposure durations but otherwise standard factors. A summary of pathways of concern for derivation of SSLs for the residential, commercial / industrial and construction settings is shown in table A5.2 (US EPA, 2002a). This list is not exhaustive but will depend on the particular site.

The 2002 supplement also provided new SSL equations for combined exposures via ingestion and dermal absorption, updated dispersion modelling data for the residential air exposure model, and new methods to develop SSLs for the migration of volatiles from subsurface sources into indoor air. These changes are important for New Zealand, because New Zealand guidance developed in the late 1990s was based on the earlier US EPA guidance. If the US EPA guidance is still to be followed, at least these updates should be considered.

Table A5.2: Summary of US EPA exposure pathways of concern for residential and	
commercial / industrial land uses, and construction for deriving SSLs	

Scenario1	Residential	Non-residential (commercial / industrial)		Construction	
Receptor	On-site resident	Outdoor worker	Indoor worker	Construction worker	
Pathways of concern	Ingestion (surface and shallow subsurface soils)	Ingestion (surface and shallow subsurface soils)	Inhalation (indoor vapours)	Ingestion (surface and subsurface soil)	
	Dermal absorption (surface and shallow subsurface soils)	Dermal absorption (surface and shallow subsurface soils) Ingestion (indoor dust) Migration to groundwater		Dermal absorption (surface and subsurface soil)	
	Inhalation (fugitive dust, outdoor vapours)	Inhalation (fugitive dust, outdoor vapours)	5	Inhalation (fugitive dust, outdoor	
	Inhalation (indoor vapours) Migration to groundwater	Migration to groundwater		vapours)	

The US EPA guidance is too extensive to attempt to summarise all the equations here. The basic equations used to derive generic SSLs are as used in the 'Timber Treatment Guidelines' (MfE and MoH, 1997) with a number of subsidiary equations to derive input values into these equations. However, the following are of note:

- Residential exposure:
 - The SSLs apply to the top 2 cm of soil, being the soil that people are mostly exposed to and generates the dust that may be inhaled or migrate into homes.
 - The default exposed group are children aged 1–6 and older children and adults aged 7–30.
 - For threshold substance the critical receptor is a child.

- For non-threshold substances age-adjusted exposure rates are calculated to account for exposure over the complete exposure duration, as an adult and child, with averaging over a default lifetime of 70 years.
- Soil ingestion and dermal exposure are combined to arrive at the SSL.
- Home-grown produce consumption is not considered, being left to site-specific consideration.
- Migration of volatiles to indoor air is not considered, because of difficulties identifying suitable default values for such things as building dimensions and the distance between the building foundation and the contamination; however, volatiles to outdoor air and inhalation of fugitive dust outdoors are allowed for. These pathways are not combined with ingestion and dermal exposure.
- Two commercial / industrial scenarios are considered, for indoor and outdoor workers, to recognise the different exposure these two groups would have to soil.
 - Indoor workers have no direct exposure to soil, but may contact indoor dust, and have exposure to volatiles. This scenario covers people such as full-time factory and shop workers.
 - Outdoor workers are those involved full-time in everyday outdoor maintenance activities involving moderate digging and landscaping (eg, the site caretaker). Such a worker is expected to have an elevated soil ingestion rate, dermal exposure and inhalation of dust or vapours.
 - Migration of volatiles to indoor air is not considered but volatiles to outdoor air and inhalation of fugitive dust outdoors has been allowed for (based on the Johnson and Ettinger (1991) model and associated (since updated) guidance (US EPA, 2002c, 2004c).
- Generic construction scenario SSLs have not been calculated because of the difficulty of defining standardised default exposure assumptions.

The RAGS process and calculation of SSLs by US EPA is based on an estimate of the reasonable maximum exposure (RME) expected to occur under both current and future land-use conditions. The RME is defined as the highest exposure that is reasonably expected to occur at a site. The intent of the RME is to estimate a conservative exposure case, that is, well above the average case, but still within the range of possible exposures. RMEs are estimated for individual pathways. If a population is exposed via more than one pathway, the combination of exposures across pathways also must represent an RME.

All soil screening level equations in the 1996 soil screening guidance were designed to be consistent with the concept of RME in the residential setting. Accordingly, the US EPA used reasonably conservative defaults for intake and exposure duration, combined with values for site-specific parameters that reflect average or typical site conditions, to develop the risk-based SSLs. The SSLs were based on RME assumptions rather than central tendency conditions because this approach results in a conservative (though not a worst-case) estimate of long-term exposure that is protective of the majority of the population (US EPA, 2002a).

In deriving the generic SSLs the bioavailability of substances is assumed to be 100 per cent (except for lead, for which a different derivation process based on a soil-blood lead model is employed) and background exposure is not subtracted from the acceptable daily intake for the contaminant concerned.

The US EPA reviewed and confirmed its use of RMEs in response to criticism that the concept has so overemphasised conservatism that most risk estimates are false or meaningless (US EPA, 2004a). The US EPA's policy position is that it uses defaults that guard against underestimating risk while also being scientifically plausible given existing uncertainty. Further, the policy is to examine and report on the upper end of a range of risks or exposures when there is uncertainty about where the particular risk lies – in other words, a precautionary approach.

The US EPA rejects the argument that combining several values results in excessive overestimates of risk (eg, combining two 95th percentile defaults results in an estimate above the 99th percentile and combining three 95th percentile defaults results in an estimate above the 99.9th percentile) as being too simplistic. It has pointed out that just multiplying the numbers as implied will not necessarily lead to the answers above, but will depend on the variability of the data and the shape of the input distributions, with different parameters having little 'influence' (eg, for narrow distributions there is little difference between high-end and central estimates). If all the input variables show the same variability, shape, etc, then the multiplicative reasoning with respect to compounding values is true, but otherwise not (US EPA, 2004a). The US EPA further notes that it is rarely the case that distributions are the same in actual situations.

An example is when three variables are involved and one has a wide distribution while the other two have narrow distributions: taking the mean of the wide distribution, instead of the 95th percentile, while taking the 95th percentile of the other two variables, will result in an outcome that is less than the intended 95th percentile (or some similarly high number) protection because of the overriding influence of the wide distribution. Following this logic, it is necessary to consider the likely variability of each exposure variable used in the risk assessment. Thus it may be appropriate to use central estimates (means) for most variables, but the 95th percentile for a variable that has a wide distribution (eg, exposure duration).

The SSL derivation has been used by a number of US EPA regional offices (eg, regions 3, 6 and 9) to produce their own sets of SSLs (called PRGs in Region 9), but these have been recently harmonised into a single set of regional guidelines (US EPA, 2008). In addition, an online PRG calculator (the Risk Assessment Information System – RAIS) developed by the United States Department of Energy's Oak Ridge National Laboratory using the RAGS guidance and the SSL equations, is available at http://rais.ornl.gov/prg/prg_document.shtml. The scenarios considered are:

- construction worker / excavation land use exposure to contaminants in soil
- industrial land use (indoor and outdoor worker) exposure to contaminants in soil
- industrial land use (indoor and outdoor worker) exposure to contaminants in groundwater
- recreational land use exposure to contaminants in soil
- recreational land use exposure to contaminants in water
- residential land use exposure to contaminants in soil
- residential land use exposure to contaminants in water
- agricultural land use exposure to contaminants in soil, groundwater and homegrown produce.

The online tool goes further in developing generic values than assumed by US EPA (2002a), in particular agricultural and recreational scenarios are considered, and multiple direct (ingestion, dermal) and indirect (inhalation of dust and vapours) pathways are combined. The documentation with the online tool notes that the combination of pathways will result in lower (more conservative) PRGs than the SSLs calculated in the Soil Screening Guidance.

The agricultural and recreational scenarios use site-specific exposure factors developed for the assessment of the Oak Ridge Reservation, in Tennessee (USDOE, 1999). The documentation warns that the default agricultural and recreational assumptions may not be applicable to other sites. The usefulness of these PRGs as generic values must therefore be questioned.

Most, if not all, states in the US have developed generic guideline values based on the US EPA guidance.

A5.4 Australia

A5.4.1 Legislative framework

Federal management of contaminated sites in Australia is provided by the National Environmental Protection (Assessment of Site Contamination) Measure (NEPM) (NEPC, 1999b). Its purpose is to establish a nationally consistent approach to assessment of site contamination. The measure was developed by the National Environment Protection Council (NEPC), a body comprising environment ministers from the Australian Government and each state and territory. The Council was brought into being by the National Environment Protection Council Act 1994, enacted by the Commonwealth Government following the Intergovernmental Agreement on the Environment in 1992. The Act provides for the NEPC making national environment protection measures in a number of defined areas, including general guidelines for the assessment of site contamination.

Various activities are subject to Commonwealth law, including activities of the Commonwealth in a participating jurisdiction. The Australian Government has implemented law to make the Commonwealth subject to relevant NEPM implementation law in the states and territories.

Responsibility for contaminated land rests with states and territories unless the site is owned by the Commonwealth, and individual states and territories have implemented specific legislation and produced guidance for contaminated site management. Detailed reporting of the various state provisions is beyond the scope of this study, but some brief examples are given here.

In New South Wales (NSW) the Environmental Planning and Assessment Act 1979 provides for the management of contaminated land to ensure that it is not put to inappropriate use; and requires local authorities to consider land contamination when making rezoning or development decisions. Councils are required to provide information on land contamination when issuing planning certificates. Land remediation is facilitated and controlled through State Environmental Planning Policy 55 – Remediation of Land (DUAP and EPA, 1998).

The NSW Environmental Protection Authority has issued a number of guidelines under the Contaminated Land Management Act 1997 and has adopted the framework of the contaminated land NEPM within these guidelines.

Victoria gazetted the State Environmental Protection Policy (Prevention and management of contaminated land) in 2002 under the Environment Protection Act 1970 (Victoria Government, 2002). The policy sets out, amongst other things, land-use categories (the same as in the NEPM) and the adoption of the investigation levels and the procedures for deriving such levels, set out in the NEPM.

Victoria initiated the use of environmental auditors within a statutory process to oversee assessment and remediation of contaminated land. New South Wales and Western Australia have adopted similar schemes.

The NEPM has recently been reviewed, with many priority and lesser priority proposals for its modification coming out of that review (NEPC, 2006). The recommendations included:

- Revise the NEPM policy framework and Schedule A (the assessment process) to improve clarity and understanding of the fundamental site assessment principles and emphasise the appropriate use of the NEPM, in particular to address the misuse of investigations levels, eg, use of investigation levels for clean up.
- Revise the existing Health-based Investigation Levels (HILs) in the light of current knowledge, leading to more accurate and often less conservative numbers, and the derivation of additional HILs for priority substances.
- Follow-up review of worldwide models and field methods for the assessment of volatiles, and adopting as interim guidance a model(s), analytical approaches and field methods, from a 'best-fit' scenario most suited to Australian conditions.
- Review current bioavailability approaches, methods and limitations to improve the basis for their application in site assessment.

The widespread practice of invoking screening or investigation levels as a clean-up standard, contrary to the intent, has led in most jurisdictions to widespread reliance upon the 'dig and dump' strategy as the most common remediation method (Fowler, 2007). This has been reinforced by environmental authorities in a number of jurisdictions transferring responsibility to environmental auditors for determining remediation approaches. Auditors have tended to act cautiously by recommending removal to landfill in most cases rather than exploring alternative approaches, largely out of a concern to avoid any possible future personal liability.

A5.4.2 The NEPM derivation methodology

The methodology for deriving HILs is mainly described in NEPM schedules B(7a) and B(7b) (NEPC, 1999a, 1999c), the former providing the derivation methodology and the latter a description of standard exposure scenarios (see below). Three exposure pathways are considered: soil ingestion, inhalation of particulates, and dermal absorption.

Exposure via home produce consumption is discussed but not provided for in the scenarios for which HILs are calculated, with site-specific assessment recommended where home produce consumption is found to be greater than 10 per cent of fruit and vegetable consumption. Exposure via consumption of home-grown produce was considered to be too variable and uncertain (due to site-specific differences and plant type) to be included on a generic basis (NEPC, 1999b). Implicitly, home produce consumption of less than 10 per cent of total produce consumption is considered an insignificant contribution to a resident's contaminant intake. None of the HILs presented in the NEPM include exposure via consumption of home-grown produce in their derivation.

However, the revised NEPM draft (NEPC, 2010) provides a default assumption for the low density residential scenario of 10% home produce ingestion. No home produce ingestion is assumed for high-density residential or any other scenario.

The derivation provided in NEPC (1999c) uses similar equations to those of other jurisdictions, being based on estimating the total exposure to a given substance:

 $Exposure = BE + (S_{ing} \times C_{ing} \times B_{ing}) + (S_{inh} \times C_{inh} \times B_{inh}) + (S_{skin} \times C_{skin} \times B_{skin})$

where:	BE	=	background exposure
	Sing	=	amount of soil ingested
	Cing	=	concentration of substance in ingested soil
	Bing	=	bioavailability of substance when ingested
	S _{inh}	=	amount of soil/dust inhaled and retained
	C_{inh}	=	concentration of substance in soil/dust inhaled and retained
	\mathbf{B}_{inh}	=	bioavailability of soil/dust inhaled and retained
	\mathbf{S}_{skin}	=	amount of soil on skin
	C_{skin}	=	concentration of substance in soil on skin
	\mathbf{B}_{skin}	=	bioavailability of substance when on skin

The bioavailability of substances is assumed to be 100 per cent if specific information is not available (NEPC, 1999a) (a conservative default assumption for dermal absorption for most substances); but it is also noted that different levels of bioavailability will occur between soil ingested, inhaled or in contact with skin (NEPC, 1999d). The revised NEPM draft (NEPC, 2010) re-evaluated this approach and now assumes the default oral bioavailability for arsenic to be 70%, not 100%. The oral bioavailability for lead is now assumed to be 50%, not 100%.

Total exposure includes that from background sources, principally food and water, and therefore less than the allowable intake of a contaminant is assigned to contaminated soil sources. The total exposure must not exceed the provisional tolerable weekly intake (PTWI), acceptable daily intake (ADI) or Guideline Dose (GD – for cancer toxic effects) of the contaminant concerned.

A specific equation is not provided in the Schedule B(7a) for the derivation of an HIL. But implicitly, the same contaminant concentration is taken to be soil that is ingested, inhaled and attached to the skin, with this concentration solved for when the total exposure is equated to the allowable intake (PTWI, ADI or GD), or some fraction of this value. However, no subsidiary equations are provided for calculation of the inhaled dust intake or dermal absorption. Allowable intakes are from WHO/FAO sources; or in the case of guideline doses for carcinogenic compounds, as set by national health advisory bodies.

Health investigation levels have been produced for four exposure scenarios, while two other exposure scenarios are discussed but left for site-specific assessment. The four exposure scenarios for which values are given are (as listed in NEPC, 1999a):

- A. standard residential (<10 per cent consumption of produce grown on-site; no poultry) includes children's daycare centres, kindergartens, preschools and primary schools
- D. residential with minimal opportunities for soil access includes dwellings with fully and permanently paved yard spaces such as high-rise apartments and flats
- E. parks, recreational open spaces and playing fields includes secondary schools
- F. commercial / industrial includes premises such as shops and offices as well as factories and industrial sites.

The other two scenarios, for which values are not given because investigation levels need to be determined on a site-specific basis, are:

- B. residential with substantial vegetable garden (contributing 10 per cent or more of vegetable and fruit intake) and/or poultry providing any egg or poultry intake
- C. residential with substantial vegetable garden (contributing 10 per cent or more of vegetable and fruit intake); poultry excluded.

While a methodology to derive HILs is nominally provided in the 1999 NEPM version, the majority of the HILs have not followed this methodology, and have been inconsistently derived. Health investigation levels for the residential scenario are nominally determined using a two-and-half-year-old child as the critical receptor. Values for each scenario are not calculated in detail; rather, those for scenarios D, E and F are simple factorings up from the residential scenario.

As noted in the previous section, the NEPM methodology has been undergoing review. The emphasis was on:

- revising the existing HILs in the light of current knowledge
- introducing a methodology to derive ecological investigation levels (EILs)
- reviewing worldwide models and field methods for the assessment of volatiles with a view to adopting those that best fit Australian conditions
- reviewing current bioavailability approaches.

All scenarios in the revised draft version of the NEPM (NEPC, 2010) are now individually modelled and seem internally consistent. The revised draft version provides a compilation of parameters that should be used in deriving the HILs (Schedule B7 Table 5).

For the final revised NEPM readers may want to consult the following website: http://ephc.gov.au/contam

A5.5 Canada

A5.5.1 Legislative framework

Within Canada, individual provinces are responsible for their own policies and regulation of contaminated sites, independent of the other provinces and the federal government. Each province has specific regulation on land contamination, but some regulatory regimes are more developed than others. Quebec and British Columbia are more active than the other provinces. Each province has a set of land-use-based generic criteria but the values differ. Each province also allows risk-based management decisions, but requirements are not the same (Beaulieu, 2007).

Many provinces have public lists (internet-based) of known contaminated sites, but the lists are not compiled in the same way. Three provinces have put in place a network of private acknowledged experts to supervise some of the assessment and remediation work. There is a trend to tackle governmental environmental liabilities, with the federal government more active than the provinces. The larger provinces have developed their own policies and guidance which may or may not draw on national guidance developed by the Canadian Council of Ministers of the Environment. CCME as an intergovernmental body has developed a number of documents (see next section) intended to be applied Canada-wide, but the CCME guidelines are sometimes adopted, sometimes transformed and sometimes ignored by the provinces (Beaulieu, 2007). The provinces are free to develop their own regulation, policies, criteria and priorities independently of the other provinces and the federal government.

There is no overarching federal legal framework for contaminated land in Canada (Fowler, 2007), but a variety of federal legislation exists governing soil contamination, with the Canadian Environmental Protection Act (CEPA), Fisheries Act, and Canadian Environmental Assessment Act (CEAA) all containing provisions that relate to contaminated sites management. In 2003, the federal government established the Federal Contaminated Sites Accelerated Action Plan (FCSAAP), a new contaminated sites initiative to help identify, assess and manage the risks at contaminated properties under the custodial care of Canadian federal government departments.

A5.5.2 Guideline derivation

The Canadian Council of Ministers of the Environment has developed separate soil quality guidelines for the protection of environmental and human health, as generic guidance for the federal and provincial governments (although as noted earlier, not all provinces have adopted them). Guidelines are developed based on four defined land-use scenarios: agricultural, residential / parkland, commercial and industrial. Other land-use scenarios may be defined by provincial jurisdictions or on a site-specific basis.

The guidelines have been produced in response to growing public concern over the potential ecological and human-health effects associated with exposure to contaminated sites in Canada. To promote consistency and provide guidance in assessing and remediating contaminated sites under the National Contaminated Sites Remediation Program, initiated by CCME in 1989, an interim set of numerical environmental quality guidelines was released in 1991 (CCME, 1991). These adopted existing criteria for soil and water used by various jurisdictions in Canada. However, many of the interim criteria for soil were based on professional judgement. To ensure that revised guidelines are scientifically defensible, a derivation protocol was developed in 1996, with an update released in 2006 (CCME, 1996, 2006).

The Canadian land-use definitions are (CCME, 2006):

- **Agricultural**: where the primary land use is growing crops or tending livestock. This also includes agricultural lands that provide habitat for resident and transitory wildlife and native flora.
- **Residential / parkland**: where the primary activity is residential or recreational activity; parkland is defined as a buffer between areas of residency, and also includes campground areas, but excludes wildlands such as national or provincial parks.
- **Commercial**: where the primary activity is commercial and not residential or manufacturing. This does not include zones where food is grown. The toddler was chosen as the critical receptor for children, as commercial facilities (such as a shopping mall) could have childcare facilities and children would have unrestricted access to the complex.
- **Industrial**: where the primary activity involves the production, manufacture, or construction of goods.

Soil quality guidelines for each chemical are developed for both ecological and human receptors. For each of the four land uses, to protect both human health and the environment, the most protective guideline is chosen as the recommended soil quality guideline. This is done after following through a complex set of checking a number of direct and indirect pathways for both ecological and human receptors. This is shown diagrammatically in figure A5.2.

Only the human health guideline derivation process will be considered here. The human health soil quality guideline (SQGHH) is determined by evaluating direct soil exposure (soil ingestion, dermal contact, and particulate inhalation), transport of contaminants through groundwater to potential potable water sources, intrusion of contaminant vapours into buildings, and human consumption of contaminated food. The specific exposure scenario is dependent on the land use, with some of the exposure pathways not evaluated for all land uses or contaminant types. The lowest of the soil concentrations deemed protective of each of these potential exposure pathways becomes the SQGHH. This contrasts with the approach taken in many other jurisdictions to combine exposure from the various pathways on the assumption the risk is additive.

The development of the soil quality guideline is a two step process (CCME, 2006). The first step considers all direct soil exposure pathways, including the ingestion of soil/dust, dermal contact, and inhalation of soil particles into lungs (as a combined value), as well as the primary indirect pathways. The primary indirect pathways that are included in the first step are dependent on the contaminant type, but may include inhalation of vapours migrating into indoor air (for volatile contaminants), ingestion of groundwater used as potable water (for soluble organic contaminants) and consumption of produce (for substances that biomagnify). The latter pathway applies principally to agricultural land use. The second step is to assess two 'check mechanisms': exposure from ingestion of food grown on contaminated soils (if not already applied in the initial step for substances which biomagnify); and the off-site migration via wind and water erosion of contaminants from commercial or industrial sites to more sensitive neighbouring properties.

The basic equations used for the derivation are similar to those used in other derivations and will not be repeated here. The equations in CCME (2006) are as presented in Health Canada (2004) and use the various exposure parameters presented in that document.



Figure A5.2: Overview of steps for derivation of a soil quality guideline in Canada

Source: CCME, 2006.

In common with many jurisdictions, for threshold contaminants the background intake (estimated daily intake, EDI) is subtracted from the tolerable daily intake (TDI) to obtain a residual TDI before calculating the soil guideline value. An unusual aspect is that the residual TDI is then, as a matter of policy, equally allocated five ways between soil, consumer products, air, water and food. This has the effect of double-counting exposure to food and water. The allocation of the residual is shown diagrammatically in figure A5.3.

Having only a small part of the residual TDI allocated to soil potentially results in quite conservative soil guideline values for some contaminants. As a result, the 2006 revision of the 1996 protocol now permits adjustment of the soil allocation factor for chemicals that can defensibly shown to be not present (or not in significant concentrations) in all media. When the EDI is greater than the TDI (residual TDI = 0), theoretically the population cannot be safely subjected to any increased exposure. In these circumstances, the provisional soil quality guideline should be set at the background soil concentration or practical quantitation limit for that contaminant (CCME, 2006).

For threshold contaminants an excess cancer risk approach is taken, with guideline values calculated for both 1×10^{-5} and a 1×10^{-6} incremental risk, with the choice being left up to the individual jurisdictions as to what risk value to use.

The protocol recognises two generic soil types to minimise the uncertainty in guideline derivation introduced by soil variability: coarse-textured soils (soils containing predominantly sand and gravel sizes) and fine-textured soils (soils containing predominantly silt and clay sizes).

The protocol does not specify the depth to which the generic soil guidelines apply, although it notes that most direct human and ecological exposure pathways apply to soil located at or near the surface and suggests: *Surface soils are often defined as those within the uppermost 1.5 m of the soil profile*.



Figure A5.3: Assumed soil allocation factor from the residual tolerable daily intake

Source: CCME, 2006.

A5.6 The United Kingdom

Note: Since this section was written the situation in the United Kingdom has changed. The legislative framework remains the same but the derivation of SGVs has changed from the probabilistic model presented in Defra and EA (2002a) to a deterministic methodology as set out in EA (2008a), supported by a revised methodology for determining toxicological criteria in EA (2008b), both published in August 2008. The existing SGs calculated in 2002 have been withdrawn. Several new SGV documents have been published (eg, EA, 2009a, 2009b, 2009c) with more in preparation.

The revisions follow a review of the methodology (Defra, 2006a) and the outcomes of that review (Defra, 2008a). The change to a deterministic model was foreshadowed in Defra (2006a). Other revisions include reconsideration of the generic land-use scenarios and default assumptions used in the CLEA model to derive SGVs including improvements in clarity, internal consistency, and practical usability of the approach. The basic exposure equations remain the same, albeit used in a deterministic fashion with single-point estimates for the various parameters. New guidance on the definition of contaminated land has also been published (Defra, 2008b).

As key decisions had already been made on the new New Zealand methodology by the Technical Advisory Group by the time the authors became aware of the changes to the UK approach and as the changes are not so significant as to require reconsideration of the New Zealand approach (and arguably reinforce some of the choices made), the decision was made not to update this section.

A5.6.1 Legislative framework

In the United Kingdom, contaminated land is regulated under the Environmental Protection Act introduced in 1990. The act initially mainly focused on preventing new contaminated land being created. It wasn't until the Environment Act was introduced in 1995 that contaminated land was specifically addressed. Section 57 of the 1995 legislation inserted Part 2A into the Environmental Protection Act 1990, although it did not come into force in England until April 2000. Part 2A has also been implemented in Wales and Scotland, with minor differences. Part 2A largely replaced and modernised regulatory powers that existed under much older public health legislation under which polluted land could be abated as a statutory nuisance.

The legislation was intended to improve the focus and transparency of the controls, to enable all problems resulting from contamination to be handled as part of the same process, to increase the consistency of approach taken by different authorities; and to provide a more tailored regulatory mechanism – including liability rules that are better able to reflect the complexity and range of circumstances found on individual sites.

Part 2A states that:

'Contaminated land' is any land which appears to the local authority in whose area it is situated to be in such a condition, by reason of substances in, on or under the land, that -

significant harm is being caused or there is a significant possibility of such harm being caused; or pollution of controlled waters is being, or is likely to be, caused.

The Environmental Protection Act is administered by the Department for Environment, Food and Rural Affairs (Defra) which has issued guidance on the Part 2A in the form of Circular 01/2006 'Contaminated land' (Defra, 2006b), which replaced an earlier 2000 circular. Regulations under the Act specify detailed provisions, including special sites, remediation notices, compensation, appeals and public registers. Section 78R requires enforcement authorities to keep a public register of all regulatory action taken by the enforcing authority in respect of the remediation of contaminated land, and will include information about the condition of land (Defra, 2006b). However, when Part 2 came into force, Section 143 was repealed (this would have required local authorities to compile registers of land which may be contaminated).

The policy objectives of Part 2A with respect to contaminated land are threefold:

- a. to identify and remove unacceptable risks to human health and the environment
- b. to seek to bring damaged land back into beneficial use
- c. to seek to ensure that the cost burdens faced by individuals, companies and society as a whole are proportionate, manageable and economically sustainable.

These three objectives underlie the 'suitable for use' approach to the remediation of contaminated land, which focuses on the risks caused by the contamination. The approach recognises that the risks presented by any given level of contamination will vary greatly according to the use of the land and a wide range of other factors, such as the underlying geology. Risks therefore need to be assessed on a site-by-site basis. The 'suitable for use' approach consists of three elements:

- a. Ensuring that land is suitable for its current use by identifying any land where contamination is causing unacceptable risks to human health and the environment, assessed on the basis of the current use and circumstances, and returning such land to a condition where such risks no longer arise, ie, remediating the land.
- b. Ensuring that land is made suitable for any new use, as planning permission is given for that new use by assessing the potential risks from contamination, on the basis of the proposed future use, before permission is given for the development and, where necessary, remediating the land before the new use commences.
- c. Limiting requirements for remediation to the work necessary to prevent unacceptable risks to human health or the environment in relation to the current use or future use of the land for which planning permission is being sought. In other words, recognising that the risks can be satisfactorily assessed only in the context of specific uses of the land (whether current or proposed), and that any attempt to guess what might be needed at some time in the future for other uses is likely to result either in premature or unnecessary work.

Local authorities have the primary regulatory role under both planning legislation and Part 2A. Land contamination is a material planning consideration within the planning regime. This means that a planning authority has to consider the potential implications of contamination both when it is developing 'structure' or 'local' plans and when it is considering individual applications for planning permission. Under Part 2A, a local authority must ensure that their area of responsibility is inspected to identify contaminated land, to determine whether any particular site is contaminated land, and, provided the site is not a 'special site', establish liability for the costs of inspection and remediation, to decide, after consultations, what remediation is required and ensure remediation takes place. The responsibility for establishing liability and enforcing remediation of designated as special sites lies with the Environment Agency (EA). There are four kinds of special sites; where 'controlled' water is being polluted, particular industrial sites which pose special problems or are subject to national legislation of some kind, defence sites, and land contaminated by radioactive material (Defra, 2006b).

In addition to acting as the enforcement authority for special sites, the Environment Agency is required to assist local authorities in identifying contaminated land, particularly in cases where water pollution is involved, report periodically on contaminated land and provide site-specific guidance to local authorities. In response to the need to assist and provide guidance, the Environment Agency and Defra have developed risk-based procedures for assessing harm from contaminated sites to humans and ecosystems. Examples are the research and development publications CLR 7–11 on human health risk assessment (Defra and EA, 2002a, 2002b, 2002e, 2002f, 2004). These set out, among other things, the requirements for risk assessment, a set of priority contaminants for development of SGVs, the framework for toxicity assessment and the model (CLEA) used for deriving generic SGV values and for carrying out site-specific assessments.

To date only 10 generic SGVs have been published and development of further SGVs has now been suspended pending the completion of a review of the underlying assumptions for the CLEA model (Defra, 2006a). A number of issues have arisen, including criticism that some values are too conservative and uncertainty as to how the generic SGVs fit under the Part 2A regime. In 2005, Defra issued a statement advising caution on applying the SGVs to determine whether land was contaminated under Part 2A (Defra, 2005). The statement included the following:

... it should be a matter for careful consideration by local authorities whether concentrations of substances in soil equal to, or not significantly greater than, an SGV would meet the legal test ... it is apparent that there is a wide body of opinion that such concentrations would not necessarily satisfy that legal test.

The problem identified was that soil concentrations below SGVs provide are 'acceptable' but do not necessarily indicate that concentrations at or just above the SGV will be 'unacceptable' in the legal context. Exceedance of SGVs only indicates that further assessment or remedial action may be required (Defra, 2005). This appears to call into question whether the CLEA model can achieve one of its key objectives, ie, to determine whether land is 'contaminated' under Part 2A.

A5.6.2 Derivation of soil guideline values

The United Kingdom uses the CLEA model (Defra and EA, 2002a) to derive soil guideline values. The model is used to estimate average daily human exposure (ADE) to soil contamination based on the conceptual exposure models for three standard land uses. These are:

- **residential** covers a wide variety of dwellings including detached, semi-detached and terraced properties up to two storeys high, and takes into account several different house designs including buildings based on suspended floors and ground-bearing slabs. Residents are assumed to have private gardens and/or access to community open space close to the home and exposure has been estimated with and without a contribution from eating home-grown vegetables
- **allotment** allows for the use of communal open space, commonly provided by the local authority, for local people to grow fruit and vegetables for their own consumption
- **commercial** / **industrial** assumes that work takes place in a permanent single-storey building, factory or warehouse where employees spend most time indoors involved in office-based or relatively light physical work. This land use is not designed to consider those sites involving 100 per cent hard cover (such as car parks), because of the implausibility of exposure from ingestion or skin contact that the scenario assumes.

The ADE is defined as the average daily amount of a contaminant to which a critical human receptor is exposed over the duration of exposure, calculated using the equation below and is reported as a function of body weight to enable direct comparison with relevant health criteria values (HCV). As noted in Defra and EA (2002a), the equation is based on a standard methodology that has been adopted internationally for such assessments. For example, for an exposure scenario with three exposure pathways – oral and inhalation intake and dermal contact – the equation is:

$$ADE = \frac{IR_{inh} \times EF_{inh} \times ET_{inh}}{AT \times BW} + \frac{IR_{oral} \times EF_{oral} \times ET_{oral}}{AT \times BW} + \frac{IR_{dermal} \times EF_{dermal} \times ET_{dermal}}{AT \times BW}$$

where: the subscripts inh, oral and dermal refer to the inhalation, ingestion and dermal contact routes respectively, and

- ADE = average daily human exposure to a chemical from soil (mg/kg BW/day)
- IR = chemical exposure rate (mg/day)
- EF = exposure frequency (days/year)
- ED = exposure duration (years)
- BW = human body weight (kg)
- AT = averaging time (day)

The chemical exposure rate is a function of the concentration of contaminant in the relevant medium and the daily human exposure rate to that medium, which in turn is a function of land use and the receptor(s) exposed.

In the usual way, ADE is equated with the relevant HCV which is either an index dose value for a non-threshold substance (see MfE, 2010b for a further explanation of an index dose) or, for a threshold substance, a tolerable daily soil intake (TDSI), which is the TDI less the background intake from sources other than soil. An index dose is set specifically for exposure to soil and does not have the background intake deducted.

The conventional approach to risk assessment has been the use of deterministic models, in which single values are inserted into the simple equations defining exposure. However, the UK uses a probabilistic model, in which some of the single-value parameters in the exposure assessment are replaced with a family of values selected from defined probability distributions (Defra and EA, 2002a). This is intended to avoid the problem in a deterministic model of having to deal with parameter uncertainty and variability by selecting values representative of a worst-case exposure scenario, a practice which Defra and EA (2002a) notes as a common practice. While this has the assumed comfort of being more protective against an unforeseen situation or risks to sensitive individuals, the problem with this approach can be that such choices, however defensible they might be individually, tend to be implausible collectively (Defra and EA, 2002a). It has the further disadvantage that the model is complex and unable to be understood by any but the most expert individual.

Eight parameters are treated probabilistically in the CLEA model, on the basis that:

- the exposure estimate is sensitive to a change in its value
- variability in the parameter is well characterised
- there is no correlation between any two independently modelled probabilistic parameters.

The eight parameters treated probabilistically are:

- body weight
- total body surface area as a function of body weight
- respiration rate as a function of body weight
- mean daily soil ingestion rate by children aged 1–6 years
- estimated ratio of the concentration of a contaminant in chosen vegetables to the contaminant concentration found in the soil
- daily vegetable consumption rate
- fraction of homegrown garden vegetables as part of daily vegetable consumption rate
- fraction of exposed skin area in contact with soil.

The output of the CLEA model, for a particular contaminant and land-use scenario is a probability distribution of average daily exposure, calculated through many iterations of inserting values from the distributions of each of the exposure parameters into the exposure equation. A value then has to be chosen to equate with the HCV to derive the SGV. The point chosen by Defra and the Environment Agency is the 95th percentile of that distribution, so as to arrive at a 'reasonable worst case'. Note that this is not the same as being protective of the 95th percentile of the population, as some practitioners in the UK have assumed (Defra, 2006a).

The CLEA model can incorporate up to 10 different environmental pathways based on the concentration of the chemical in soil, with the choice dependent on the particular land-use category and specific considerations on the fate and transport, and toxicological properties of the contaminant of concern (eg, a vapour pathway would be chosen in the case of a volatile organic compound but not in the case of a heavy metal).

The 10 exposure pathways are:

- ingestion of soil
- ingestion of household dust
- ingestion of contaminated vegetables
- ingestion of soil attached to vegetables
- dermal contact with soil
- dermal contact with household dust
- inhalation of fugitive soil dust
- inhalation of fugitive household dust
- inhalation of vapours outside
- inhalation of vapours inside.

The model also uses 18 age intervals (or age classes) to break down the exposure characteristics of a human lifetime, allowing the model flexibility to consider exposure periods of a year or more. The first 16 intervals correspond to the first 16 years of life, the 17th interval is typical of an adult working life (age 16–59), and the 18th represents retirement (age 60–70). The intervals have been chosen to represent those stages in life where the most significant differences in site use are likely to occur. In deriving SGVs for the standard residential and allotment land-uses, a young female child from birth to six years is assumed to be the critical receptor. For the standard commercial and industrial land-use, a working adult is assumed to be the critical receptor. To date, 10 soil guideline values have been published.

Because of the probabilistic approach, it is difficult to compare the exposure parameter values with those adopted in deterministic models. However, of note is that exposure frequencies are not a constant across all exposure pathways, as is common with some deterministic models. For example, dermal exposure to indoor dust and outdoor soil are treated separately, with indoor dust exposure for 365 days a year, but outdoor soil exposure is treated as a less frequent occurrence. In addition, indoor dust has a lower concentration of chemical than outdoor soil, to reflect the fact that the indoor dust is only partly made up of outdoor soil tracked inside. This sort of refinement is not included in, for example, Australian and New Zealand guideline derivations.

A number of updates to the model have been issued as Contaminated Land Briefing Notes (EA, 2004a, 2004b, 2005a, 2005b) in respect of the dermal exposure route, vapour intrusion into buildings and combining exposure pathways.

For the dermal exposure route, the Environment Agency took into account work by the US EPA (2001a). Skin areas exposed to contaminants and soil adherence factors were revised (including differentiating between indoor and outdoor activities). In addition, dermal exposure was changed from being explicitly time-based (the default had been 12 hours per day) to an exposure per event, using a contaminant-specific absorbed fraction per event. These changes had no effect on the SGVs published to that point (arsenic and cadmium) because dermal intake was considered to be negligible (EA, 2005a).

The changes to the vapour intrusion algorithm within the CLEA model were triggered by a report commissioned by the Environment Agency in 2002. The original model considers vapour intrusion for both concrete slab-on-grade and concrete and wooden suspended floor construction. This report (Evans et al, 2002) examined 10 models for modelling vapour intrusion for the slab-on-grade situation and selected four for detailed assessment against case study results. The report concluded that the existing vapour intrusion models should be replaced by the Johnson and Ettinger (1991) model. This model assumes partitioning of vapours from the soil to the soil gas, migration of these vapours up through the soil to the underside of the slab, and then migration into the building through dust-filled cracks around the perimeter of the building. As a result of the recommendation, the CLEA model was changed and at the same time a review of typical British construction details resulted in modifications to the default parameters in the model (EA, 2004a, 2004b). However, the adoption of the Johnson and Ettinger approach meant that the previous suspended floor options were dispensed with in the CLEA model. The details of the adopted algorithms are beyond the scope of this report.

The CLEA model has been the subject of criticism within the contaminated-land assessment community in Britain, and the model and its underpinning policy is currently undergoing review. A discussion document (Defra, 2006a) was released in late 2006 which noted that concerns had been expressed about the limited number of SGVs; and that the SGVs that existed were not proportionate or realistic. There was an overall perception that the values were too stringent. The discussion document proposed making some immediate changes to the derivation method and proposed further study on possible changes. Immediate recommended changes included:

- replacing the child's soil ingestion parameter from a probability distribution with means of 100 mg/day and 95th percentile of 300 mg/day to a single-point estimate of 100 mg/day
- changing home-grown produce consumption from being based on a self-sufficient subgroup of the population to being based on a reasonable worst case for the whole population
- reviewing plant uptake models, with the objective of taking the conclusions of that review into the CLEA model
- improving guidance for estimating vapour intrusion into buildings the perception was that the models overestimated vapour intrusion and were not consistent with British building practices
- changing the CLEA model to being fully deterministic.

With respect to plant uptake and vapour intrusion, UK practitioners had expressed concern during the review that these pathways are very difficult to predict on a generic basis as a result of both scientific uncertainty and site variability. This was noted as consistent with the opinion of the US EPA, although most other countries still include estimation of these routes within their guidance because there are few practical alternatives (Defra, 2006a). Excluding these pathways was considered potentially problematic as they often represent significant mechanisms for human exposure for some chemicals. For plant uptake, CLEA considers both uptake into roots and above-ground-parts of vegetables, and divides produce intake into different types of above-ground and below-ground vegetables with uptake factors for each; a single consumption-weighted factor is derived for each group of above-ground and below-ground vegetables. Factors are derived for each contaminant on a case-by-case basis. The following hierarchy is used in selecting uptake factors (Defra, 2006a):

- empirical studies of plant uptake for the specific vegetables and chemicals of concern (that is, measured uptake factors under typical growing conditions)
- generic values for plant uptake recommended by authoritative bodies from the UK and other countries (often includes measured uptake factors from a range of plants and under varying soil and climate conditions)
- generic screening models based on good scientific principles.

For plant uptake of organics, the current CLEA model uses the relationship of Ryan et al (1988) which built on earlier work of Briggs et al (1982, 1983) to predict root and shoot concentrations of organic compounds (Defra and EA, 2002a; EA, 2006). The relationship assumes that plant uptake is proportional to the partitioning of an organic chemical between the soil and soil solution (pore water), based on experimental work on barley in growth solutions using a small range of organic compounds. The partitioning is a function of the organic matter present and the relative hydrophobicity of the compound, as measured by the octanol-water partition coefficient, K_{ow} . The greater the fraction of soil organic carbon (f_{oc}) and the higher the hydrophobicity, the lower is the plant uptake. The Ryan et al (1988) approach applies to non to weakly polar and relatively hydrophilic compounds rather than hydrophobic organic contaminants (log K_{ow} 0 to 4). Care should be taken for log K_{ow} outside the range for which the relationship was developed and it should not be used for compounds that ionise in the soil (Defra and EA, 2002d; EA, 2006)

Defra (2006a) notes that the Environment Agency recognised the limitations of the Briggs and Ryan relationships and was reviewing available models for uptake of organics. This review (EA, 2006) was released at about the same time as the Defra review. The review concluded that none of the plant uptake models was adequate as a general screening tool, with all resulting in over-estimates of root uptake, in some cases by up to five orders of magnitude. The situation with uptake into above-ground parts was more confused, with individual models both under-and over-predicting. The simplest models (eg, Travis and Arms, 1988) were just as effective as the more complex models. The report concluded with recommendations for further study.

A5.7 The Netherlands

A5.7.1 Legislative and policy framework

The main piece of legislation pertaining to soil contamination in the Netherlands is the Soil Protection Act 1987. This Act aims to prevent new pollution and requires that soil contaminated after 1987 is remediated as much as can be reasonably achieved so the land can be returned to 'multifunctional use', ie, for all land uses. For soil contaminated before 1987, the seriousness of contamination is determined and the necessary action identified. Two generic risk-based soil screening values are used to determine the seriousness of contamination: intervention values (IVs) and target values (TVs). Both screening values are based on potential risks to ecosystems, while intervention values are based on the potential risk to humans and ecosystems.

Intervention values are used to determine cases of serious soil contamination, while target values are viewed as sustainable-soil-quality objectives. Additionally, an intermediate value, which is arbitrarily set as the average of the target value and intervention value, is used to assist the process of site investigation (Swartjes, 1999; Carlon, 2007). These values are applied independent of soil use, whether residential, industrial or other use, and have statutory standing through the Soil Protection Act.

From a site investigation, which is conducted following standardised procedures, four outcomes are determined:

- 1. If the average soil concentration is below target values, no action is required.
- 2. If average soil concentrations exceed target values but fall below intermediate values, no further investigation is required but minor restrictions on land use are put in place.
- 3. If average soil concentrations exceed the intermediate values but are below intervention values, further investigation is required to confirm these concentrations, after which restrictions on land use are put in place.
- 4. For soils in which the average soil concentrations in 25 m^3 of soil exceed the intervention value, remediation will be necessary in principle, but the urgency of remediation has to be determined.

Remediation urgency is determined using a standardised computer-based methodology (CSOIL), to distinguish between urgent and non-urgent cases of serious soil contamination. Non-urgent cases are taken up in the provincial soil remediation programme without a defined time for starting the remediation. The determination of remediation urgency is based on actual (ie, site-specific) risks to human health, the ecosystem and risk due to contaminant migration.

Before 1997, all soils in the Netherlands were required to be remediated to multifunctional use (essentially to the target values) unless there were site-specific reasons not to do so. However, this 'strict' remediation goal result in a standstill in site remediation operations due to the perceived expense and necessity in relation to benefit and subsequent land use; it also prevented the development of urban land, revitalisation of business sites and sale of companies (VROM, 1999). Because of this, the Dutch Government reviewed the soil remediation policy to establish how the impediments to soil remediation could be removed.

In addition to the screening values, reference values have been derived for acceptable soil quality after remediation (Swartjes and Walthaus, in Carlon, 2007). Reference values represent sustainable soil quality of the upper soil layer, being the top layer of soil from 0.5 to 1 metre depth depending on land use. A distinction is made between land-use specific national reference values and local reference values, which can be derived on a site-specific basis. National Reference Values have been derived for several specific land uses, for immobile contaminants only. Mobile contaminants should be removed, as far as this is economically possible (Carlon, 2007).

A5.7.2 Calculation of guideline values

The target values for soil are determined for negligible risk to ecosystems, which is assumed to be 1 per cent of the maximal permissible risk level for ecosystems (MPR_{eco}). MPR_{eco} is defined as the hazardous concentration for 5 per cent of the species in the ecosystem (HC5), ie, 95 per cent protection. The HC5 is derived on an empirical basis by statistical interpretation of observed NOECs (no observed effect concentrations) and LOECs (lowest observed effect concentrations). For metals the added risk approach was followed for the derivation of target values by adding the natural background concentration in soils to the risk-based concentration as calculated above.

The human exposure model CSOIL is used to derive intervention values for soil and groundwater (Swartjes, 1999). The exposure is calculated using a standardised scenario based on all possible exposure pathways for a residential situation. The exposure routes are shown in figure A5.4. CSOIL is also used for derivation of remediation objectives, determination of the urgency of remediation and calculation of site-specific exposure.

Current IVs and indicative values are available in VROM (2000) although more recent review has resulted in proposals for new IVs (Bars et al, 2001; Lijzen et al, 2001) for what are known as first-series compounds (a group of most important compounds). However, these values have yet to be adopted into law (the Soil Protection Act). There are also second, third and fourth series of compounds currently being reviewed. These were originally developed by Crommentuijn et al (1994), Kreule et al (1995) and Kreule and Swartjes (1998), and still apply as legal intervention values for these contaminants.

The current version of CSOIL is described in detail in Brand et al (2007).



Figure A5.4: Schematic lay-out of the CSOIL exposure model

Source: Brand et al, 2007.

Calculations by Otte et al (2001) determined that three of the exposure routes are responsible for at least 90 per cent of the total exposure for almost all compounds. These routes are the human exposure:

- via the ingestion of contaminated soil particles
- to volatile compounds in the indoor air
- via the consumption of contaminated crops.

The other most significant pathways, depending on contaminant are:

- dermal uptake via soil contact (1–7 per cent for 18 compounds)
- drinking-water intake due to permeation through plastic pipes (1–13 per cent for 29 compounds)
- dermal uptake during bathing (1–5 per cent for 20 compounds).

Although not every exposure route has a significant contribution to the total human exposure, the basic principle of the model is that all possible exposure routes are taken into account.

The model concept consists of roughly three parts:

- 1. the description of the behaviour of the compound in the soil and the partitioning over the soil phases
- 2. the transfer processes and parameterisation of the different exposure routes (direct and indirect)
- 3. the quantification of the lifetime average exposure (Otte et al, 2001).

Details of the model are too complex to describe here, however, in common with all such models, there are a number of input parameters such as:

- compound-specific input parameters; mainly physicochemical properties
- site and soil properties, related to potential exposure
- exposure parameters which describe the receptor characteristics and behaviour such as breathing volume or ingestion frequency (Otte et al, 2001).
A5.8 New Zealand

A5.8.1 Policy and legislative framework

The New Zealand policy and legislative framework is outlined in MfE (2006b) and will not be repeated here. The document is available on MfE's website at: http://www.mfe.govt.nz/issues/hazardous/contaminated/direction-land-management.html.

A5.8.2 Industry-based guidelines

Four industry-based guideline documents have been developed by the Ministry for the Environment to provide good practice guidance for risk assessment of four types of site considered as a priority because of their prevalence in New Zealand. The guidelines have not been developed with any statutory basis, but as a matter of administrative expediency: to assist local government and contaminated-site practitioners to assess the risk posed by sites. Adoption of the guidelines is voluntary, unless referred to in operative district or regional plans, in which case they have regulatory force. The guidelines are ostensibly industry-based. Nevertheless they can be, and routinely are, used to assess the particular contaminants within each guideline on other sorts of sites that may happen to have the contaminants. The four guidelines are:

- *Health and environmental guidelines for selected timber treatment chemicals* (MfE and MoH, 1997), (the 'Timber Treatment Guidelines'). These present soil and water criteria for the timber preservation chemicals copper, chromium, arsenic, boron and pentachlorophenol. An interim guideline for dioxins was also adopted (as represented by 2,3,7,8- tetrachlorodibenzo-p-dioxin) but no details were given to support its derivation.
- *Guidelines for assessing and managing contaminated gasworks sites in New Zealand* (MfE, 1997) (the *Gasworks Guidelines*), which present human-health soil acceptance criteria for selected polycyclic aromatic hydrocarbons (PAHs), the mono-aromatic hydrocarbons benzene, ethyl benzene, toluene and xylene (collectively, BTEX), the phenol compounds phenol and cresol, and inorganic cyanide compounds in both free and complexed forms.
- *Guidelines for assessing and managing petroleum hydrocarbon contaminated sites in New Zealand* (MfE, 1999), (the 'Oil Industry Guidelines'). This presents human health-guideline values for soil and groundwater and, indirectly for indoor and outdoor air, for petroleum hydrocarbons (represented as three carbon ranges of mixed aliphatic hydrocarbons), the BTEX group of mono-aromatics and the PAHs naphthalene, benzo(a)pyrene and pyrene. Guidelines are presented for several different generic soil texture types, to account for the different vapour migration behaviour with different soil texture (clay, silt, sand, etc).
- Identifying, investigating and managing risks associated with former sheep-dip sites: A guide for local authorities (MfE, 2006a) (the 'Sheep-dip Guide'), which presents human-health guideline values for arsenic (as derived in the TTG, above) and the chlorinated pesticides DDT (as the sum of DDE, DDD and DDT ∑DDT), dieldrin (including aldrin) and lindane.

All these guidelines draw on principles set down by the US EPA (US EPA 1989a; 1991a; 1996a; 1996b) in developing values primarily for the protection of human health. The 'Timber Treatment Guidelines' methodology, perhaps because it was the first and generally more detailed document in terms of derivation methodology of the four guidelines, is generally considered by practitioners to be current New Zealand policy with respect to derivation. The other New Zealand guidelines have similar derivations but with some significant differences that would result in different values for the same substances (and do result in different values for a few organic compounds that are in common). The detail of the 'Timber Treatment Guidelines' is presented below, with differences of the other guidelines presented in separate sections, where relevant.

Background exposure is generally not explicitly included in the methodology provided in the existing New Zealand guidelines, and typically 100 per cent of the TDI is assigned to exposure from soil sources, following US EPA practice. An exception is for copper (*Timber Treatment Guidelines*), where only 10 per cent of the TDI is assigned to soil sources "due to the relatively high intake of copper from other sources" (MfE and MoH, 1997). This appears to be overly conservative in relation to exposure from soil sources.

The underlying premise in existing New Zealand industry-based guidelines is that protection of on-site ecosystems is only required to the extent necessary to facilitate the use of the land, eg, plant growth and livestock grazing. The 'Timber Treatment Guidelines' provide numeric values for the protection of plant life and livestock health. Some of the values that have been adopted (eg, for copper) are specifically based on protection of plant life as opposed to human health. The 'Oil Industry' and 'Gasworks Guidelines' consider that the nature of the contaminants (ie, volatile, readily degradable) is such that soil guideline values based on human health will also protect plant and livestock health.

The exposure scenarios used for the derivation within existing New Zealand guidelines is summarised in table A5.3 and the detailed description are given in following sections.

Guideline	Scenario			
Timber Treatment	Agricultural / horticultural			
Guidelines	Residential			
	Industrial – paved, unpaved			
	Subsurface maintenance workers			
Gasworks Guidelines	Agricultural / horticultural			
	Standard residential			
	High-density residential			
	Commercial / industrial			
	Parkland / recreational			
	Maintenance workers			
'Oil Industry Guidelines'	Agricultural / horticultural			
	Residential			
	Commercial / industrial			
	Maintenance			
Sheep-dip Guide	Lifestyle block			
	Standard residential			
	High-density urban residential			
	Parks / recreation			
	Commercial / industrial			

Table A5.3: Exposure scenarios in existing New Zealand industry-based guidelines

For a given exposure scenario, the estimated exposure – and therefore the derived soil numeric value – is dependent on the exposure pathways applicable for that scenario, and the parameters (eg, soil ingestion rate, exposure frequency) selected to define the extent of exposure via those pathways.

Timber treatment guidelines

The 'Timber Treatment Guidelines' (MfE and MoH, 1997) provide for four exposure scenarios with the residential and commercial / industrial pathway each having two subscenarios. These allow for different proportions of home-grown produce consumption and paved and unpaved commercial / industrial sites. The scenario definitions are:

- **Agricultural** includes all agricultural and horticultural uses, particularly those involved in the production of food for human consumption. The general public is protected by ensuring that soil contamination would not give rise to concentrations of contaminants in produce that would pose a concern to public health. The health of residents at any farm property is also considered, assuming that residents may be exposed via the consumption of home-grown livestock and produce, and through direct contact with the contaminated soil, eg, ingestion of contaminated soil.
- **Residential** includes low-density residential use and rural residential use, where a considerable proportion of the total amount of produce consumed may be grown at the site. If livestock for human consumption are kept at a site then it should be assessed against the agricultural criteria, in the first instance. The small size of many residential developments within urban areas limits the amount of produce that may be grown, reducing the potential exposure for some contaminants. Recommended acceptance criteria have been derived for two rates of home produce consumption, reflecting the differences between urban residential use and rural residential use.
- **Commercial / industrial** reflects exposure conditions at a largely unpaved industrial site where workers may come into direct contact with contaminated soil. This scenario is not designed to include consideration of workers actively involved in excavation or similar activities, for which separate criteria are derived. Where a site is largely paved, higher contaminant concentrations may be acceptable, as outlined in the guidelines.
- **Maintenance** for each of the above site uses, human exposure to ground contamination may be associated with subsurface maintenance works, eg, repair and replacement of services. While the duration of such works is generally much shorter than the other exposure scenarios considered, the rate of exposure is likely to be much higher and this may be significant where the work is undertaken routinely by the same person.

The 'Timber Treatment Guidelines' provide a set of equations to estimate the chronic daily intake of individual contaminants for four exposure pathways: soil ingestion, produce consumption, inhalation of particulates, and dermal absorption.¹⁹ These equations are based on the same generic equations described previously and follow US EPA practice.

Intake = <u>concentration × contact rate × exposure frequency × exposure duration</u> averaging time × body weight

¹⁹ The equations in the 'Timber Treatment Guidelines' are in error, as presented in Chapter 5 of the document, but the correct equations are used in the derivation calculations. This error is separate from other errors in the produce pathway calculations. Details are given in Cavanagh and Proffitt (2005).

The pathway-specific equations for threshold contaminants are, using the nomenclature of the document:

Soil ingestion:		ion	$CDI = \frac{C_{S} \times IR \times CF \times ED \times EF \times MF}{AT \times BW}$
Produce ingestion:			tion: $CDI = \frac{C_P \times IP \times ED \times EF \times Pg}{AT \times BW}$
Inhalation of dust:			lust: $CDI = \frac{C_S \times IH \times ED \times EF \times MF \times R}{AT \times BW \times PEF}$
Derr	nal ab	sorj	ption: $CDI = \frac{C_S \times CF \times AR \times AH \times AF \times ED \times EF}{AT \times BW}$
$\begin{array}{rcl} CDT &= & chrome daily intake in terms of body weight (hig/kg B w/day)\\ C_{s} &= & contaminant concentration in soil (mg/kg)\\ IR &= & soil ingestion rate (mg/day)\\ CF &= & conversion factor (10^{-6} kg/mg)\\ ED &= & exposure duration (years)\\ EF &= & exposure frequency (days/year)\\ MF &= & matrix factor, typically set to 1\\ AT &= & averaging time ED \times 365 days for a threshold substance, or 70\\ & for non-threshold\\ BW &= & body weight (kg) \end{array}$			
IP = produc $Pg = propor$ $IH = inhalat$ $R = propor$ $PEF = particle$ $AR = expose$ $AH = soil ad$ $AF = absorp$			produce ingestion rate expressed as dry weight (kg DW/day) proportion of home-grown produce (dimensionless) inhalation rate (m^3 /day) proportion retained in lungs (dimensionless) particle emission factor (m^3 /kg) exposed skin surface area (cm ²) soil adherence factor (mg /cm ²) absorption factor (dimensionless)

For threshold substances, there is a single critical receptor and the exposure duration in days is the averaging time, ie, averaging time = $ED \times 365$. ED in the numerator and denominator of each equation cancel and the contact rate is just that for the critical receptor, with, for example, the equation for the soil ingestion pathway becoming:

$$CDI = \frac{C_{S} \times IR \times CF \times EF \times MF}{365 \times BW}$$

The equations for the other three exposure pathways are similar.

For non-threshold substances there is no critical receptor, as exposure at any time could cause an adverse effect. Using one or more receptor groups to better represent varying intake rates, body weights and other age-dependent parameters, the total chronic intake is obtained by summing over a total period of exposure for the different receptor groups.

For each exposure pathway the total intake becomes:

 $CDI = CDI_1 + CDI_2 + CDI_3 + ... CDI_n = \sum CDI_i$

where i represents each receptor age group.

It is possible to have all the various parameters being age-dependent but in the US EPA protocol that New Zealand adopted at the time, exposure frequencies for each age range are taken to be common across all age ranges. Similarly, dermal soil adherence has been taken as common to all age groups (no longer the case in US EPA practice). Thus, by substituting the pathway exposure equations into Equation 3.9 and collecting common terms, only the exposure rate, exposure duration and body weight remain in the summation, as an 'age-adjusted' rate, with, using soil ingestion as an example, the total chronic intake rate becoming:

$$CDI = \frac{C_{s} \times IR_{adj} \times CF \times EF \times MF}{AT}$$

With IR_{adj} being represented by:

$$IR_{adj} = \sum \frac{I_i \times ED_i}{BW_i}$$

where

Σ signifies summation over receptor groups i = 1 to n

- Iri = soil ingestion (or produce ingestion rates, inhalation rates or skin surface areas, as appropriate for each of the equations, for receptor group i
- BW_i = body weight for receptor group i.

Following US EPA practice for generic guidelines, by convention the averaging time for nonthreshold substances is a 70-year lifetime and there are two age ranges: a child aged 1–6 and an adult aged 7–30. Total exposure therefore becomes 30 years and this duration is divided by 70.

The 'Timber Treatment Guidelines' combine the pathway-specific guideline values using an equation of the following form. This follows from the total exposure from each pathway being additive.



The generic equations above, except the final equation for combining pathways, are used in all New Zealand guideline documents, although there can be slight differences in their application. This is discussed in subsequent sections.

Subsidiary equations are required for each of the produce exposure and inhalation equations, to determine contaminant concentrations in produce in terms of soil concentration and to determine a particle emission factor for calculating dust inhalation. Vapour inhalation is not considered in the Timber Treatment Guidelines.

Calculation of produce concentrations uses bioconcentration factors to relate soil concentration to produce concentration.

 $C_P = BCF \times C_S$

Where: C_P = concentration in produce

BCF = bioconcentration factor, expressed as the ratio of the contaminant content in produce (mg/kg dry weight) and soil (mg/kg dry weight)

 C_s = concentration in soil

In addition, the 'Timber Treatment Guidelines' divide produce into root and leafy vegetables, with the root BCF taken to be five times that of the 'Timber Treatment Guidelines' leafy BCF. Fruit are ignored on the basis that contaminants do not significantly translocate to fruit. In the calculations, fruit is ignored not by ignoring the fruit component of the typical diet – the totally daily produce consumption (kg/day) is specified with fruit included – but by only using the proportions of the total for leafy and root vegetables. The two proportion factors sum to less than 1, in effect reducing the total daily produce consumption. The same effect could have been achieved by specifying only the daily vegetable intake, with the two proportions summing to 1.

The detail of the equations is not provided in the 'Timber Treatment Guidelines', but it is apparent from the 'Oil Industry Guidelines' (which adopt a similar approach) that an average concentration in produce is calculated from:

$$C_{P} = P_{leafy} \times C_{P \ leafy} + P_{root} \times C_{P \ root}$$

Where: the subscripts leafy and root refer to above-ground edible vegetation and roots, respectively

P is the proportion of leafy or root vegetables

 $C_{P \text{ leafy}}$ and $C_{P \text{ root}}$ are related to $C_{S \text{ by}}$ bioconcentration factors (BCF) for leafy parts and roots, respectively, ie, using $C_{P \text{ leafy}} = BCF_{\text{leafy}} \times C_S$, and similar for roots

Substituting in the BCFs and rearranging in order to represent C_P in terms of C_S should yield:

$$C_{P} = C_{S} (BCF_{root} \times P_{root} + BCF_{leafy} \times P_{leafy})$$

However, as Cavanagh (2004b) pointed out, the 'Timber Treatment' and 'Gasworks' guidelines incorrectly used an equation equivalent to:

$$C_{P} = C_{S} (BCF_{root} \times P_{leafy} + BCF_{leafy} \times P_{root})$$

The actual effect of this error is small because P_{leafy} and P_{root} have similar values, being 31 per cent for leafy vegetables (referred to as 'stem' in the guideline) and 29 per cent for root vegetable based on 1985 Australian dietary surveys (MfE and MoH, 1997).

Typically the BCF for metals is based on field or laboratory experiments, while the BCF for organics is often estimated from the octanol-water partition coefficient (K_{ow}). For metals, the 'Timber Treatment Guidelines' relate BCF_{root} to a soil-plant distribution coefficient, K_d (unit: ml/mg):

$$0.85 \ln BCF_{root} = 3.02 - \ln K_d$$

This is attributed to ECETOC (1990); however, as Cavanagh (2004b) has noted, ECETOC (1990) indicates this equation is applicable to above-ground parts not root. As BCF_{root} is assumed to be five times that of BCF_{leafy} in the *Timber Treatment Guidelines*, the effect is an underestimation of both BCFs by a factor of five (assuming the equation above is valid for the particular metals). A further error pointed out by Cavanagh (2004b) is that the BCF (and produce concentration) are expressed as dry weight, but the amount of produce consumed is expressed as wet weight, requiring BCFs expressed on a dry-weight basis to be converted to are wet-weight basis (or alternatively, the amount of produce consumed to be converted to dry weight). This correction does not appear to have been undertaken for the inorganic contaminants.

For organic compounds (only pentachlorophenol is considered) the 'Timber Treatment Guidelines' use an empirical relationship developed by Travis and Arms (1988) to describe the uptake into above-ground parts, based on the octanol-water partition coefficient (K_{ow}). It should be noted that the US EPA (2003) has criticised the Travis and Arms relationship as being based on few data, some of which are at variance from the source documents they cite.

 $log \; BCF_V = 1.588 \text{--} 0.578 \; log \; K_{\rm ow}$

where: $BCF_V =$ bioconcentration factor for vegetation $K_{ow} =$ octanol-water partition coefficient

 BCF_v is based on the dry weight of vegetation, but as noted early, the guidelines assume the fresh weight concentration can be estimated using a moisture content of 80 per cent (ie, divide by a factor of 5). No justification is given for this moisture content.

Finally, the values derived as above are not necessarily the values adopted in the *Timber Treatment Guidelines*. Protection of plant health is also considered and the value adopted is the lowest for protection of human health or plant health. Plant health values were only adopted for the agricultural and residential scenarios, but in some cases this results in very much lower guidelines than would have been the case if only human health had been considered. The plant health values have been taken from the literature as a worst-case value for the onset of phytotoxicity for acid (sandy) soils.

The human-health guideline values for copper are unusually low relative to international values because only 10 per cent of the TDI was assigned to soil sources (in other words, 90 per cent was assigned to background intake). This has caused some controversy within the contaminated-land assessment industry (and additional remediation expenditure in some cases, although generally arsenic has driven the assessment for timber treatment sites and orchard land, the guidelines often being applied to the latter site type also).

Gasworks and oil industry guidelines

Two other guidelines, the 'Gasworks' and 'Oil Industry' guidelines, were developed shortly after the 'Timber Treatment Guidelines'. These two guidelines use essentially the same exposure scenarios as before, with an additional scenario, parkland / recreation in the *Gasworks Guidelines*. The new definitions are repeated below, as are clarifications to the agricultural / horticultural and residential scenarios (clarifications shown in italics below) contained within these documents.

- Agricultural / horticultural deemed to include all agriculture and horticulture, particularly those related to food production. The general public is protected by ensuring that soil contamination does not give rise to a concentration in produce *that exceeds a published maximum residue level (MRL), although MRLs have not been nominated for most contaminants of concern. Consideration is given to the risk associated with consuming 100 per cent of produce from a contaminated source.* Consideration is also given to protecting the health of residents at any farm property, assuming that residents may be exposed by consuming home-grown livestock and produce, and through direct contact with contaminated soil. It is assumed that residences do not incorporate basements.
- **Standard residential** based on low-density residential use, including rural residential use, where a considerable proportion of the total amount of produce consumed is grown at the site. *No consideration is given to livestock uptake of contaminants. It is assumed that residences do not incorporate basements.*

- **High-density residential** assumes there are limited soil access opportunities, therefore there is significantly less soil and dust exposure by ingestion compared with a standard residential site. This scenario does not include consuming of produce grown at the site. (Values derived in 'Gasworks Guidelines' only, but commentary provided in 'Oil Industry Guidelines' for deriving values from generic pathway values).
- **Commercial / industrial** as for 'Timber Treatment Guidelines'.
- Maintenance workers as for 'Timber Treatment Guidelines'.
- **Parkland / recreational** ('Gasworks Guidelines' only) reflects shorter exposure times but potentially on a regular basis. Opportunities for contact with soil will arise and children are the key concern in these areas.

While the base exposure scenarios are essentially the same, the 'Oil Industry Guidelines' examine a number of more complex pathway scenarios for each exposure scenario than the 'Gasworks Guidelines', involving the inhalation pathway to indoor and outdoor air for different soil types. This results in very many more soil guideline values for each contaminant, and makes for quite a complex document. In addition, the 'Oil Industry Guidelines' contain soil guideline values for the protection of groundwater. Examination of this pathway is beyond the scope of this report.

Previous reviews (Cavanagh, 2004b, 2004c, 2004d, 2005a; Cavanagh and Proffitt, 2005) have identified differences and inconsistencies between the earlier 'Timber Treatment Guidelines' and the subsequent 'Gasworks' and 'Oil Industry' guidelines. Rather than deriving a combined value from each of the relevant human health exposure pathways, as described in the previous section, both the 'Gasworks' and 'Oil Industry' guidelines take the lowest value derived from the individual human health pathways; however, this is not adhered to consistently with the 'Gasworks Guidelines' as some values appear to have been derived by combining pathways. Not combining pathways is a less conservative approach than combining the values and it is not clear why this method has been chosen; it may have been because the inhalation pathway for some volatile organic compounds and the produce ingestion pathway for some polycyclic aromatic hydrocarbons produce very low values. Combining these with the soil ingestion would have resulted in even lower, and potentially untenable, values. In other instances, there is clearly a much lower value for a particular pathway; with the other pathways have only a small effect if pathways had been combined.

Ecological receptors are considered separately in a qualitative way in both of these guidelines.

As mentioned above, the 'Oil Industry Guidelines' provide guideline values for several different soil types. However, soil type only influences the values derived for the inhalation pathway for volatile substances. Where vapour inhalation is not an exposure route, or not critical, the adopted guideline is the same regardless of soil type.

Further differences occur in the derivation of soil guideline values related to the produce consumption pathway and partitioning of volatiles to indoor air. For the produce consumption pathway, the 'Gasworks Guidelines' use the same methodology (including the errors) and assumptions used in the 'Timber Treatment Guidelines' to derive soil guideline values for the produce-consumption pathway, with the exception that 50 per cent of the produce consumed are vegetables and all are assumed to be root crops. The source of percentage of vegetables consumed is Langley (1993) using Australian data. Plant uptake factors for organics are predicted using the Travis and Arms (1988) relationship.

The 'Oil Industry Guidelines' are also different from the 'Timber Treatment Guidelines' with respect to produce uptake. Notably, the equation for determining soil concentrations from produce concentration is correct (MfE 1999: appendix 4F), and instead of Travis and Arms (1988), the model of Ryan et al (1988) is used to estimate plant uptake of contaminants. This model determines a fresh-weight plant uptake factor for roots and stems from contaminant concentrations in pore water. However, the 'Oil Industry Guidelines' do not indicate the relationship between pore water concentrations and soil concentrations, which become the soil acceptance criteria. The 'Oil Industry Guidelines' also assume a different composition of root and leafy vegetables to that in both the 'Gasworks' and 'Timber Treatment' guidelines, notably 10 per cent are assumed to be root vegetables, 50 per cent leafy (stem) vegetables or fruit, and 40 per cent tree-fruits, where tree-fruits are assumed not to take up contaminants.

All three guidelines assumed that 100 per cent of produce consumed was home-grown for the agricultural scenario to be protective of farming families and also, by default, consumers of farm produce. This was later considered to be unrealistic for the agricultural scenario, and was explicitly changed by the time the 'Sheep-dip Guide' was issued. Guideline values were derived for two standard residential scenarios: for 50 per cent and for 10 per cent home-grown produce, the latter suggested for larger urban areas. The higher percentage was routinely adopted in the early days of the 'Timber Treatment Guidelines' but by the time the later two guidelines had been issued, the 10 per cent home-grown produce was generally accepted as the standard urban scenario regardless of location. For example, while both 10 and 50 per cent produce values are presented in route-specific tables within the 'Oil Industry Guidelines', in the most commonly used residential 'All Pathways' tables it is the 10 per cent home-grown produce values that are given when the produce pathway is critical, rather than the 50 per cent values. This suggests an official acceptance that the 10 per cent values were the residential default.

Few details are given within the 'Gasworks Guidelines' with respect to the vapour intrusion into buildings, other than that a volatilisation factor was determined by modelling. Two scenarios were considered, surface soil (<1 m) and subsurface soil, and one soil type a sand / sandy loam, on the presumption that that would be conservative. Mention is made of the 'Oil Industry Guidelines' within the section on volatile modelling and it is presumed a similar approach was adopted, although lower values have been derived within the 'Oil Industry Guidelines' for the same soil type and contaminant.

The 'Oil Industry Guidelines' used a modified form of the Jury et al (1983, 1984) model for migration of vapours from the soil to the outdoor air. For migration into buildings the Johnson and Ettinger (1991) model was used. As noted previously, this model was developed for migration into buildings with concrete slab-on-grade floor, or into basements. It is not applicable to the common New Zealand construction of a house on piles with a crawl space below the floor, although Davis et al (2008) report that the model has recently been modified to cope with crawl-space construction.

The completely underground basement, a common residential feature in the United States, is uncommon in New Zealand for standard low-density residences. Further, the assumption within the Johnson and Ettinger model is of a perimeter crack between the floor and walls. However, in New Zealand it is common for the concrete floor to be poured integrally with the perimeter footings (typically just a thickening of the floor) with no perimeter crack being present, with timber framed walls being fixed to the concrete. In such construction vapours would have to go to the outside of the building before migrating through walls and/or under the wall/floor connection, or migrate through floor penetrations, the latter not apparently allowed for in the Johnson and Ettinger model. Further, the 'Oil Industry Guidelines' assume a crack to floor area ratio of 0.01 (1 per cent).

In comparison, the US EPA default recommendation of a 1 mm crack width (US EPA, 2004b) translates to a crack ratio of 0.38 percent for a typical slab-on-grade house in the US, the 2 mm default crack width in UK's CLEA model (EA, 2004a, 2004b), translates to a crack ratio of 0.1 per cent, only one-tenth of the assumption in the 'Oil Industry Guidelines', while the Dutch VOLASOIL model – concerning soils contaminated with volatile compounds – (Waitz et al, 1996) uses a default crack ratio of 0.01 per cent for a 'bad' floor and 0.001 per cent for a 'normal' floor. Using the 'Oil Industry Guidelines' value, a 1 per cent crack ratio for a 10 m × 15 m building (150 m²) is 1.5 m² of floor openings and would mean a perimeter crack 30 mm wide, clearly unrealistic for even poor construction.

Sheep-dip guide

The general methodology provided in the 'Timber Treatment Guidelines' (MfE and MoH, 1997) and the 'Oil Industry Guidelines' (MfE, 1997) was used for soil guideline values in the 'Sheep-dip Guide' (MfE 2006a).

In contrast to the earlier guidelines, the 'Sheep-dip Guide' explicitly includes lifestyle-block land use as a typical New Zealand land use, rather than the agricultural or horticultural scenarios. This scenario assumes that 50 per cent of the produce consumed by residents is grown on site, but consumption of meat, milk and eggs of animals raised on site is excluded. Previously this land use has been a subset of residential land use (high home-grown produce in the *Timber Treatment Guidelines*). The standard residential land-use category within the 'Sheep-dip Guide' assumes that 10 per cent of the produce consumed by residents is grown on site, while the remaining categories do not consider consumption of produce grown on site.

The following five land-use categories were adopted.

- Lifestyle block residential property where 50 per cent of vegetables consumed are assumed to be grown on site. The consumption of products (eggs, milk, meat) from animals raised on site is excluded and should be considered on a site-specific basis.
- **Standard residential** low-density residential property with home-grown vegetables contributing 10 per cent of the total intake.
- **High-density urban residential** residential with minimum opportunity for exposure to soil; no produce consumption; includes daycare centres, kindergartens, preschools and primary schools, where no gardens are present.
- **Parks / recreation** parks, recreational open space, playing fields; includes secondary schools.
- **Commercial / industrial** (unpaved) unpaved commercial and industrial properties. Where paving is present, its integrity and likely effectiveness in reducing exposure must be considered on a site-specific basis. No consideration of the protection of plant life has been included.

The exposure scenarios considered are largely based on those provided in the *Gasworks Guidelines*, while the parameter values used are based on those contained in both the *Timber Treatment* and *Gasworks Guidelines*. However, different parameters were used for the dermal exposure and produce consumption pathways. The equations used to derive the soil guideline values were as provided in Cavanagh and Proffitt (2005), which are essentially corrected versions of those appearing in the *Timber Treatment Guidelines*. In addition, the produce consumption values were derived using a weighted average of the consumption of different types of vegetables, using fresh weight to dry weight conversions for each. This approach, while

not making a large difference to the derived soil guideline values, is technically more robust and uses New Zealand estimates for daily vegetable consumption, rather than Australian estimates.

The 'Sheep-dip Guide' is important in clarifying that the standard urban residential scenario had only a small percentage of home-grown produce consumption (chosen to be 10 per cent), whereas with the 'Timber Treatment Guidelines' there was initial (and perhaps continuing) confusion amongst practitioners as to whether a large percentage (50 per cent) or a small percentage (10 per cent) should be chosen as the urban residential default. In addition, the dropping of the agricultural scenario (which used 100 per cent home-grown produce), which was intended to represent a rural residential scenario (eg, a farm), in favour of the lifestyle block scenario, resulted in a home-grown produce percentage that was more likely to be representative of an average to high-end for most rural residents. Dropping the agricultural scenario also removes consideration of having to protect the productive capacity of the land (by setting phytotoxicity based values) and not exceeding food maximum residual levels; both are considerations that go beyond protection of site users.

A5.8.3 Summary

The basic methodology of the New Zealand guidelines compares well with the overseas guidelines reviewed. This is not surprising given their US EPA origins. While less complex than both, the Dutch and UK derivation methodology, the added complexity provides dubious advantages given the uncertainty of calculating some of the primary pathways (eg, plant uptake and indoor inhalation of volatiles). The supposed additional refinement offered for some pathways and/or the addition of minor pathways will tend to be overshadowed by the uncertainties in the main pathways. The bigger advantage is possibly to remain relatively simple, for generic guideline value derivation, with the greater complexity reserved for site-specific assessment if warranted on a case-by-case basis.

It is appropriate to persevere with the US EPA-based general 'Timber Treatment Guidelines' methodology, as recommended by the Technical Review Group for the NES (MfE, 2005). However, it is clear that the inconsistencies and errors need to be corrected and a greater use be made of New Zealand-specific scenarios and parameters where possible.

The findings of this comparison have been incorporated into the *Methodology for Deriving Standards for Contaminants in Soil to Protect Human Health.*

Appendix 6: Natural Background Topsoil Datasets for Arsenic and Cadmium

A6.1 Introduction

The toxicological and technical advisory group recommended that:

"Where SGVs for cadmium and arsenic are exceeded by the 99th percentile of the arsenic and cadmium national natural background dataset, then the SGV will be adjusted to align with it."

In accordance with this recommendation:

- 1. The arsenic soil guideline value for the rural residential (25 per cent produce consumption) land-use scenario, as determined in table A6.5 of this document has been adjusted as the value **is exceeded** by the 99th percentile of the arsenic national natural background dataset.
- 2. The cadmium soil guideline value for the rural residential (25 per cent produce consumption) land use scenario, as determined in table A6.6 has not been adjusted as the value is not exceeded by the 99th percentile of the cadmium national natural background dataset.

This appendix contains the national background topsoil datasets for arsenic and cadmium and summary statistics of the datasets including the 99th percentile. Also included is a description of the method used to prepare the datasets, the data sources used, and data limitations. A regional breakdown of background topsoil concentrations for arsenic and cadmium is also provided for comparative purposes. The text and analysis in this appendix is mostly extracted from an unpublished paper (Moore & Taylor, 2009) presented at the WasteMINZ 2009 Annual Conference.

A6.2 Arsenic topsoil dataset

A6.2.1 Method used to prepare the dataset

Preparation and manipulation of the national background dataset followed the methodology in Soil Maps of Cadmium in New Zealand (Taylor et al, 2007). Topsoil soil samples data was collated from the sources outlined in table A6.1. Sampling strategy and protocol varied with the purpose of sample collection.

Four hundred and twenty-three samples were considered for natural background classification as they came from sites with land uses recorded as reserve, parks, schools, native forest, urban background or background. Upon screening for outliers and data anomalies a selection of 385 of these samples were appropriate to derive the average natural background level of arsenic in New Zealand topsoils.

Many of the samples in the database were collected for soil quality monitoring or state of the environment reporting and sampling was based on composite sampling (Hill & Sparling, 2009; Kim & Taylor, 2009; Sparling et al, 2004). Other samples were collected to determine regional background concentrations for arsenic and other elements or to determine the status of pollutant issues.

Several samples were recorded as none detected in the original data sources (<2 mg/kg). For data analysis these were incorporated into the dataset as half the detection level.

Region	Original data source
Auckland	Background Concentrations of Inorganic Elements in Soils from the Auckland Region, ARC Technical Publication No.153
SG Auckland	Rural soil reports
SG Tasman	TDC Rural Soil Reports Final June 2003
Marlborough	Soil quality monitoring
SG Waikato	Rural Soil Reports
Waikato	Concentrations of various elements in Waikato surface soils and suggested default cleanfill thresholds. Kim, 2005
Hawke's Bay	SOE soil monitoring data 2007
Wellington	Annual soil quality monitoring
URS Wellington	URS report for Greater Wellington (2003) Determination of common pollutant background soil concentrations for the Wellington Region
Canterbury	Background concentrations of selected trace elements in Canterbury soils Report No. R07/1 Prepared for Environment Canterbury by Tonkin and Taylor Ltd July 2006
Bay of Plenty	Soil quality monitoring
NZ wide Longhurst et al, 2004	Concentrations of arsenic, cadmium, copper, lead, and zinc in New Zealand pastoral topsoil and herbage. New Zealand Journal of Agricultural Research 2004, Vol. 47: 23–32

Table A6.1: Regional council supplied data

A6.2.2 National assessment of natural background concentrations for arsenic

The national natural background dataset for arsenic in topsoil is provided in table A6.5. The summary statistics from the national dataset are shown in table A6.2 and broken down by region in table A6.3.

The 99th percentile of the national natural background concentrations for arsenic is 17.4 mg/kg.

Regional assessment of natural background arsenic topsoil concentrations showed no statistical difference (p < 0.05) between regions (see table A6.3), however, the small samples size (<30) for some regions makes it difficult for statistical differentiation.

Summary statistics						
Count	372					
Mean mg/kg	4.5					
Median mg/kg	3.7					
Minimum mg/kg	0.4					
Max mg/kg	0.77					
Upper 95 percentile mg/kg	10.0					
Upper 97.5 percentile mg/kg	11.8					
Upper 98 percentile mg/kg	16.0					
Upper 99 percentile mg/kg	17.4					

Table A6.2: National statistics of the natural arsenic background concentration in topsoil

Table A6.3: Statistics of the natural arser	nic background concentration in topsoil by
region	

Region	Count	Mean (mg/kg)	Median (mg/kg)	Minimum (mg/ kg)	Max (mg/kg)
Auckland	96	4.4	4.0	0.4	11.5
Bay of Plenty	17	5.9	5.0	1.0	16.0
Canterbury	73	4.3	3.0	0.9	36.9
Hawke's Bay	26	4.8	3.9	1.1	11.5
Marlborough	6	3.7	3.5	2.0	5.0
Waikato	38	4.7	3.8	0.7	21.9
Wellington	80	4.1	4.0	1.0	12.3

A6.3 Cadmium topsoil dataset

A6.3.1 Method used to prepare the Cadmium dataset

Table A6.6 presents the national natural background dataset for cadmium in topsoil. Data of background topsoil samples were collated from Soil Maps of Cadmium in New Zealand (Taylor et al, 2007) and additional data collected by regional councils since this report. Data for 486 samples were used to derive natural background levels of cadmium in New Zealand topsoil. These samples were collected from sites with park, native, tussock and unfertilised land uses.

Several samples were recorded as none detected in the original data sources (<0.1, <0.2, <0.3 mg/kg). For data analysis these were incorporated into the dataset as half the detection level. No integration of data sources or other collation methodology has been undertaken for this dataset.

A6.3.2 National assessment of natural background concentrations for cadmium

The natural background dataset for cadmium is provided in table A6.6. The summary statistics from the national dataset are shown in table A6.4.

The 99th percentile of the national natural background concentrations for cadmium is 0.65 mg/kg.

Summary statistics						
Count	486					
Mean mg/kg	0.15					
Median mg/kg	0.1					
Minimum mg/kg	0.01					
Max mg/kg	0.77					
Upper 95 percentile mg/kg	0.47					
Upper 97.5 percentile mg/kg	0.60					
Upper 98 percentile mg/kg	0.601					
Upper 99 percentile mg/kg	0.65					

Table A6.4: Summary statistics of the national natural background cadmium topsoil dataset

A6.4 Data limitations

The following limitations in the national arsenic and cadmium background datasets were identified:

- 1. Variation in the sampling and analysis methodology used by different councils and Crown research institutes.
- 2. Poor land-use classification.
- 3. Representative data could not be obtained for large areas of New Zealand and sample density was concentrated in Auckland, Waikato, Wellington and Canterbury. Natural geological processes, such as geothermal activities and processes associated with gold seams, are known to occur in some regions of New Zealand, which may result in the formation of hotspot areas of high arsenic concentrations. Regions that were able to contribute few or no samples include Gisborne, Manawatu-Wanganui, Marlborough, Nelson, Northland, Otago, Southland, Taranaki and Westland.
- 4. Variation due to soil type is known to occur (eg, higher arsenic concentrations occur in peat rich topsoil (Kabata-Pendias & Pendias, 2000), but this was not investigated due to the limited number of data points available and the quality of supporting information.
- 5. The effect of soil bulk density has not been considered in collation of this dataset due to limited bulk density data. Taylor & Hill (2010) showed soils of low bulk density can appear to have extremely high contaminant concentrations compared to soils of high bulk density. Without consideration of bulk density, very light organic soil appears to contain extremely high amounts of arsenic and cadmium compared with mineral soil. However, when converted onto a volumetric basis, both organic soil and mineral soil can have similar amounts of arsenic and cadmium.

No.	Data source	Arsenic (mg/kg)	No.	Data source	Arsenic (mg/kg)
1	Auckland	3.68	51	Auckland	5.17
2	Auckland	1.67	52	Auckland	2.29
3	Auckland	1.99	53	Auckland	1.7
4	Auckland	2.02	54	Auckland	6.07
5	Auckland	0.41	55	Auckland	10.16
6	Auckland	1.08	56	Auckland	3.1
7	Auckland	0.48	57	Auckland	3.98
8	Auckland	2.15	58	Auckland	8.2
9	Auckland	7.8	59	Auckland	10.1
10	Auckland	6.61	60	Auckland	10.6
11	Auckland	4.5	61	Auckland	8.3
12	Auckland	3.7	62	Auckland	6.97
13	Auckland	4.8	63	Auckland	3.11
14	Auckland	4.8	64	Auckland	7.07
15	Auckland	1.92	65	Auckland	1.91
16	Auckland	5.82	66	Auckland	3.76
17	Auckland	2.8	67	Auckland	2.75
18	Auckland	4.33	68	Auckland	6.09
19	Auckland	3.55	69	Auckland	7.6
20	Auckland	5.29	70	Auckland	9.11
21	Auckland	0.69	71	Auckland	5.33
22	Auckland	4.14	72	Auckland	8.34
23	Auckland	2.27	73	Auckland	6.73
24	Auckland	1.99	74	Auckland	3.4
25	Auckland	4.12	75	Auckland	4.02
26	Auckland	6.61	76	Auckland	7.44
27	Auckland	1.93	77	Auckland	5.23
28	Auckland	1.6	78	Auckland	3.92
29	Auckland	0.62	79	Auckland	5.49
30	Auckland	7.77	80	Auckland	5.09
31	Auckland	6.85	81	Auckland	7.56
32	Auckland	6.83	82	Auckland	7.22
33	Auckland	8.45	83	Auckland	9.78
34	Auckland	0.78	84	Auckland	4.74
35	Auckland	2.41	85	Auckland	6.98
36	Auckland	2.56	86	Auckland	3.91
37	Auckland	3.17	87	Auckland	8.6
38	Auckland	3.15	88	Auckland	7.01
39	Auckland	2.63	89	Auckland	1.72
40	Auckland	4.57	90	Auckland	2.3
41	Auckland	1.58	91	Auckland	1.19
42	Auckland	2.48	92	Auckland	1.57
43	Auckland	1.36	93	Auckland	3.03
44	Auckland	4.02	94	Auckland	4.88
45	Auckland	2.12	95	Auckland	1.49
46	Auckland	1.93	96	Auckland	4.92
47	Auckland	1.32	97	Bay of Plenty	3
48	Auckland	5	98	Bay of Plenty	2
49	Auckland	11.54	99	Bay of Plenty	3
50	Auckland	2.93	100	Bay of Plenty	1

Table A6.5: Arsenic background sites in native bush

No.	Data source	Arsenic (mg/kg)]	No.	Data source	Arsenic (mg/kg)
101	Bay of Plenty	16	1	151	Canterbury	2.7
102	Bay of Plenty	16		152	Canterbury	2.8
103	Bay of Plenty	5		153	Canterbury	11.5
104	Bay of Plenty	5		154	Canterbury	3
105	Bay of Plenty	6		155	Canterbury	3.7
106	Bay of Plenty	1		156	Canterbury	7
107	Canterbury	2.3		157	Canterbury	5
108	Canterbury	2.7		158	Canterbury	2.9
109	Canterbury	4.3		159	Canterbury	2.2
110	Canterbury	2.5		160	Canterbury	2.8
111	Canterbury	2.5		161	Canterbury	4.3
112	Canterbury	2.3		162	Canterbury	1.7
113	Canterbury	2.5		163	Canterbury	2.9
114	Canterbury	2.5		164	Canterbury	6.1
115	Canterbury	2.3		165	Canterbury	3.9
116	Canterbury	2.1		166	Canterbury	1.6
117	Canterbury	3.3		167	Canterbury	4.6
118	Canterbury	3.9		168	Canterbury	4.1
119	Canterbury	2.6		169	Canterbury	0.9
120	Canterbury	4.6		170	Canterbury	1.8
121	Canterbury	2.7		171	Canterbury	1.7
122	Canterbury	4.6		172	Canterbury	8.7
123	Canterbury	2.3		173	Canterbury	2.1
124	Canterbury	4.1		174	Canterbury	7.8
125	Canterbury	3		175	Canterbury	2.4
126	Canterbury	2.9		176	Canterbury	5.1
127	Canterbury	3		177	Canterbury	4.7
128	Canterbury	3.6		178	Canterbury	1.13
129	Canterbury	3.9		179	Hawkes Bay	6.6
130	Canterbury	4.6		180	Hawkes Bay	4.1
131	Canterbury	3.4		181	Hawkes Bay	4.1
132	Canterbury	2.6		182	Hawkes Bay	7.9
133	Canterbury	3		183	Hawkes Bay	7.5
134	Canterbury	2.9		184	Hawkes Bay	9.9
135	Canterbury	4.2		185	Hawkes Bay	9.6
136	Canterbury	2.9		186	Hawkes Bay	5.4
137	Canterbury	2.2		187	Hawkes Bay	6.2
138	Canterbury	6.2		188	Hawkes Bay	5
139	Canterbury	6.8		189	Hawkes Bay	7.7
140	Canterbury	2.5		190	Hawkes Bay	2.5
141	Canterbury	36.9		191	Hawkes Bay	3
142	Canterbury	4.3		192	Hawkes Bay	3.6
143	Canterbury	2.1		193	Hawkes Bay	3
144	Canterbury	16.9		194	Hawkes Bay	3.3
145	Canterbury	3.7		195	Hawkes Bay	11.5
146	Canterbury	4.1		196	Hawkes Bay	3.1
147	Canterbury	3.8		197	Hawkes Bay	2.8
148	Canterbury	3.1		198	Hawkes Bay	3
149	Canterbury	3.9		199	Hawkes Bay	1.9
150	Canterbury	8.4]	200	Hawkes Bay	1.3

No.	Data source	Arsenic (mg/kg)		No.	Data source	Arsenic (mg/kg)
201	Hawkes Bay	1.7		251	NZ wide Longhurst	2
202	Hawkes Bay	1.1		252	NZ wide Longhurst	2
203	Hawkes Bay	3.4		253	NZ wide Longhurst	4
204	Hawkes Bay	5.4		254	NZ wide Longhurst	3.66
205	Marlborough	5		255	NZ wide Longhurst	0.568
206	Marlborough	4		256	NZ wide Longhurst	1.604
207	Marlborough	3		257	NZ wide Longhurst	3
208	NZ wide Longhurst	4.67		258	NZ wide Longhurst	1
209	NZ wide Longhurst	7.34		259	NZ wide Longhurst	0.835
210	NZ wide Longhurst	5.33		260	NZ wide Longhurst	9.97
211	NZ wide Longhurst	3		261	NZ wide Longhurst	2
212	NZ wide Longhurst	7.268		262	NZ wide Longhurst	17.98
213	NZ wide Longhurst	3.67		263	NZ wide Longhurst	10.64
214	NZ wide Longhurst	3.66		264	NZ wide Longhurst	2.66
215	NZ wide Longhurst	2.33		265	NZ wide Longhurst	2.66
216	NZ wide Longhurst	4.33		266	NZ wide Longhurst	2
217	NZ wide Longhurst	1.129		267	NZ wide Longhurst	2.33
218	NZ wide Longhurst	3		268	NZ wide Longhurst	2
219	NZ wide Longhurst	5		269	NZ wide Longhurst	1
220	NZ wide Longhurst	7.35		270	NZ wide Longhurst	1.33
221	NZ wide Longhurst	3.33		271	NZ wide Longhurst	1.33
222	NZ wide Longhurst	25.73		272	NZ wide Longhurst	1.67
223	NZ wide Longhurst	10.756		273	NZ wide Longhurst	2
224	NZ wide Longhurst	5.66		274	NZ wide Longhurst	3.34
225	NZ wide Longhurst	3		275	NZ wide Longhurst	1.637
226	NZ wide Longhurst	3.34		276	NZ wide Longhurst	9.03
227	NZ wide Longhurst	1.802		277	NZ wide Longhurst	1.33
228	NZ wide Longhurst	5.33		278	NZ wide Longhurst	2.307
229	NZ wide Longhurst	17.304		279	NZ wide Longhurst	2.34
230	NZ wide Longhurst	16.472		280	NZ wide Longhurst	0.767
231	NZ wide Longhurst	5.35		281	NZ wide Longhurst	1.604
232	NZ wide Longhurst	4.34		282	NZ wide Longhurst	2
233	NZ wide Longhurst	3		283	NZ wide Longhurst	5.66
234	NZ wide Longhurst	3.67		284	NZ wide Longhurst	7.34
235	NZ wide Longhurst	0.732		285	NZ wide Longhurst	1.67
236	NZ wide Longhurst	6.01		286	NZ wide Longhurst	0.933
237	NZ wide Longhurst	10.02		287	NZ wide Longhurst	4.33
238	NZ wide Longhurst	5.34		288	NZ wide Longhurst	3.67
239	NZ wide Longhurst	5.33		289	NZ wide Longhurst	1
240	NZ wide Longhurst	2.33		290	NZ wide Longhurst	4
241	NZ wide Longhurst	4.67		291	NZ wide Longhurst	6
242	NZ wide Longhurst	4.67		292	NZ wide Longhurst	5.34
243	NZ wide Longhurst	0.934		293	NZ wide Longhurst	3.68
244	N∠ wide Longhurst	3		294	SG Walkato	7
245	NZ wide Longhurst	2		295	SG Walkato	4
246	NZ wide Longhurst	3.33		296	SG Walkato	8
247	NZ wide Longhurst	4		297		6
248	NZ wide Longhurst	1.33		298		3
249	NZ wide Longhurst	2		299		1
250	IN∠ WIDE LONGNUIST	2.67	J	300	UKS Weilington	5

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No.	Data source	Arsenic (mg/kg)	No.	Data source	Arsenic (mg/kg)
301	URS Wellington	2	344	URS Wellington	4.9
302	URS Wellington	3	345	URS Wellington	5.3
303	URS Wellington	4	346	URS Wellington	4.3
304	URS Wellington	7	347	URS Wellington	3.5
305	URS Wellington	4	348	URS Wellington	4
306	URS Wellington	4	349	URS Wellington	4
307	URS Wellington	3	350	URS Wellington	4
308	URS Wellington	3	351	URS Wellington	1
309	URS Wellington	1	352	URS Wellington	3.8
310	URS Wellington	6	353	URS Wellington	2.6
311	URS Wellington	3	354	URS Wellington	2.6
312	URS Wellington	7	355	URS Wellington	2.4
313	URS Wellington	6	356	URS Wellington	2.5
314	URS Wellington	5	357	URS Wellington	4.1
315	URS Wellington	3	358	Waikato	0.7
316	URS Wellington	7	359	Waikato	1.8
317	URS Wellington	5	360	Waikato	2
318	URS Wellington	5	361	Waikato	2
319	URS Wellington	4	362	Waikato	3.8
320	URS Wellington	3	363	Waikato	4.8
321	URS Wellington	2	364	Waikato	5.8
322	URS Wellington	3	365	Waikato	8
323	URS Wellington	7	366	Waikato	9
324	URS Wellington	5	367	Waikato	9
325	URS Wellington	5	368	Waikato	21.9
326	URS Wellington	5	369	Wellington	10
327	URS Wellington	4	370	Wellington	3
328	URS Wellington	7	371	Wellington	2
329	URS Wellington	1	372	Wellington	6
330	URS Wellington	4	373	Wellington	7
331	URS Wellington	5	374	Wellington	3
332	URS Wellington	2	375	Wellington	1
333	URS Wellington	3	376	Wellington	3
334	URS Wellington	7	377	Wellington	5
335	URS Wellington	4	378	Wellington	5
336	URS Wellington	3	379	Wellington	3
337	URS Wellington	4	380	Wellington	6
338	URS Wellington	7	381	Wellington	1
339	URS Wellington	3	382	Wellington	1
340	URS Wellington	1.8	383	Wellington	5
341	URS Wellington	12.3	384	Wellington	5
342	URS Wellington	7.6	385	Wellington	3
343	URS Wellington	3.8			

Summary statistics

Count	385
Mean mg/kg	4.5
Median mg/kg	3.7
Minimum mg/kg	0.4
Max mg/kg	37

Upper 95 percentile mg/kg	10.0
Upper 97.5 percentile mg/kg	11.8
Upper 98 percentile mg/kg	16.0
Upper 99 percentile mg/kg	17.4

No.	Land use	Cadmium (mg/kg)	No.	Land use	Cadmium (mg/kg)
1	Unfertilised	0.00	51	Unfertilised	0.10
2	Unfertilised	0.00	52	Unfertilised	0.10
3	Unfertilised	0.01	53	Unfertilised	0.10
4	Unfertilised	0.01	54	Unfertilised	0.10
5	Unfertilised	0.02	55	Unfertilised	0.10
6	Unfertilised	0.02	56	Unfertilised	0.10
7	Unfertilised	0.02	57	Unfertilised	0.10
8	Unfertilised	0.02	58	Unfertilised	0.10
9	Unfertilised	0.03	59	Unfertilised	0.10
10	Unfertilised	0.03	60	Unfertilised	0.10
11	Unfertilised	0.03	61	Unfertilised	0.10
12	Unfertilised	0.03	62	Unfertilised	0.10
13	Unfertilised	0.03	63	Unfertilised	0.11
14	Unfertilised	0.03	64	Unfertilised	0.11
15	Unfertilised	0.04	65	Unfertilised	0.12
16	Unfertilised	0.04	66	Unfertilised	0.12
17	Unfertilised	0.04	67	Unfertilised	0.13
18	Unfertilised	0.05	68	Unfertilised	0.13
19	Unfertilised	0.05	69	Unfertilised	0.13
20	Unfertilised	0.05	70	Unfertilised	0.13
21	Unfertilised	0.05	71	Unfertilised	0.13
22	Unfertilised	0.05	72	Unfertilised	0.13
23	Unfertilised	0.05	73	Unfertilised	0.14
24	Unfertilised	0.05	74	Unfertilised	0.14
25	Unfertilised	0.05	75	Unfertilised	0.15
26	Unfertilised	0.05	76	Unfertilised	0.15
27	Unfertilised	0.05	77	Unfertilised	0.15
28	Unfertilised	0.05	78	Unfertilised	0.15
29	Unfertilised	0.05	79	Unfertilised	0.16
30	Unfertilised	0.05	80	Unfertilised	0.17
31	Unfertilised	0.05	81	Unfertilised	0.17
32	Unfertilised	0.05	82	Unfertilised	0.17
33	Unfertilised	0.05	83	Unfertilised	0.17
34	Unfertilised	0.06	84	Unfertilised	0.17
35	Unfertilised	0.06	85	Unfertilised	0.17
36	Unfertilised	0.06	86	Unfertilised	0.17
37	Unfertilised	0.06	87	Unfertilised	0.17
38	Unfertilised	0.06	88	Unfertilised	0.17
39	Unfertilised	0.06	89	Unfertilised	0.17
40	Unfertilised	0.06	90	Unfertilised	0.20
41	Unfertilised	0.06	91	Unfertilised	0.20
42	Unfertilised	0.07	92	Unfertilised	0.20
43	Unfertilised	0.07	93	Unfertilised	0.20
44	Unfertilised	0.07	94	Unfertilised	0.20
45	Unfertilised	0.08	95	Unfertilised	0.20
46	Unfertilised	0.08	96	Unfertilised	0.20
47	Unfertilised	0.08	97	Unfertilised	0.20
48	Unfertilised	0.08	98	Unfertilised	0.21
49	Unfertilised	0.08	99	Unfertilised	0.23
50	Unfertilised	0.09	100	Unfertilised	0.23

Table A6.6: Cadmium background sites

No.	Land use	Cadmium (mg/kg)	No.	Land use	Cadmium (mg/kg)
101	Unfertilised	0.27	151	Forest	0.05
102	Unfertilised	0.27	152	Forest	0.05
103	Unfertilised	0.27	153	Forest	0.05
104	Unfertilised	0.30	154	Forest	0.05
105	Unfertilised	0.30	155	Forest	0.05
106	Unfertilised	0.30	156	Forest	0.05
107	Unfertilised	0.30	157	Forest	0.05
108	Unfertilised	0.32	158	Forest	0.06
109	Unfertilised	0.33	159	Forest	0.06
110	Unfertilised	0.33	160	Forest	0.06
111	Unfertilised	0.37	161	Forest	0.06
112	Unfertilised	0.40	162	Forest	0.07
113	Unfertilised	0.40	163	Forest	0.08
114	Unfertilised	0.43	164	Forest	0.08
115	Unfertilised	0.47	165	Forest	0.08
116	Unfertilised	0.47	166	Forest	0.08
117	Unfertilised	0.50	167	Forest	0.08
118	Unfertilised	0.60	168	Forest	0.09
119	Unfertilised	0.60	169	Forest	0.09
120	Unfertilised	0.63	170	Forest	0.10
121	Unfertilised	0.63	171	Forest	0.10
122	Unfertilised	0.67	172	Forest	0.10
123	Unfertilised	0.73	173	Forest	0.10
124	Unfertilised	0.77	174	Forest	0.10
125	Forest	0.02	175	Forest	0.10
126	Forest	0.02	176	Forest	0.10
127	Forest	0.03	177	Forest	0.10
128	Forest	0.03	178	Forest	0.10
129	Forest	0.03	179	Forest	0.10
130	Forest	0.04	180	Forest	0.10
131	Forest	0.04	181	Forest	0.10
132	Forest	0.04	182	Forest	0.10
133	Forest	0.04	183	Forest	0.10
134	Forest	0.04	184	Forest	0.10
135	Forest	0.04	185	Forest	0.12
136	Forest	0.05	186	Forest	0.13
137	Forest	0.05	187	Forest	0.14
138	Forest	0.05	188	Forest	0.15
139	Forest	0.05	189	Forest	0.17
140	Forest	0.05	190	Forest	0.18
141	Forest	0.05	191	Forest	0.20
142	Forest	0.05	192	Forest	0.20
143	Forest	0.05	193	Forest	0.30
144	Forest	0.05	194	Forest	0.49
145	Forest	0.05	195	Forest	0.60
146	Forest	0.05	196	Forest	0.60
147	Forest	0.05	197	Forest	0.65
148	Forest	0.05	198	Tussoc	0.07
149	Forest	0.05	199	Tussoc	0.08
150	Forest	0.05	200	Tussoc	0.08

No.	Land use	Cadmium (mg/kg)]	No.	Land use	Cadmium (mg/kg)
201	Tussoc	0.09	1	251	Native	0.05
202	Native	0.01		252	Native	0.05
203	Native	0.01		253	Native	0.05
204	Native	0.01		254	Native	0.05
205	Native	0.01		255	Native	0.05
206	Native	0.01		256	Native	0.05
207	Native	0.03		257	Native	0.06
208	Native	0.03		258	Native	0.06
209	Native	0.03		259	Native	0.06
210	Native	0.03		260	Native	0.06
211	Native	0.03		261	Native	0.06
212	Native	0.03		262	Native	0.06
213	Native	0.04		263	Native	0.06
214	Native	0.04		264	Native	0.07
215	Native	0.04		265	Native	0.08
216	Native	0.04		266	Native	0.08
217	Native	0.04		267	Native	0.10
218	Native	0.05		268	Native	0.10
219	Native	0.05		269	Native	0.10
220	Native	0.05		270	Native	0.10
221	Native	0.05		271	Native	0.10
222	Native	0.05		272	Native	0.10
223	Native	0.05		273	Native	0.10
224	Native	0.05		274	Native	0.10
225	Native	0.05		275	Native	0.10
226	Native	0.05		276	Native	0.10
227	Native	0.05		277	Native	0.10
228	Native	0.05		278	Native	0.10
229	Native	0.05		279	Native	0.10
230	Native	0.05		280	Native	0.10
231	Native	0.05		281	Native	0.10
232	Native	0.05		282	Native	0.10
233	Native	0.05		283	Native	0.10
234	Native	0.05		284	Native	0.10
235	Native	0.05		285	Native	0.10
236	Native	0.05		286	Native	0.11
237	Native	0.05		287	Native	0.11
238	Native	0.05		288	Native	0.11
239	Native	0.05		289	Native	0.12
240	Native	0.05		290	Native	0.12
241	Native	0.05		291	Native	0.13
242	Native	0.05		292	Native	0.13
243	Native	0.05		293	Native	0.13
244	Native	0.05		294	Native	0.13
245	Native	0.05		295	Native	0.14
246	Native	0.05		296	Native	0.14
247	Native	0.05		297	Native	0.15
248	Native	0.05		298	Native	0.15
249	Native	0.05		299	Native	0.17
250	INATIVE	0.05	J	300	INATIVE	0.18

901Native0.20361Park0.05302Native0.20352Park0.05304Native0.20354Park0.05305Native0.20356Park0.05306Native0.20356Park0.05307Native0.20358Park0.05308Native0.20359Park0.06309Native0.20359Park0.06301Native0.30360Park0.10312Native0.30362Park0.10313Native0.33363Park0.10314Native0.30366Park0.10315Native0.40366Park0.10316Native0.47366Park0.10317Native0.06368Park0.10318Native0.06368Park0.10319Native0.06369Park0.10320Native0.12371Park0.10321Native0.05373Park0.10322Park0.05374Park0.10323Park0.05376Park0.10324Park0.05376Park0.10325Park0.05378Park0.10326Park0.05 <td< th=""><th>No.</th><th>Land use</th><th>Cadmium (mg/kg)</th><th>No.</th><th>Land use</th><th>Cadmium (mg/kg)</th></td<>	No.	Land use	Cadmium (mg/kg)	No.	Land use	Cadmium (mg/kg)
302Native0.20352Park0.05303Native0.20353Park0.05305Native0.20356Park0.05306Native0.20356Park0.05307Native0.20356Park0.05308Native0.20358Park0.05309Native0.20359Park0.06309Native0.20359Park0.10311Native0.30361Park0.10312Native0.33362Park0.10313Native0.33362Park0.10314Native0.33366Park0.10315Native0.47366Park0.10316Native0.47368Park0.10317Native0.06369Park0.10320Native0.11370Park0.10321Native0.05373Park0.10322Park0.05374Park0.10323Park0.05375Park0.10324Park0.05376Park0.10325Park0.05378Park0.10326Park0.05378Park0.10331Park0.05378Park0.10332Park0.05384	301	Native	0.20	351	Park	0.05
333Native0.20353Park0.05344Native0.20355Park0.05306Native0.20356Park0.05307Native0.20357Park0.05308Native0.20358Park0.06309Native0.20359Park0.06309Native0.20359Park0.06310Native0.30360Park0.10311Native0.30362Park0.10313Native0.33363Park0.10314Native0.33364Park0.10315Native0.40365Park0.10316Native0.47366Park0.10317Native0.40366Park0.10318Native0.41370Park0.10321Native0.12371Park0.10322Park0.06368Park0.10323Park0.05374Park0.10324Park0.05374Park0.10325Park0.05376Park0.10326Park0.05376Park0.10327Park0.05376Park0.10328Park0.05376Park0.10339Park0.05376 </td <td>302</td> <td>Native</td> <td>0.20</td> <td>352</td> <td>Park</td> <td>0.05</td>	302	Native	0.20	352	Park	0.05
304 Native 0.20 354 Park 0.05 305 Native 0.20 355 Park 0.05 307 Native 0.20 357 Park 0.05 307 Native 0.20 358 Park 0.06 308 Native 0.20 358 Park 0.06 308 Native 0.30 360 Park 0.10 311 Native 0.30 362 Park 0.10 313 Native 0.30 362 Park 0.10 313 Native 0.30 363 Park 0.10 314 Native 0.39 364 Park 0.10 315 Native 0.47 366 Park 0.10 316 Native 0.05 367 Park 0.10 318 Native 0.05 371 Park 0.10 320 Park 0.05	303	Native	0.20	353	Park	0.05
305 Native 0.20 356 Park 0.05 306 Native 0.20 356 Park 0.05 308 Native 0.20 358 Park 0.05 308 Native 0.20 358 Park 0.06 309 Native 0.30 360 Park 0.10 311 Native 0.30 362 Park 0.10 313 Native 0.33 363 Park 0.10 313 Native 0.33 366 Park 0.10 314 Native 0.33 366 Park 0.10 315 Native 0.47 366 Park 0.10 316 Native 0.47 366 Park 0.10 317 Native 0.47 366 Park 0.10 318 Native 0.11 370 Park 0.10 321 Park 0.05	304	Native	0.20	354	Park	0.05
306 Native 0.20 356 Park 0.05 307 Native 0.20 357 Park 0.05 309 Native 0.20 359 Park 0.06 309 Native 0.20 359 Park 0.06 310 Native 0.30 360 Park 0.10 311 Native 0.30 361 Park 0.10 311 Native 0.33 363 Park 0.10 313 Native 0.33 366 Park 0.10 315 Native 0.40 365 Park 0.10 315 Native 0.47 366 Park 0.10 318 Native 0.47 366 Park 0.10 320 Native 0.12 371 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.05	305	Native	0.20	355	Park	0.05
307 Native 0.20 357 Park 0.05 308 Native 0.20 358 Park 0.06 308 Native 0.20 358 Park 0.06 310 Native 0.30 361 Park 0.10 311 Native 0.30 361 Park 0.10 313 Native 0.30 362 Park 0.10 313 Native 0.33 363 Park 0.10 314 Native 0.39 364 Park 0.10 315 Native 0.47 366 Park 0.10 318 Native 0.05 367 Park 0.10 319 Native 0.06 368 Park 0.10 320 Native 0.11 370 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.05	306	Native	0.20	356	Park	0.05
308 Native 0.20 358 Park 0.06 309 Native 0.20 359 Park 0.06 310 Native 0.30 360 Park 0.10 311 Native 0.30 361 Park 0.10 313 Native 0.33 363 Park 0.10 313 Native 0.33 363 Park 0.10 315 Native 0.39 364 Park 0.10 315 Native 0.47 366 Park 0.10 317 Native 0.06 368 Park 0.10 318 Native 0.06 368 Park 0.10 321 Native 0.11 370 Park 0.10 322 Park 0.06 373 Park 0.10 323 Park 0.05 374 Park 0.10 324 Park 0.05 <td< td=""><td>307</td><td>Native</td><td>0.20</td><td>357</td><td>Park</td><td>0.05</td></td<>	307	Native	0.20	357	Park	0.05
309 Native 0.20 359 Park 0.06 310 Native 0.30 360 Park 0.10 311 Native 0.30 361 Park 0.10 311 Native 0.30 362 Park 0.10 313 Native 0.33 363 Park 0.10 314 Native 0.40 366 Park 0.10 315 Native 0.47 366 Park 0.10 316 Native 0.06 368 Park 0.10 316 Native 0.12 371 Park 0.10 320 Native 0.12 371 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.05 375 Park 0.10 322 Park 0.05 376 Park 0.10 324 Park 0.05 <td< td=""><td>308</td><td>Native</td><td>0.20</td><td>358</td><td>Park</td><td>0.05</td></td<>	308	Native	0.20	358	Park	0.05
310 Native 0.30 360 Park 0.10 311 Native 0.30 361 Park 0.10 312 Native 0.30 362 Park 0.10 313 Native 0.33 363 Park 0.10 314 Native 0.39 364 Park 0.10 315 Native 0.40 365 Park 0.10 315 Native 0.47 366 Park 0.10 317 Native 0.05 367 Park 0.10 318 Native 0.06 368 Park 0.10 319 Native 0.11 370 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.05 374 Park 0.10 322 Park 0.05 375 Park 0.10 324 Park 0.05 <td< td=""><td>309</td><td>Native</td><td>0.20</td><td>359</td><td>Park</td><td>0.06</td></td<>	309	Native	0.20	359	Park	0.06
311 Native 0.30 361 Park 0.10 312 Native 0.30 362 Park 0.10 313 Native 0.33 364 Park 0.10 313 Native 0.33 364 Park 0.10 315 Native 0.40 365 Park 0.10 315 Native 0.47 366 Park 0.10 316 Native 0.06 367 Park 0.10 317 Native 0.06 368 Park 0.10 320 Native 0.12 371 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.05 373 Park 0.10 324 Park 0.05 376 Park 0.10 324 Park 0.05 377 Park 0.10 325 Park 0.05 3	310	Native	0.30	360	Park	0.10
312Native0.30362Park0.10313Native0.33363Park0.10314Native0.39364Park0.10315Native0.40366Park0.10316Native0.05367Park0.10317Native0.06366Park0.10318Native0.06367Park0.10319Native0.06368Park0.10320Native0.11370Park0.10321Native0.12371Park0.10322Park0.05373Park0.10323Park0.05373Park0.10324Park0.05374Park0.10325Park0.05376Park0.10326Park0.05376Park0.10327Park0.05376Park0.10328Park0.05376Park0.10329Park0.05380Park0.10330Park0.05380Park0.10331Park0.05381Park0.10332Park0.05382Park0.10333Park0.05382Park0.10344Park0.05384Park0.10355Park0.05386Park<	311	Native	0.30	361	Park	0.10
313 Native 0.33 363 Park 0.10 314 Native 0.39 364 Park 0.10 315 Native 0.40 366 Park 0.10 315 Native 0.47 366 Park 0.10 317 Native 0.05 367 Park 0.10 318 Native 0.06 368 Park 0.10 319 Native 0.08 369 Park 0.10 320 Native 0.11 370 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.05 374 Park 0.10 323 Park 0.05 375 Park 0.10 324 Park 0.05 376 Park 0.10 325 Park 0.05 377 Park 0.10 326 Park 0.05 380	312	Native	0.30	362	Park	0.10
314Native0.39364Park0.10315Native0.40365Park0.10316Native0.047366Park0.10317Native0.05367Park0.10318Native0.06368Park0.10319Native0.06368Park0.10320Native0.11370Park0.10321Native0.12371Park0.10322Park0.04372Park0.10323Park0.05373Park0.10324Park0.05373Park0.10325Park0.05374Park0.10326Park0.05376Park0.10327Park0.05376Park0.10328Park0.05376Park0.10329Park0.05376Park0.10330Park0.05380Park0.10331Park0.05384Park0.10332Park0.05384Park0.10333Park0.05384Park0.10334Park0.05386Park0.10335Park0.05386Park0.10336Park0.05386Park0.10337Park0.05386Park <td>313</td> <td>Native</td> <td>0.33</td> <td>363</td> <td>Park</td> <td>0.10</td>	313	Native	0.33	363	Park	0.10
315Native0.40365Park0.10316Native0.05366Park0.10317Native0.06368Park0.10318Native0.06368Park0.10319Native0.06368Park0.10320Native0.11370Park0.10321Native0.12371Park0.10322Park0.04372Park0.10323Park0.05373Park0.10324Park0.05374Park0.10325Park0.05376Park0.10326Park0.05376Park0.10327Park0.05376Park0.10328Park0.05377Park0.10329Park0.05378Park0.10330Park0.05378Park0.10331Park0.05380Park0.10332Park0.05381Park0.10333Park0.05382Park0.10334Park0.05384Park0.10335Park0.05386Park0.10336Park0.05386Park0.10337Park0.05386Park0.10338Park0.05386Park <t< td=""><td>314</td><td>Native</td><td>0.39</td><td>364</td><td>Park</td><td>0.10</td></t<>	314	Native	0.39	364	Park	0.10
316Native0.47366Park0.10317Native0.05367Park0.10318Native0.06368Park0.10319Native0.08369Park0.10321Native0.11370Park0.10322Park0.04372Park0.10323Park0.05373Park0.10324Park0.05374Park0.10325Park0.05376Park0.10326Park0.05376Park0.10327Park0.05376Park0.10328Park0.05378Park0.10329Park0.05378Park0.10330Park0.05380Park0.10331Park0.05381Park0.10332Park0.05382Park0.10333Park0.05384Park0.10344Park0.05384Park0.10355Park0.05386Park0.10366Park0.05387Park0.10374Park0.05386Park0.10375Park0.05386Park0.10376Park0.05387Park0.10376Park0.05389Park0.	315	Native	0.40	365	Park	0.10
317Native0.05367Park0.10318Native0.06368Park0.10319Native0.08369Park0.10320Native0.11370Park0.10321Native0.12371Park0.10322Park0.04372Park0.10323Park0.05373Park0.10324Park0.05374Park0.10325Park0.05376Park0.10326Park0.05376Park0.10327Park0.05376Park0.10328Park0.05376Park0.10329Park0.05378Park0.10330Park0.05380Park0.10331Park0.05381Park0.10332Park0.05382Park0.10333Park0.05383Park0.10344Park0.05384Park0.10355Park0.05386Park0.10366Park0.05387Park0.10377Park0.05386Park0.10384Park0.05386Park0.10375Park0.05386Park0.10384Park0.05389Park0.	316	Native	0.47	366	Park	0.10
318 Native 0.06 368 Park 0.10 319 Native 0.08 369 Park 0.10 320 Native 0.11 370 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.04 372 Park 0.10 323 Park 0.05 373 Park 0.10 324 Park 0.05 374 Park 0.10 325 Park 0.05 375 Park 0.10 326 Park 0.05 377 Park 0.10 327 Park 0.05 378 Park 0.10 328 Park 0.05 380 Park 0.10 330 Park 0.05 381 Park 0.10 331 Park 0.05 382 Park 0.10 332 Park 0.05 384	317	Native	0.05	367	Park	0.10
319 Native 0.08 369 Park 0.10 320 Native 0.11 370 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.04 372 Park 0.10 323 Park 0.05 373 Park 0.10 324 Park 0.05 374 Park 0.10 325 Park 0.05 375 Park 0.10 326 Park 0.05 376 Park 0.10 327 Park 0.05 377 Park 0.10 328 Park 0.05 378 Park 0.10 329 Park 0.05 380 Park 0.10 330 Park 0.05 381 Park 0.10 333 Park 0.05 382 Park 0.10 333 Park 0.05 385	318	Native	0.06	368	Park	0.10
320 Native 0.11 370 Park 0.10 321 Native 0.12 371 Park 0.10 322 Park 0.04 372 Park 0.10 323 Park 0.05 373 Park 0.10 324 Park 0.05 374 Park 0.10 324 Park 0.05 375 Park 0.10 325 Park 0.05 376 Park 0.10 326 Park 0.05 377 Park 0.10 327 Park 0.05 377 Park 0.10 328 Park 0.05 379 Park 0.10 329 Park 0.05 380 Park 0.10 330 Park 0.05 381 Park 0.10 333 Park 0.05 384 Park 0.10 334 Park 0.05 386	319	Native	0.08	369	Park	0.10
321 Native 0.12 371 Park 0.10 322 Park 0.04 372 Park 0.10 323 Park 0.05 373 Park 0.10 324 Park 0.05 374 Park 0.10 325 Park 0.05 376 Park 0.10 326 Park 0.05 376 Park 0.10 326 Park 0.05 376 Park 0.10 327 Park 0.05 377 Park 0.10 328 Park 0.05 379 Park 0.10 330 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 332 Park 0.05 382 Park 0.10 333 Park 0.05 385 Park 0.10 334 Park 0.05 386 <t< td=""><td>320</td><td>Native</td><td>0.11</td><td>370</td><td>Park</td><td>0.10</td></t<>	320	Native	0.11	370	Park	0.10
322 Park 0.04 372 Park 0.10 323 Park 0.05 373 Park 0.10 324 Park 0.05 374 Park 0.10 325 Park 0.05 375 Park 0.10 326 Park 0.05 376 Park 0.10 326 Park 0.05 377 Park 0.10 328 Park 0.05 377 Park 0.10 329 Park 0.05 378 Park 0.10 330 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 333 Park 0.05 382 Park 0.10 334 Park 0.05 385 Park 0.10 335 Park 0.05 386 Park 0.10 336 Park 0.05 386	321	Native	0.12	371	Park	0.10
323 Park 0.05 373 Park 0.10 324 Park 0.05 374 Park 0.10 325 Park 0.05 375 Park 0.10 326 Park 0.05 376 Park 0.10 327 Park 0.05 376 Park 0.10 328 Park 0.05 377 Park 0.10 329 Park 0.05 378 Park 0.10 330 Park 0.05 379 Park 0.10 331 Park 0.05 380 Park 0.10 333 Park 0.05 381 Park 0.10 334 Park 0.05 382 Park 0.10 335 Park 0.05 383 Park 0.10 336 Park 0.05 385 Park 0.10 337 Park 0.05 386 Park 0.10 338 Park 0.05 389 Park <td< td=""><td>322</td><td>Park</td><td>0.04</td><td>372</td><td>Park</td><td>0.10</td></td<>	322	Park	0.04	372	Park	0.10
324 Park 0.05 374 Park 0.10 325 Park 0.05 375 Park 0.10 326 Park 0.05 376 Park 0.10 327 Park 0.05 377 Park 0.10 328 Park 0.05 378 Park 0.10 329 Park 0.05 379 Park 0.10 330 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 332 Park 0.05 381 Park 0.10 333 Park 0.05 382 Park 0.10 334 Park 0.05 385 Park 0.10 335 Park 0.05 386 Park 0.10 335 Park 0.05 386 Park 0.10 336 Park 0.05 390	323	Park	0.05	373	Park	0.10
325 Park 0.05 375 Park 0.10 326 Park 0.05 376 Park 0.10 327 Park 0.05 377 Park 0.10 328 Park 0.05 377 Park 0.10 328 Park 0.05 378 Park 0.10 329 Park 0.05 379 Park 0.10 330 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 332 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 385 Park 0.10 337 Park 0.05 386 Park 0.10 338 Park 0.05 389	324	Park	0.05	374	Park	0.10
326 Park 0.05 376 Park 0.10 327 Park 0.05 377 Park 0.10 328 Park 0.05 378 Park 0.10 329 Park 0.05 379 Park 0.10 330 Park 0.05 379 Park 0.10 331 Park 0.05 380 Park 0.10 332 Park 0.05 381 Park 0.10 333 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 386 Park 0.10 336 Park 0.05 388 Park 0.10 337 Park 0.05 389 Park 0.10 338 Park 0.05 390	325	Park	0.05	375	Park	0.10
327 Park 0.05 377 Park 0.10 328 Park 0.05 378 Park 0.10 329 Park 0.05 379 Park 0.10 330 Park 0.05 379 Park 0.10 331 Park 0.05 380 Park 0.10 332 Park 0.05 381 Park 0.10 333 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 385 Park 0.10 336 Park 0.05 386 Park 0.10 337 Park 0.05 388 Park 0.10 338 Park 0.05 389 Park 0.10 340 Park 0.05 389 Park 0.10 341 Park 0.05 390 Park <td< td=""><td>326</td><td>Park</td><td>0.05</td><td>376</td><td>Park</td><td>0.10</td></td<>	326	Park	0.05	376	Park	0.10
328 Park 0.05 378 Park 0.10 329 Park 0.05 379 Park 0.10 330 Park 0.05 380 Park 0.10 331 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 332 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 386 Park 0.10 336 Park 0.05 386 Park 0.10 337 Park 0.05 389 Park 0.10 341 Park 0.05 390 Park 0.10 341 Park 0.05 393	327	Park	0.05	377	Park	0.10
329 Park 0.05 379 Park 0.10 330 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 332 Park 0.05 381 Park 0.10 332 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 385 Park 0.10 336 Park 0.05 386 Park 0.10 337 Park 0.05 388 Park 0.10 338 Park 0.05 389 Park 0.10 341 Park 0.05 390 Park 0.11 342 Park 0.05 393	328	Park	0.05	378	Park	0.10
330 Park 0.05 380 Park 0.10 331 Park 0.05 381 Park 0.10 332 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 383 Park 0.10 335 Park 0.05 384 Park 0.10 336 Park 0.05 385 Park 0.10 337 Park 0.05 386 Park 0.10 338 Park 0.05 387 Park 0.10 339 Park 0.05 388 Park 0.10 341 Park 0.05 389 Park 0.10 342 Park 0.05 391 Park 0.11 343 Park 0.05 392 Park 0.12 344 Park 0.05 393 Park 0.12 344 Park 0.05 395 Park <td< td=""><td>329</td><td>Park</td><td>0.05</td><td>379</td><td>Park</td><td>0.10</td></td<>	329	Park	0.05	379	Park	0.10
331 Park 0.05 381 Park 0.10 332 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 384 Park 0.10 336 Park 0.05 385 Park 0.10 337 Park 0.05 386 Park 0.10 338 Park 0.05 387 Park 0.10 339 Park 0.05 389 Park 0.10 341 Park 0.05 389 Park 0.10 342 Park 0.05 390 Park 0.11 342 Park 0.05 393 Park 0.12 344 Park 0.05 394 Park 0.12 344 Park 0.05 395 Park <td< td=""><td>330</td><td>Park</td><td>0.05</td><td>380</td><td>Park</td><td>0.10</td></td<>	330	Park	0.05	380	Park	0.10
332 Park 0.05 382 Park 0.10 333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 384 Park 0.10 336 Park 0.05 385 Park 0.10 336 Park 0.05 386 Park 0.10 337 Park 0.05 386 Park 0.10 338 Park 0.05 387 Park 0.10 339 Park 0.05 388 Park 0.10 341 Park 0.05 390 Park 0.10 341 Park 0.05 391 Park 0.11 343 Park 0.05 392 Park 0.12 344 Park 0.05 393 Park 0.12 345 Park 0.05 395 Park 0.12 346 Park 0.05 396 Park <td< td=""><td>331</td><td>Park</td><td>0.05</td><td>381</td><td>Park</td><td>0.10</td></td<>	331	Park	0.05	381	Park	0.10
333 Park 0.05 383 Park 0.10 334 Park 0.05 384 Park 0.10 335 Park 0.05 385 Park 0.10 336 Park 0.05 385 Park 0.10 336 Park 0.05 386 Park 0.10 337 Park 0.05 386 Park 0.10 338 Park 0.05 387 Park 0.10 339 Park 0.05 388 Park 0.10 340 Park 0.05 389 Park 0.10 341 Park 0.05 390 Park 0.11 342 Park 0.05 392 Park 0.11 343 Park 0.05 393 Park 0.12 344 Park 0.05 395 Park 0.12 345 Park 0.05 396 Park 0.12 346 Park 0.05 397 Park <td< td=""><td>332</td><td>Park</td><td>0.05</td><td>382</td><td>Park</td><td>0.10</td></td<>	332	Park	0.05	382	Park	0.10
334 Park 0.05 384 Park 0.10 335 Park 0.05 385 Park 0.10 336 Park 0.05 386 Park 0.10 337 Park 0.05 387 Park 0.10 338 Park 0.05 387 Park 0.10 338 Park 0.05 387 Park 0.10 339 Park 0.05 388 Park 0.10 340 Park 0.05 389 Park 0.10 341 Park 0.05 390 Park 0.10 342 Park 0.05 391 Park 0.11 343 Park 0.05 392 Park 0.12 344 Park 0.05 393 Park 0.12 345 Park 0.05 395 Park 0.12 345 Park 0.05 396 Park 0.12 346 Park 0.05 397 Park <td< td=""><td>333</td><td>Park</td><td>0.05</td><td>383</td><td>Park</td><td>0.10</td></td<>	333	Park	0.05	383	Park	0.10
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	350	Park	0.05	400	Park	0.15

No.	Land use	Cadmium (mg/kg)		No.	Land use	Cadmium (mg/kg)
401	Park	0.15		444	Park	0.26
402	Park	0.15		445	Park	0.27
403	Park	0.15		446	Park	0.27
404	Park	0.15		447	Park	0.27
405	Park	0.15		448	Park	0.27
406	Park	0.15		449	Park	0.27
407	Park	0.16		450	Park	0.28
408	Park	0.16		451	Park	0.28
409	Park	0.17		452	Park	0.29
410	Park	0.17		453	Park	0.29
411	Park	0.17		454	Park	0.30
412	Park	0.18		455	Park	0.30
413	Park	0.18		456	Park	0.30
414	Park	0.18		457	Park	0.30
415	Park	0.18		458	Park	0.31
416	Park	0.19		459	Park	0.31
417	Park	0.19		460	Park	0.31
418	Park	0.19		461	Park	0.32
419	Park	0.19		462	Park	0.32
420	Park	0.19		463	Park	0.32
421	Park	0.20		464	Park	0.33
422	Park	0.20		465	Park	0.34
423	Park	0.20		466	Park	0.37
424	Park	0.20		467	Park	0.39
425	Park	0.20		468	Park	0.39
426	Park	0.20		469	Park	0.40
427	Park	0.20		470	Park	0.41
428	Park	0.20		4/1	Park	0.42
429	Park	0.21		472	Park	0.43
430	Park	0.21		473	Park	0.44
431	Park	0.21		474	Park	0.45
432	Park	0.22		475	Park	0.46
433	Park	0.22		476	Park	0.47
434	Park	0.23		4//	Park	0.48
435	Park	0.23		470	Park	0.40
430	Park	0.23		479	Park	0.50
437	Park	0.23		400	Park	0.55
439	Park	0.20		482	Park	0.59
440	Park	0.24		483	Park	0.60
441	Park	0.25		484	Park	0.63
442	Park	0.25		485	Park	0.65
443	Park	0.25		486	Park	0.77
		0.20	l	100		0

Summary statistics

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Count	486
Mean mg/kg	0.14787
Median mg/kg	0.1
Minimum mg/kg	0.003
Max mg/kg	0.77

Upper 95 percentile mg/kg	0.466
Upper 97.5 percentile mg/kg	0.6
Upper 98 percentile mg/kg	0.6006
Upper 99 percentile mg/kg	0.65

Abbreviations and Glossary

AD _{adj}	Age-adjusted dermal absorption factor
ADE	Average daily human exposure
ADI	Acceptable daily intake
AF	Contaminant-specific dermal absorption factor
AH	Soil adherence factor
AR	Skin area
AT	Averaging time
ATSDR	Agency for Toxic Substances and Disease Control (US)
BAF	Bioaccumulation factor
BaP	Benzo(a)pyrene
BaP _{eq}	BaP equivalence concentration
BCF	Bioconcentration factor
BI	Background intake
BTEX	Benzene, ethyl benzene, toluene and xylene
BW	Body weight
CCME	Canadian Council of Ministers of the Environment
CEAA	Canadian Environmental Assessment Act
CEPA	Canadian Environmental Protection Act
CERCLA	Comprehensive Environmental Response Compensation and Liability Act (US)
CLEA	Contaminated Land Exposure Assessment soil guideline derivation model (UK)
CSOIL	Contaminated soil exposure model (the Netherlands)
C _P	Concentration in produce
Cs	Concentration in soil
DDD	Dichloro diphenyl-dichloroethane
DDE	Dichloro diphenyl-dichloroethylene
DDT	Dichloro diphenyl-trichloroethane
Defra	Department for the Environment, Food and Rural Affairs (UK)
DER	Default exposure ratio
DW	Dry weight
EA	The Environment Agency (UK)
ECO-SSL	Ecological soil screening limit (USA)
ED	Exposure duration (years)
EDI	Estimated daily intake
EF	Exposure frequency (days/year)
EPA	Environmental Protection Agency/Authority (US)
ERMA	Environmental Risk Management Authority (NZ)
EFSA	European Food Safety Authority

FAO	Food and Agriculture Organization
FW	Fresh weight
GD	Guidance dose for cancer toxic effects
GRI	Gas Research Institute
GWG	Gasworks guidelines
HHEM	Human health evaluation manual
HIL	Health-based investigation levels (Australia)
HpCDD	Heptachloro dibenzo-p-dioxin
HQ	Hazard quotient
HRS	Hazard ranking system
IH	Inhalation rate
$\mathrm{IH}_{\mathrm{adj}}$	Age-adjusted inhalation rate
IP	Produce ingestion rate
IP_{adj}	Age-adjusted produce ingestion rate
IR	Soil ingestion rate
IR _{adj}	Age-adjusted soil ingestion rate
IV	Intervention value (The Netherlands)
JECFA	Joint Experts Committee on Food Additives
K _h	Henry's Law coefficient
K _{ow}	Octanol-water partition coefficient
LEED	Linked employer-employee data
ln	Natural logarithm (logarithm to the base e)
LOEC	Lowest observed effect concentration
log	Logarithm to the base 10
MfE	Ministry for the Environment
MoH	Ministry of Health
MPR _{eco}	Maximum permissible risk level for ecosystems (The Netherlands)
MRL	Maximum residue level
NCP	National contingency plan
NEPC	National Environmental Protection Council (Australia)
NEPM	National environmental protection (assessment of site contamination) measure (Australia)
NES	National environmental standard
NOEC	No observed effect concentration
NPL	National priorities list (US)
OCDD	Octachloro dibenzo-p-dioxin
OIG	'Oil Industry Guidelines' (also known as Petroleum Industry Guidelines)
PAH	Polycyclic aromatic hydrocarbons
PCB	Polychlorinated biphenyl
PCDD	Polychlorinated dibenzo-p-dioxin
PCDF	Polychlorinated dibenzofuran

PCP	Pentachlorophenol
PeCDD	Pentachloro dibenzo-p-dioxin
PEF	Particle emission factor or Potency equivalency factor
Pg	Proportion of home-grown produce
pН	Measure of acidity and alkalinity
p_i	Proportion of total vegetable consumption of vegetable type i
PRG	Preliminary remediation goals
PTWI	Provisional tolerable weekly intake
RAGS	Risk assessment guidance for superfund (US)
RAIS	Risk assessment information system (US)
RCRA	Resource Conservation and Recovery Act (US)
RHS	Reference health standard
RME	Reasonable maximum exposure (US)
SCS _(health)	Soil contaminant standard protective of human health having regulatory status under the NES
SDG	Sheep-dip Guide
SGV	Soil guideline value
SGV _(health)	Soil guideline value protective of human health
SSGV	Site-specific soil guideline value derived according to the NES methodology
SL	Soil loading factor
SQGHH	Human health soil quality guideline (Canada)
SSL	Soil screening level (US)
TCDD	Tetrachlorodibenzo-p-dioxin or 2,3,7,8-tetrachlorodibenzo-p-dioxin
TDI	Tolerable daily intake
TDSI	Tolerable daily soil intake
TEF	Toxic equivalence factor
TEQ	Toxic equivalency (for dioxins)
TTG	Timber Treatment Guidelines
TV	Target value
UK	United Kingdom
US EPA	United States Environmental Protection Agency
UST	Underground storage tanks (US)
WHO	World Health Organization
∑DDT	Total DDT (sum of DDT, DDE, DDD)

References

Alloway BJ. 1995. Cadmium. In BJ Alloway (ed) *Heavy Metals in Soils*. Chapter 6, pp. 122–147, 2nd edition. Blackie.

Alloway BJ, Thornton I, Smart GA, Sherlock JC, Quinn MJ. 1988. Metal availability. *Science of the Total Environment* 75: 41–69.

Anonymous. 1987. Environmental Health Criteria 71 Pentachlorophenol. International Program on Chemical Safety. Geneva, World Health Organization.

ANZECC. 1992. Australian and New Zealand Guidelines for the Assessment and Management of Contaminated Sites. Australian and New Zealand Environment and Conservation Council.

ATSDR. 1992. *Public Health Assessment Guidance Manual*. Agency for Toxic Substances and Disease Registry, US Department of Health and Human Services, Atlanta, Georgia.

Auckland Regional Council 2001: *Background concentrations of inorganic elements from the Auckland region*, Auckland Regional Council Technical Publication No. 153, October 2001.

Australian exposure factor guidance handbook. 2011. In press.

Baes CF, Sharp RD, Sjoreen AL, Shor RW. 1984. A Review and Analysis of Parameters for Assessing Transport of Environmentally Released Radionuclides Through Agriculture. ORNL-5786, Health and Safety Research Division, Oak Ridge National Laboratory, Oak Ridge Tennessee.

Baars AJ, Theelan RMC, Janssen PJCM, Hesse JM, van Apeldoorn ME, Meijerink MCM, Verdam L, Zeilmaker MJ. 2001. *Re-evaluation of Human Toxicological Maximum Permissible Risk Levels*. RIVM report 711701025, National Institute for Public Health and the Environment (RIVM), Bilthoven, The Netherlands.

Beaulieu M. 2007. International Overview of the Practices on Contaminated Sites in Canada. Presentation at the 7th Meeting of the International Committee On Contaminated Land, Stockholm, 10–11 September 2007.

Bechtel-Jacobs. 1998. *Empirical Models for the Uptake of Inorganic Chemicals from Soil by Plants*. Report BJC/OR-133, Bechtel Jacobs Company LLC, Oak Ridge, Tennessee.

Bellin CA, O'Connor GA. 1990. Plant uptake of pentachlorophenol from sludge-amended soils. *Journal of Environmental Quality* 19: 598–602.

Binder S, Sokal D, Maughan D. 1986. Estimating soil ingestion: the use of tracer elements in estimating the amount of soil ingested by young children. *Archives of Environmental Health* 41(6): 341–345.

Blevins DG, Lukaszewski KM. 1998. Boron in plant structure and function. *Annual Reviews of Plant Physiology and Plant Molecular Biology* 49: 481–500.

Brand E, Otte PF, Lijzen, JPA. 2007. CSOIL 2000; an exposure model for human risk assessment of soil contamination. A model description. RIVM report 711701054/2007. National Institute for Public Health and the Environment (RIVM), Bilthoven, The Netherlands.

Briggs GG, Bromilow RH, Evans AA. 1982. Relationship between lipophilicity and root uptake and translocation of non-ionised chemicals in barley. *Pesticide Science* 13: 495–504.

Briggs GG, Bromilow RH, Evans, AA, Williams, M. 1983. Relationships between lipophilicity and the distribution of non-ionised chemicals in barley shoots following uptake by the roots. *Pesticide Science* 14: 492–500.

Brindson S. 2004. *Simulated Typical Diets for the 2003/004 New Zealand Total Diet Survey*. Report prepared for the New Zealand Food Safety Authority.

Calabrese EJ. 2003. Letter from Edward Calabrese to K Holtzclaw, the General Electric Company, re: Soil Ingestion Rates. 23 July 2003.

Calabrese EJ, Pastides H, Barnes R, Edwards C, Kostecki PT, Stanek EJ, Veneman P, Gilbert CE. 1989. How much soil do young children ingest: an epidemiologic study. In: *Petroleum Contaminated Soils*. Lewis Publishers, Chelsea, MI. 16 pp. 363–397.

Calabrese EJ, Stanek EJ. 1991. A guide to interpreting soil ingestion studies: II. Quantitative and qualitative evidence of soil ingestion. *Regulatory Toxicology and Pharmacology* 13: 278–292.

Calabrese EJ, Stanek EJ, Pekow P, Barnes RM. 1997. Soil ingestion estimates for children residing on a Superfund site. *Ecotoxicological and Environmental Safety* 36: 258–268.

Carlon C (ed). 2007. Derivation Methods of Soil Screening Values in Europe. A Review and Evaluation of National Procedures Towards Harmonization. European Commission, Joint Research Centre, Ispra, Italy EUR 22805-EN.

Casterline JL, Barnett NM, Ku Y. 1985. Uptake, translocation and transformation of pentachlorophenol in soybean and spinach plants. *Environmental Research* 37: 101–118.

Cavanagh JE. 2003. Critique of National and International Methodologies Used to Derive Numeric Values Representing Soil Contaminant Concentrations Protective of Human Health. Contract Report: LC0203/146 for the Ministry for the Environment, Landcare Research, Lincoln.

Cavanagh JE. 2004a. *Proposed Exposure Scenarios, Pathways and Parameters for Use in Deriving Soil Guideline Values.* Unpublished report, Ministry for the Environment, Wellington.

Cavanagh JE. 2004b. *Review of Soil Acceptance Criteria for Copper and DDT*. Contract Report LC0405/043 for the Auckland Regional Council, Landcare Research, Lincoln.

Cavanagh JE. 2004c. *Review of Soil Acceptance Criteria for Dieldrin*. Contract Report LC0405/057 for the Auckland Regional Council, Landcare Research, Lincoln.

Cavanagh JE. 2004d. *Review of Soil Acceptance Criteria for Lead*. Contract Report LC0405/050 for the Auckland Regional Council, Landcare Research, Lincoln.

Cavanagh JE. 2005a. *Review of Soil Acceptance Criteria for Arsenic*. Landcare Research Contract Report LC0405/086 for the Auckland Regional Council.

Cavanagh JE. 2005b. *Basis for Deriving Soil Guideline Values Protective of Human Health for the Proposed National Environmental Standards for Contaminated Land*. Contract Report: LC04050/120 for the Ministry for the Environment, Landcare Research, Lincoln.

Cavanagh JE, O'Halloran K. 2003. International Legislative Frameworks for Managing Contaminated Land – Derivation and Application of Numeric Values. Contract Report LC0203/110 for the Ministry for the Environment, Landcare Research, Lincoln.

Cavanagh JE, Proffitt GT. 2005. Soil Acceptance Criteria for Sandilands Residential Area. Contract Report: LC0405/074 for Christchurch City Council, Landcare Research, Lincoln.

CCME. 1991. The National Contaminated Sites Remediation Program. Canadian Council of Ministers of the Environment, Winnipeg, Manitoba.

CCME. 1996. Protocol for the Derivation of Environmental and Human Health Soil Quality Guidelines. Canadian Council of Ministers of the Environment, Winnipeg, Manitoba.

CCME. 1997. Canadian Soil Quality Guidelines for Pentachlorophenol: Environmental and Human Health. Canadian Council of Ministers of the Environment, Winnipeg, Canada.

CCME. 2006. Protocol for the Derivation of Environmental and Human Health Soil Quality Guidelines. Canadian Council of Ministers of the Environment, Winnipeg, Manitoba.

Clausing P, Brunekreef B, Van Wijnen JH. 1987. A method for estimating soil ingestion by children. *International Archives of Occupational and Environmental Health* 59(1): 73–82.

Collins C, Fryer M, Grosso A. 2006. Plant uptake of non-ionic organic chemicals. *Environmental Science and Technology* 40: 45–52.

Crommentuijn GH, Van den Plassche EJ, Canton JH. 1994. *Guidance Document on the Derivation of Ecotoxicological Criteria for Serious Soil Contamination in View of the Intervention Value for Soil Clean-up*. RIVM Report 955001003, National Institute of Public Health and the Environment, Bilthoven, The Netherlands.

Crosby DG. 1981. Environmental chemistry of pentachlorophenol. *Pure and Applied Chemistry* 53: 1051–1080.

Davis S, Mirick D. 2006. Soil ingestion in children and adults in the same family. *Journal of Exposure Analysis and Environmental Epidemiology* 16: 63–75.

Davis S, Waller P, Buschbon R, Ballou J, White P. 1990. Quantitative estimates of soil ingestion in normal children between the ages of 2 and 7 years: population based estimates using aluminium, silicon, and titanium as soil tracer elements. *Archives of Environmental Health* 45: 112–122.

Davis GB, Trefy MG, Patterson BM. 2004. *Petroleum and Solvent Vapours: Quantifying Their Behaviour, Assessment and Exposure*. A report to the Western Australian Department of Environment, CSIRO Land and Water Report.

Davis GB, Trefry MG, Patterson BM. 2008. *Petroleum Vapour Model Comparison: Interim Report for CRC CARE*. Technical Report No. 1, CRC for Contamination Assessment and Remediation of the Environment Pty Ltd, Adelaide.

Defra. 2005. *CLAN 2/05 – Soil Guideline Values and the Determination of Land as Contaminated Land under Part II*. Contaminated Land Note 2/05, Contaminated Land Branch, Department for Environment, Food and Rural Affairs. Retrieved from http://www.defra.gov.uk/environment/quality/land/contaminated/ technical-guidance.htm (May 2008).

Defra. 2006a. Assessing Risks From Land Contamination – A Proportionate Approach, Soil Guideline Values: The Way Forward. Report CLAN 6/06, Department for Environment, Food and Rural Affairs, London.

Defra. 2006b. *Circular 01/2006 Environmental Protection Act 1990: Part 2A Contaminated Land*. Department for Environment, Food and Rural Affairs, London.

Defra. 2008a. Improvements to Contaminated Land Guidance: Outcome of the "Way Forward" Exercise on Soil Guideline Values. Department for Environment, Food and Rural Affairs, London.

Defra. 2008b. *Guidance on the Legal Definition of Contaminated Land*. Department for Environment, Food and Rural Affairs, London.

Defra and EA. 2002a. *The Contaminated Land Exposure Assessment (CLEA) Model: Technical basis and algorithms.* R&D Publication CLR10, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

Defra and EA. 2002b. Assessment of Risks to Human Health from Land Contamination: An Overview of the Development of Soil Guideline Values and Related Research. R&D Publication CLR7, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

Defra and EA. 2002c. *Soil Guideline Values for Cadmium Contamination*. R&D Publication SGV3, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

Defra and EA. 2002d. *Soil Guideline Values for Chromium Contamination*. R&D Publication SGV4, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

Defra and EA. 2002e. *Potential Contaminants for the Assessment of Land*. R&D Publication CLR8, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

Defra and EA. 2002f. Contaminants in Soil: Collation of Toxicological Data and Intake Values for Humans. R&D Publication CLR9, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

Defra and EA. 2004. *Model Procedures for the Management of Land Contamination*, R&D Publication CLR11, Department for Environment, Food and Rural Affairs and the Environment Agency, Bristol, UK.

DUAP and EPA. 1998. *Managing Land Contamination Planning Guidelines; State Environmental Planning Policy 55 (SEPP 55) – Remediation of Land*, New South Wales Department of Urban Affairs and Planning and New South Wales Environment Protection Authority.

EA. 2004a. Update on Estimating Vapour Intrusion into Buildings. CLEA Briefing Note 2: Version 1.1 (July 2004), Environment Agency. Retrieved from http://www.environment-agency.gov.uk/subjects/landquality (May 2008).

EA. 2004b. Update of Supporting Values and Assumptions Describing UK Building Stock. CLEA briefing Note 3: Version 1.1 (July 2004), Environment Agency. Retrieved from http://www.environment-agency.gov.uk/subjects/landquality (May 2008).

EA. 2005a. *Update on the Dermal Exposure Pathway*. CLEA Briefing Note 1: Version 1.1 (March 2005), Environment Agency. Retrieved from http://www.environment-agency.gov.uk/subjects/landquality (May 2008).

EA. 2005b. An Update on Deriving Soil Guideline Values Based on Combined Intake from Individual Routes of Exposure. CLEA Briefing Note 4: Version 1.0 (November 2005), Environment Agency. Retrieved from http://www.environment-agency.gov.uk/subjects/landquality (May 2008).

EA. 2006. *Evaluation of Models for Predicting Plant Uptake of Chemicals from Soil*. Science Report SC050021/SR, Environment Agency, Bristol, UK.

EA. 2008a. Updated Technical Background to the CLEA Model. Science Report SC050021/SR3, Environment Agency, Bristol, UK. Retrieved from http://www.environment-agency.gov.uk/subjects/landquality/113813/ (October 2008).

EA. 2008b. Human Health Toxicological Assessment of Contaminants in Soil. Science Report SC050021/SR2, Environment Agency, Bristol, UK. Retrieved from http://www.environment-agency.gov.uk/subjects/landquality/113813/ (October 2008).

EA. 2009a. Supplementary Information for the Derivation of SGV for Arsenic. Science Report SC050021, Environment Agency, Bristol, UK.

EA. 2009b. Supplementary Information for the Derivation of SGV for Cadmium. Science Report SC050021/Technical review cadmium, Environment Agency, Bristol, UK.

EA. 2009c. *Supplementary Information for the Derivation of SGV for Mercury*. Science report: SC050021 Environment Agency, Bristol, UK

ECETOC. 1990. *Hazard Assessment of Chemical Contaminants in Soil*. Technical Report No. 40. European Chemical Industry Ecology and Toxicology Centre, Brussels.

Effroysom RA, Sample BE, Suter GW. 2004. Bioaccumulation of inorganic chemicals from soil by plants: spiked soils vs field contamination or background. *Human and Ecological Risk Assessment* 10: 1117–1127.

El Saadi O, Langley A (eds). 1991. *The Health Risk Assessment and Management of Contaminated Sites*. Workshop proceedings of the National Workshop on the Health Risk, Assessment and Management of Contaminated Sites.

Evans D, Hers I, Wolters RM, Boddington RTB, Hall DH. 2002. *Vapour Transfer of Soil Contaminants*. R&D Technical Report P5-018/TR, prepared by Golder Associates, Environment Agency, Bristol, UK.

FAO. 2000. Assessing Soil Contamination: A Reference Manual. FAO Pesticide Disposal Series 8, Food and Agriculture Organization of the United Nations, Rome.

FAO/WHO. 2006. *Summary and Conclusions*. 64th meeting of the Joint FAO/WHO Expert Committee on Food Additives. World Health Organization, Geneva.

Fowler R. 2007. *Site Contamination Law and Policy in Europe, North America and Australia – Trends and Challenges.* Proceedings 8th Meeting of the International Committee on Contaminated Land, Stockholm, 10–11 September 2007. Retrieved from http://www.iccl.ch/meeting_stockholm.html (May 2008).

Frame GM, Cochran JW. Boewadt SS. 1996. Complete PCB congener distributions for 17 Aroclor mixtures determined by 3 HRGC systems optimised for comprehensive, quantitative, congener-specific analysis. *Journal of High Resolution Chromatography* 19: 657–668.

Gaw SK, Kim ND, Northcott GL, Wilkins AL, Robinson G. 2008. Uptake of DDT, arsenic, cadmium, copper and lead by lettuce and radish grown in contaminated horticultural soils. *Journal of Agricultural and Food Chemistry* 56: 6584–6593.

GRI. 1988. *The Management of Manufactured Gas Plant Sites*. Vol III, Risk Assessment. Gas Research Institute, Chicago.

Harnley ME, Petreas MX, Flattery J, Goldman LR. 2000. Polychlorinated dibenzo-*p*-dioxin and polychlorinated dibenzofuran contamination in soil and home-produced eggs near pentachlorophenol sources. *Environmental Science and Technology* 34: 1143–1149.

Haque A, Gruttke H, Kratz W, Kielhorn U, Weigmann G, Meyer G, Bornkamm R, Schupan I, Ebing W. 1988. Environmental fate and distribution of sodium [14C] pentachlorophenate in a section of urban wasteland ecosystem. *The Science of the Total Environment* 68: 127–139.

Hawley JK. 1985. Assessment of health risk from exposure to contaminated soil. *Risk Analysis* 5: 289–302.

He Y, Xu J, Tang C, Wu Y. 2005. Facilitation of pentachlorophenol degradation in the rhizosphere of ryegrass (*Lolium perenne* L). *Soil Biology and Biochemistry* 37: 2017–2024.

Health Canada. 2004. Federal Contaminated Site Risk Assessment in Canada – Part I: Guidance on Human Health Preliminary Quantitative Risk Assessment (PQRA). Health Canada, Environmental Health Assessment Services, Safe Environments Program, Ottawa.

Hill RB, Sparling GP. 2009. *Soil quality monitoring*. In: Land Monitoring Forum. Land and Soil Monitoring: A guide for SoE and regional council reporting. Hamilton: Land Monitoring Forum: 27–88.

Hofshi R, Boreham D, Arpia ML. 2000. 'Lamb Hass' Maturity and Fruit Quality Study. Presented at the 2000 Avocado Research Symposium. California Avocado Commission.

Hulster A, Marschner H. 1993. Transfer of PCDD/PCDF from contaminated soils to food and fodder crops. *Chemosphere* 27: 439–446.

Hulster A, Muller J, Marschner H. 1994. Soil-plant transfer of polychlorinated dibenzo-*p*-dioxins and dibenzofurans to vegetables of the cucumber family (Cucurbitaceae). *Environmental Science and Technology* 28: 1110–1115.

Inui H, Wakai T, Gion K, Kim Y-S, Eun H. 2008. Differential uptake for dioxin-like compounds by zucchini subspecies. *Chemosphere* 73: 1602–1607.

Johnson PC, Ettinger RA. 1991. Heuristic model for predicting the intrusion rate of contaminant vapors in buildings. *Environmental Science and Technology* 25: 1445–1452.

Jones KC, Duarte-Davidson R. 1997. Transfers of airborne PCDD/Fs to bulk deposition collectors and herbage. *Environmental Science and Technology* 31: 2937–2943.

Jones KC, Johnston AE. 1991. Significance of atmospheric inputs of lead to grassland at one site in the United Kingdom since 1860. *Environmental Science and Technology* 25: 1174–1178.

Jury WA, Spencer WF, Farmer WJ. 1983. Behaviour assessment model for trace organics in soil: 1. Model Description. *Journal of Environmental Quality* 12(4).

Jury WA, Spencer WF, Farmer WJ. 1984. Behaviour assessment model for trace organics in soil: 11. Chemical classification and parameter sensitivity. *Journal of Environmental Quality* 13(4).

Jury WA, Russo D, Streile G, Abd HE. 1990. Evaluation of volatilisation by organic chemicals residing below the soil surface. *Water Resources Research* 26(1): 13–20.

Kabata-Pendias A, Pendias H. 2000. *Trace Elements in Soils and Plants*, 3rd ed. CRC Press, Boca Raton, Florida USA.

Kim ND, Taylor MD. 2009. *Trace element monitoring*. In: Land Monitoring Forum. Land and Soil Monitoring: A guide for SoE and regional council reporting. Hamilton: Land Monitoring Forum: 117–178.

Kim ND. 2005. Concentrations of various elements in Waikato surface soils and suggested default cleanfill thresholds. Environment Waikato.

Kipopoulou AM, Manoli E, Samara C. 1999. Bioconcentration of polycyclic aromatic hydrocarbons in vegetables grown in an industrial area. *Environmental Pollution* 106: 369–380.

Kreule P, Van den Berg R, Waitz MFW, Swartjes FA. 1995. *Calculation of human toxicological serious soil contamination concentrations and proposals for intervention values for clean-up of soil and groundwater: Third series of compounds*. RIVM report 715810010, National Institute of Public Health and the Environment, Bilthoven, The Netherlands.

Kreule P, Swartjes FA. 1998. Proposals for Intervention Values for soil and groundwater, including the calculation of the human-toxicological serious soil contamination concentrations: Fourth series of compounds. RIVM report 711701005, National Institute of Public Health and the Environment, Bilthoven, The Netherlands.

Langley A. 1993. Refining exposure assessment. In: Langley AJ, Van Alphen M (eds) *The Health Risk Assessment and Management of Contaminated Sites*. Proceedings of the Second National Workshop. Monograph No. 2. South Australian Health Commission, Adelaide.

Langley AJ, El Saadi O (eds). 1991. Protocol for the Health Risk Assessment and Management of Contaminated Sites. Summary of a National Workshop on the Health Risk and Management of Contaminated Sites. South Australian Health Commission, Adelaide.

Langley AJ, Markey BR, Hill HS (eds). 1996. *The Health Risk Assessment and Management of Contaminated Sites*. Proceedings of the Third National Workshop. Monograph No. 5. South Australian Health Commission, Adelaide.

Langley AJ, Imray P, Lock W, Hill H (eds). 1998. *The Health Risk Assessment and Management of Contaminated Sites*. Proceedings of the Fourth National Workshop. Monograph No. 7. South Australian Health Commission, Adelaide.

Langmuir D, Chrowtowski P, Vigneault B, Chaney R. 2004. *Issue Paper on the Environmental Chemistry of Metals*. Report submitted to US EPA by ERG, Lexington MA.

Lijzen JPA, Baars AJ, Otte PF, Rikken MGJ, Swartjes FA, Verbruggen EMJ, van Wezel AP. 2001. *Technical Evaluation of the Intervention Values for Soil/Sediment and Groundwater*. RIVM report 711701 023, National Institute of Public Health and the Environment, Bilthoven, The Netherlands.

Lin Q, Wang Z, Ma S, Chen Y. 2006. Evaluation of dissipation mechanisms by *Lolium perenne* L, and *Raphanus sativa* for pentachlorophenol (PCP) in copper co-contaminated soil. *Science of the Total Environment* 368: 814–822.

Longhurst R, Roberts A, Waller J. 2004. Concentrations of arsenic, cadmium, copper, lead, and zinc in New Zealand pastoral topsoil and herbage, *New Zealand Journal of Agricultural Research* 47: 23-32.

McBride MB. 2002. Cadmium uptake by crops estimated from soil total Cd and pH. *Soil Science* 167: 62–67.

McLachlan MS. 1997. A simple model to predict accumulation of PCDD/Fs in an agricultural food chain. *Chemosphere* 34: 1263–1276.

McGrath SP. 1995. Chromium and nickel. In: *Heavy Metals in Soils*. Alloway BJ (ed) Chapter 7, pp. 152–178. 2nd edition. Blackie.

McKone TE. 1990. Dermal uptake of organic chemicals from a soil matrix. Risk Analysis 10: 407-419.

Meneses M, Schuhmacher M, Domingo JL. 2002. A design of two simple models to predict PCDD/F concentrations in vegetation and soils. *Chemosphere* 46: 1393–1402.

MfE. 1997. Guidelines for Assessing and Managing Contaminated Gasworks sites in New Zealand. Ministry for the Environment, Wellington.

MfE. 1999. *Guidelines for Assessing and Managing Petroleum Hydrocarbon Contaminated Sites in New Zealand*. Ministry for the Environment, Wellington.

MfE. 2003. Contaminated Land Management Guidelines No. 2: Hierarchy and Application in New Zealand of Environmental Guideline Values. Ministry for the Environment, Wellington.

MfE. 2004. Contaminated Land Management Guidelines No. 5: Site Investigation and Analysis of Soils, Ministry for the Environment, Wellington.

MfE. 2005. *National Environmental Standard for Contaminated Land*. Notes of Technical Review Workshop. Ministry for the Environment, Wellington.

MfE. 2006a. Identifying, Investigating and Managing Risks Associated with Former Sheep-dip Sites: A Guide for Local Authorities. Ministry for the Environment, Wellington.

MfE. 2006b. Working Towards a Comprehensive Policy Framework Managing Contaminated Land New Zealand: A Discussion Paper. Retrieved from http://www.mfe.govt.nz/issues/hazardous/contaminated /direction-land-management.html (October 2007).

MfE. 2007. Working Towards a Comprehensive Policy Framework for Managing Contaminated Land in New Zealand, Position Paper. Ministry for the Environment, Wellington.

MfE. 2010a. Proposed National Environmental Standard for Assessing and Managing Contaminants in Soil: Discussion Document. Ministry for the Environment, Wellington.

MfE. 2010b. *Plant bioconcentration factors for selected priority contaminants*. Unpublished report prepared by Landcare Research for the Ministry for the Environment, Wellington.

MfE. 2011. *Toxicological Intake Values for Priority Contaminants in Soil*. Ministry for the Environment, Wellington.

MfE and MoH. 1997. *Health and Environmental Guidelines for Selected Timber Treatment Chemicals*. Ministry for the Environment and Ministry of Health, Wellington.

MoH. 2008. Drinking-water Standards for New Zealand 2005 (revised 2008). Ministry of Health, Wellington.

Mikes O, Cupr P, Trapp S, Klanova. 2009. Uptake of polychlorinated biphenyls and organochlorine pesticides from soil and air into radishes (*Raphanus sativus*). *Environmental Pollution* 157: 488–496.

Moore DW & Taylor MD. 2009. A comparison of the Ministry for the Environment's proposed draft SGVs_(health) for arsenic and cadmium with New Zealand natural background soil concentrations. Unpublished paper presented at the WasteMINZ's 2009 Annual Conference.

Nable RO, Banuelos GS, Paull JG. 1997. Boron toxicity. Plant and Soil 193: 181-198.

NEPC. 1999a. National Environment Protection (Assessment of Site Contamination) Measure 1999, Schedule B(7b) Exposure settings and exposure scenarios. National Environment Protection Council, Adelaide.

NEPC. 1999b. National Environmental Protection (Assessment of Site Contamination) Measure (1999). National Environment Protection Council, Adelaide.

NEPC. 1999c. National Environment Protection (Assessment of Site Contamination) Measure 1999, Schedule B(7a) Guideline on Health-Based Investigation Levels. National Environment Protection Council, Adelaide.

NEPC. 1999d. National Environment Protection (Assessment of Site Contamination) Measure 1999, Schedule B(4) Guideline on Health Risk Assessment Methodology. National Environment Protection Council, Adelaide.

NEPC. 2006. Review Report – National Environmental Protection (Assessment of Site Contamination) Measure Review. National Environment Protection Council, Adelaide.

NEPC. 2010. National Environmental Protection (Assessment of Site Contamination) Measure 2011. National Environment Protection Council, Adelaide. Draft for Public Consultation (revised 1999 version).

OMEE. 1994. Scientific Criteria Document for Multimedia Environmental Standards Development – Lead. Ontario Ministry for Environment and Energy, Ontario.

Otte PF, Lijzen JPA, Otte JG, Swartjes FA, Versluijs CW. 2001. *Evaluation and Revision of the C Soil Parameter Set.* RIVM report 711701021, National Institute of Public Health and the Environment, Bilthoven, The Netherlands.

Paustenbach DJ. 2000. The practice of exposure assessment: a state-of-the-art review. *Journal of Toxicology and Environmental Health*, Part B, 3: 179–291.

Paustenbach DJ, Fehling K, Scott P, Harris M, Kerger BD. 2006. Identifying soil clean-up criteria for dioxins in urban residential soils: how have 20 years of research and risk assessment experience affected the analysis? *Journal of Toxicology and Environmental Health*, Part B, 9: 87–145.

Pirard C, de Pauw E. 2005. Uptake of polychlorinated-p-dioxins, polychlorinated dibenzofurans and coplanar polychlorobiphenyls in chickens. *Environment International* 31: 585–591.

Rikken MGJ, Lijzen JPA, Cornelese AA. 2001. *Evaluation of Model Concepts on Human Exposure*. RIVM Report 711701022, National Institute of Public Health and the Environment (RIVM), Bilthoven, The Netherlands.

Russell D, Parnell W, Wilson N. 1999. NZ Food: NZ People – Key Results from the 1997 National Nutrition Survey. Ministry of Health, Wellington.

Ryan JA, Bell RM, Davidson JM, O'Connor GA. 1988. Plant uptake of non-ionic organic chemicals from soils. *Chemosphere* 17: 2299–2323.

Sanson R, Cook A, Fairweather J. 2004. *A Study of Smallholdings and their Owners*. MAF Information Paper No: 53, Ministry of Agriculture and Forestry, Wellington.

Scheunert I, Qiao Z, Korte F. 1986. Comparative studies of the fate of atrazine-14C and pentachlorophenol-14C in various laboratory and outdoor soil-plant systems. *Journal of Environmental Science and Health*, Part B, 21: 457–485.

Schuler F, Schmid P, Schlatter C. 1997. The transfer of polychlorinated dibenzo-*p*-dioxins and dibenzofurans from soil into eggs of foraging chicken. *Chemosphere* 34: 711–718.

Sheppard SC, Evenden WG. 1992. Contaminant enrichment of sparingly soluble contaminants (U, Th and Pb) by erosion and by soil adhesion to plants and skin. *Environmental Geochemistry and Health* 14: 121–131.

Sparling G, Schipper LA. 2002. Soil quality at a national scale in New Zealand. *Journal of Environmental Quality* 31: 1848–1857.

Sparling G, Schipper LA. 2004. Soil quality monitoring arising in New Zealand: trends and issues arising from a broad-scale survey. *Agriculture, Ecosystems and Environment* 104: 545–552.

Sparling G, Schipper L, Bettjeman W & Hill R. 2004. Soil quality monitoring in New Zealand: practical lessons from a 6-year trial, *Agriculture, Ecosystems & Environment* 104: 523–534.

Sparling G, Shepherd TG, Schipper LA. 2000. Topsoil characteristics of three contrasting New Zealand soils under four long-term land uses. *New Zealand Journal of Agricultural Research* 43: 569–583.

Stanek EJ, Calabrese EJ. 1991a. A guide to interpreting soil ingestion studies: I. Development of a model to estimate the soil ingestion detection level of soil ingestion studies. *Regulatory Toxicology and Pharmacology* 13: 263–277.

Stanek EJ, Calabrese EJ. 1991b. Methodological considerations in soil ingestion estimation. *Chemical Speciation and Bioavailability* 3(314): 65–67.

Stanek EJ, Calabrese EJ. 1995a. Daily estimates of soil ingestion in children. *Environmental Health Perspectives* 103: 276–285.

Stanek EJ, Calabrese EJ. 1995b. Soil ingestion estimates for use in site evaluations based on the best tracer method. *Human Ecological Risk Assessment* 1: 133–156.

Stanek, EJ, Calabrese EJ. 2000. Daily soil ingestion estimates for children at a Superfund site. *Risk Analysis* 20(5): 627–635.

Stanek EJ, Calabrese EJ, Barnes RM, Pekow P. 1997. Soil ingestion in adults – results of a second pilot study. *Ecotoxicology and Environmental Safety* 36(3): 249–257.

Stanek EJ, Calabrese EJ, Zorn M. 2001. Soil ingestion distributions for Monte Carlo risk assessment in children. *Human and Ecological Risk Assessment* 7(2): 357–368.

Statistics New Zealand. Undated. *New Zealand: An urban/rural profile*. Statistics New Zealand. Retrieved from http://www.stats.govt.nz/Publications/BusinessPerformanceEnergyAndAgriculture/urban-rural-profile.aspx (February 2008).

Statistics New Zealand. 1999. New Zealand Time Use Survey: Users' Guide. Statistics New Zealand, Wellington.

Statistics New Zealand. 2006. A History of Survival in New Zealand: Cohort Life Tables 1876–2004, Statistics New Zealand, Wellington.

Statistics New Zealand. 2008. Demographic Trends: 2007. Statistics New Zealand, Wellington.

Stephens RD, Petreas MX, Hayward DG. 1995. Biotransfer and bioaccumulation of dioxins and furans from soil: chickens as a model for foraging animals. *Science of the Total Environment* 175: 253–273.

Swartjes FA. 1999. Risk based assessment of soil and groundwater quality in the Netherlands: Standards and remediation urgency. *Risk Analysis* 19: 1235–1250.

Swartjes FA, Walthaus H. 2007. *Risk-based Assessment of Soil and Groundwater Quality in The Netherlands (Dutch Soil Protection Act)*. In: Carlon C (ed). Derivation methods of soil screening values in Europe. A review and evaluation of national procedures towards harmonisation. European Commission, Joint Research Centre, Ispra, Italy EUR 22805-EN.

Swartjes FA, Dirven-Van Breemen EM, Otte PF, Van Beelen P, Rikken MGJ, Tuinstra J, Spijker J, Lijzen JPA. 2007. *Human Health Risks Due to Consumption of Vegetables from Contaminated Sites. Towards a Protocol for Site-specific Assessment.* RIVM report 711701040 / 2007, National Institute of Public Health and the Environment (RIVM), Bilthoven, The Netherlands.

Taylor ER. 1991. *How Much Soil Do Children Eat. The Health Risk Assessment and Management of Contaminated Sites.* Proceedings of a National Workshop on the Health Risk Assessment and Management of Contaminated Sites. El Saadi O and Langley A (eds). South Australian Health Commission. 72–83 (Appendix I).

Taylor MD, Hill RB. 2010. Should Targets for Soil Quality Monitoring be Gravimetric or Volumetric Based? *Soil News* 58: 66–69.

Taylor MD, Gibb R, Willoughby J, Hewitt A, & Arnold G. 2007. *Soil Maps of Cadmium in New Zealand*. Landcare Research Contract Report: LC0607/84.

T&T and SPHERE. 2008. Assessment of Dioxin Contamination at Sawmill Sites. Report to the Ministry for the Environment, Tonkin and Taylor Limited and SPHERE. Retrieved from http://www.mfe.govt.nz/publications/hazardous/assessment-dioxin-contamination-sawmill-sites-2008-10/index.html (July 2009).

Trapp S, Matthies M. 1995. Generic one-compartment model for uptake of organic chemicals by foliar vegetation. *Environmental Science and Technology* 29: 2333–2338.

Travis CC, Arms AD. 1988. Bioconcentration of organics in beef, milk, and vegetation. *Environmental Science and Technology* 22(3): 271–274.

USDOE. 1999. *Guidance for Conducting Risk Assessments and Related Risk Activities for the DOE-ORO* Environmental Management Program, BJC/OR-271, Office of Environmental Management, United States Department of Energy.

US EPA. 1989a. Risk Assessment Guidance for Superfund (RAGS): Volume I. Human Health Evaluation Manual (HHEM) (Part A, Baseline Risk Assessment), Interim Final, EPA/540/1-89/002. United States Environmental Protection Agency, Washington DC.
US EPA. 1989b. *Exposure Factors Handbook*. EPA/600/8-89/043, Office of Health and Environmental Assessment, Office of Research and Development, United States Environmental Protection Agency, Washington DC.

US EPA. 1991a. *Risk Assessment Guidance for Superfund, Volume 1, Human Health Evaluation Manual, Part B, Development of Preliminary Remediation Goals*, EPA/540/R-92/003, Office of Emergency and Remedial Response, United States Environmental Protection Agency, Washington DC.

US EPA. 1991b. *Risk Assessment Guidance for Super Fund, Volume I, Part C: Risk Evaluation of Remedial Alternatives*, Office of Emergency and Remedial Response, United States Environmental Protection Agency, Washington DC.

US EPA. 1991c. Risk Assessment Guidance for Superfund (RAGS), Volume I, Human Health Evaluation Manual Supplemental Guidance: "Standard Default Exposure Factors." Interim Final, Office of Emergency and Remedial Response, United States Environmental Protection Agency, Washington DC.

US EPA. 1992. *Guidelines for Exposure Assessment*, EPA/600/Z-92/00, Risk Assessment Forum United States Environmental Protection Agency, Washington DC.

US EPA. 1994. Method 200.2, Sample Preparation Procedure for Spectrochemical Determination of Total Recoverable Elements, Revision 2.8, Environmental Monitoring Systems Laboratory, Office of Research and Development, United States Environmental Protection Agency, Cincinnati, Ohio. Retrieved from http://water.epa.gov/scitech/swguidance/methods/methods_index.cfm (July 2010).

US EPA. 1996a. *Soil Screening Guidance: User's Guide*, Second Edition Office of Emergency and Remedial Response, Publication 9355.4-23, United States Environmental Protection Agency, Washington DC.

US EPA. 1996b. *Soil Screening Guidance: Technical Background Document*, 2nd edition, EPA/540/R95/128, Office of Emergency and Remedial Response, US Environmental Protection Agency, Washington DC.

US EPA. 1997. *Exposure Factors Handbook*. Office of Research and Development, US Environmental Protection Agency, Washington DC. Retrieved from http://www.epa.gov/ncea/efh/report.html (February 2008).

US EPA. 2001a. *Risk Assessment Guidance for Super Fund Volume I, Part E: Supplemental Guidance for Dermal Risk Assessment*, United States Environmental Protection Agency, Washington DC.

US EPA. 2001b. *Risk Assessment Guidance for Superfund: Volume III – Part A, Process for Conducting Probabilistic Risk Assessment*. Office of Emergency and Remedial Response, United States Environmental Protection Agency, Washington DC.

US EPA. 2001c. Risk Assessment Guidance for Super Fund, Volume II, Part D: Standardised Planning, Reporting, and Review of Superfund Risk Assessments. Office of Emergency and Remedial Response, United States Environmental Protection Agency, Washington DC.

US EPA. 2002a. Supplemental Guidance for Developing Soil Screening Levels for Superfund Sites, OSWER 9355.4-24. Office of Emergency and Remedial Response, United States Environmental Protection Agency, Washington DC.

US EPA. 2002b. *Child Specific Exposure Factors Handbook, Interim Report.* EPA-600-P-00-002B, Office of Research and Development, US Environmental Protection Agency, Washington DC.

US EPA. 2002c. Draft Guidance for Evaluating the Vapour Intrusion to Indoor Air Pathway from Groundwater and Soils (Subsurface Vapour Intrusion Guidance). Office of Emergency and Remedial Response, US Environmental Protection Agency, Washington DC. Retrieved from http://www.epa.gov/correctiveaction/eis/vapor.htm (May 2008).

US EPA. 2003. *Guidance for Developing Ecological Soil Screening Levels. Attachment 4.1 Exposure Factors and Bioaccumulation Models for Derivation of Wildlife Eco-SSL*. OSWER Directive 928.7-55, Office of Research and Development, US Environmental Protection Agency, Washington DC.

US EPA. 2004a. An Examination of EPA Risk Assessment Principles and Practices. Staff Paper prepared for the US Environmental Protection Agency by members of the Risk Assessment Task Force, EPA/100/B-04/001, Office of the Science Advisor, United States Environmental Protection Agency, Washington DC.

US EPA. 2004b. User's Guide for Evaluating Subsurface Vapoir Intrusion into Buildings. Prepared by Environmental Quality Management Inc, Office Of Emergency and Remedial Response, US Environmental Protection Agency, Washington DC, Revised February 2004. Retrieved from http://www.epa.gov/oswer/riskassessment/airmodel/johnson_ettinger.htm (May 2008).

US EPA. 2004c. *Risk Assessment Guidance for Superfund Volume I: Human Health Evaluation Manual (Part E, Supplemental Guidance for Dermal Risk Assessment)*, Final. EPA/540/R/99/005, Office of Superfund Remediation and Technology Innovation, US Environmental Protection Agency, Washington DC.

US EPA. 2006a. *Child Specific Exposure Factors Handbook (external review draft)*. EPA/600/R/06/096A. Office of Research and Development, US Environmental Protection Agency, Washington DC.

US EPA. 2006b. RCRA Orientation Manual 2006. Office of Solid Waste, US Environmental Protection Agency.

US EPA. 2007. Guidance for Developing Ecological Soil Screening Levels (Eco-SSLs), Attachment 4-1, Exposure Factors and Bioaccumulation Models for Derivation of Wildlife Eco-SSLs. Revised April 2007, OSWER Directive 9285.7-55, Office of Solid Waste and Emergency Response, US Environmental Protection Agency, Washington DC.

US EPA. 2008. *Preliminary Remediation Goals User's Guide*. United States Environmental Protection Agency. Retrieved from http://www.epa.gov/reg3hwmd/risk/human/rb-concentration_table/ usersguide.htm (September 2008).

Van den Berg M, Birnbaum L, Denison M, de Vito M, Farland W, Feeley M, Fiedler H, Hakansson H, Hanberg A, Haws L, Rose M, Safe S, Schrenk D, Tohyama C, Tritscher A, Yuomisto J, Tyskland M, Walker N, Peterson RE. 2006. The 2005 World Health Organization reevaluation of human and mammalian toxic equivalency factors for dioxins and dioxin-link compounds. *Toxicological Sciences* 93: 223–241.

Van Holderbeke M, Cornelis C, Bierkens J, Torts R. 2007. *Review of the Soil Ingeston Pathway in Human Exposure Assessment*. Study in support of the BenNeKempen project, Flemish Institute for Technological Research (VITO).

Van Wijnen JH, Clausing P, Brunekreff B. 1990. Estimated soil ingestion by children. *Environmental Research* 51: 147–162.

Vannoort R, Cressey P, Silvers K. 2000. *1997/1998 New Zealand Total Diet Survey. Part 2: Elements.* Ministry of Health, Wellington. Retrieved from http://www.moh.govt.nz (May 2008)

Vannoort RW. 2003. 2003/2004 New Zealand Total Diet Survey: Analytical Results – Q1. Report for New Zealand Food Safety Authority, Wellington.

Van Overmeire I, Pussemier L, Hanot V, De Temmerman L, Hoenig M, Goeyens L. 2006. Chemical contamination of free-range eggs from Belgium. *Food Additives and Contaminants* 23: 1109–1122.

Van Overmeire I, Wageneers N, Sioen I, Bilau M, De Henauw S, Goeyens L, Pussemier L, Eppe G. 2009. PCDD/Fs and dioxin-like PCBs in home-produced eggs from Belgium: Levels, contamination sources and health risks. *Science of the Total Environment* 407: 4419–4429.

Versar. 2001. Technical Issue Paper: Age Group Recommendations for Assessing Childhood Exposure and the Adequacy of Existing Exposure Factors Data for Children. Prepared for: Risk Assessment Forum, US Environmental Protection Agency, Washington DC, by Versar Inc. Springfield, VA.

Versluijs CW, Otte PF. 2001. Accumulation of Metals in Plants. A contribution to the technical evaluation of the intervention values and site-specific risk assessment of contaminated sites (in Dutch). RIVM report 711701024. National Institute for Public Health and the Environment, Bilthoven, The Netherlands.

Victoria Government. 2002. State Environment Protection Policy, Prevention and Management of Contamination of Land. Victoria Government Gazette, No. S 95, Tuesday 4 June 2002.

VROM. 1999. From Funnel to Sieve. Ministry of Housing, Spatial Planning and the Environment, Bilthoven, The Netherlands.

VROM. 2000. *Ministerial Circular on Target and Intervention Values*. DBO/1999226863. 4 February 2000, Ministry of Housing, Spatial Planning and Environment, The Hague.

Waitz MFW, Freijer JI, Kreule P, Swartjes FA. 1996. *The VOLASOIL Risk Assessment Model Based on CSOIL for Soils Contaminated with Volatile Compounds*. RIVM report 715810014, National Institute for Public Health and the Environment, Bilthoven, The Netherlands.

Waegeneers N, De Steur H, De Temmerman L, Van Steenwinkel S, Gellynck X, Vianene J. 2009. Transfer of soil contaminants to home-produced eggs and preventative measures to reduce contamination. *Science of the Total Environment* 407: 4438–4446.

Wester RC, Maibach H I, Sedik L, Melendres J, Wade M, DiZio S. 1993. Percutaneous absorption of pentachlorophenol from soil. *Fundamental and Applied Toxicology* 20: 68–71.

Wild SR, Berrow ML, McGrath S, Jones KC. 1992. Polynuclear aromatic hydrocarbons in crops from long-term field experiments amended with sewage sludge. *Environmental Pollution* 76: 25–32.

WHO. 1998. Environmental Health Criteria 202. Selected Non-heterocyclic Polycyclic Aromatic Hydrocarbons. International Programme on Chemical Safety, World Health Organization: Geneva.