

Strategy & Policy Directorate

Land Quality Assessment (LQA)

Land Quality Management Practitioner Guide 2017/01

Document Aim:

MOD Policy on land contamination is set out in the Land Contamination Leaflet in Volume 2 of JSP 418. This Guide provides the framework for carrying out MOD Land Quality Assessments (LQA) and should be used by MOD practitioners, Project Teams, PFI Partners, Industry Service Providers and contractors responsible for the assessment and management of land contamination and associated liabilities. This guide sets out the LQA process together with guidance and signposts to key MOD policies, legislation and Industry Guides. The level and detail of the guidance reflects MOD's experience to date and highlights areas where attention and care is needed in applying industry standards. It also identifies the points at which decisions are required and provides guidance on current good practice within the context of the MOD LQA process as applied to the defence estate. This guide supersedes Practitioner Guide 07/2012.

Document Synopsis:

This Practitioner Guide sets out the approach that should be taken to MOD LQA practice and process together with useful guidance.

The guide covers every LQA phase and provides details of reporting formats that are to be used as well as the risk assessment process that is to be followed.

This guide is aimed at experienced practitioners, be they MOD or Industry Service Providers. It pulls together good practice from across the industry and integrates key principles into a single coherent document which is focussed on the defence estate and some of its unique aspects.

Points of contact are listed where advice can be sought from the relevant MOD subject matter experts.

Defence Infrastructure Organisation



Estate Management





MINISTRY OF DEFENCE

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Equality And Diversity Impact Assessment

This policy has been Equality and Diversity Impact Assessed in accordance with the Department's Equality and Diversity Impact Assessment Tool against:

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Related Documents

JSP 418 Volume 2 leaflet 2	
PI 02/15 Asbestos Management on the defence estate	
PI 03/15 CDM (construction (design & management))	
regulations	

GLOSSARY

AGS	Association of Geotechnical and Geoenvironmental Specialists
CDM	Construction, Design and Management Regulations
CESO	Chief Environment and Safety Officer
CIEH	Chartered Institute of Environmental Health
CIRIA	Construction Industry Research and Information Association
CIWEM	Chartered Institution of Water and Environmental Management
CL:AIRE	Contaminated Land: Applications in Real Environments
CLEA	Contaminated Land Exposure Assessment
CLR	Contaminated Land Report
CoP	Code of Practice
CRA	Closure Risk Assessment
CSM	Conceptual Site Model
CUGs	Clean-Up Goals
DEFRA	Department for Environment, Food and Rural Affairs
DIO	Defence Infrastructure Organisation
DQRA	Detailed Quantitative Risk Assessment
DWS	Drinking Water Standards
EA	Environmental Agency
EIC	Environmental Industries Commission
EOLM	Environmental and Ordnance Liability Management
EM	Environmental Manager
EMS	Environmental Management System
EOC	Explosive Ordnance Clearance
EOD	Explosive Ordnance Disposal
EPS	Environmental Planning Support
EQS	Environmental Quality Standards
GAC	Generic Assessment Criteria
GIS	Geographic Information System
GQRA	Generic Quantitative Assessment
GSV	Gas Screening Values
HSE	Health and Safety Executive
IEMA	The Institute of Environmental Management and Assessment
LCF	Land Condition File
LCR	Land Condition Record
LMS	Land Management Services
LQA	Land Quality Assessment
LQM	Land Quality Management Ltd
LQS	Land Quality Statement

MCERTS	Monitoring Certification Scheme
MMP	Materials Management Plan
MOD	Ministry of Defence
NAAFI	Navy Army Air Force Institute
NDPB	Non-Departmental Public Body
NGO	Non-Governmental Organisation
NHBC	National House Building Council
NIEA	Northern Ireland Environment Agency
OCE	Order of Cost Estimate
OME	Ordnance, Munitions and Explosives
PFI	Private Finance Initiative
PPC	Pollution Prevention and Control
PRA	Preliminary (Qualitative) Risk Assessment
PTS	Professional Technical Services
QP	Qualified Person
QRA	Quantitative Risk Assessment
RCLEA	Radioactively Contaminated Land Exposure Assessment
RPL	Relevant Pollutant Linkage
RSGV	Radioactivity in Soil Guideline Values
RTM	Remedial Targets Methodology
SEPA	Scottish Environment Protection Agency
SGV	Soil Guideline Values
SiLC	Specialist in Land Condition
SME	Subject Matter Expert
SoBRA	Society of Brownfield Risk Assessment
SoKH	Statement of Known Hazards
SOR	Statement of Requirement
SPOSH	Significant Possibility of Significant Harm
SRPs	Safety Rules and Procedures
SSAC	Site Specific Assessment Criteria
SSSI	Sites of Special Scientific Interest
SSV	Soil Screening Value
SQEP	Suitably Qualified and Experienced Persons
SURF UK	United Kingdom's Sustainable Remediation Forum
TLB	Top Level Budget Holder
UKAS	United Kingdom's Accreditation Service
UXO	Unexploded Ordnance
UXOPRA	UXO Preliminary Risk Assessment
VFM	Value for Money

VOC Volatile Organic Compound

WSV Water Screening Value

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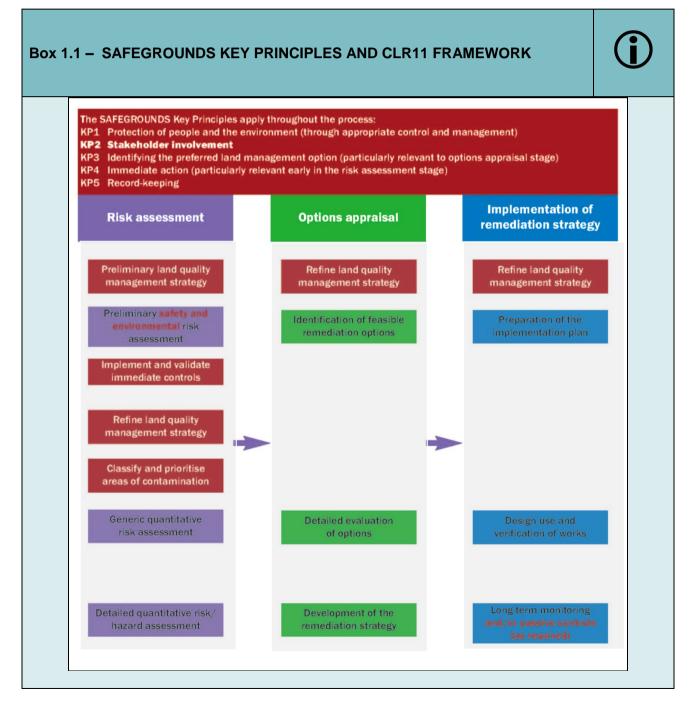
Legend

Legal Requirements and Mandatory Practice These boxes identify the relevant aspects of the EU Directives and Legislation and complying with MOD Mandatory Practice.	ик
Hints and Tips These boxes provide hints and tips for complying with Advisory pra	actice.
Illustrative Examples Generic examples have been used to illustrate the recommended a within the guidance. These examples have been drawn from real N to aid their interpretation.	
Key Information These boxes provide key information of relevance to the assessme management of land contamination on the defence estate.	ent and
Key Guidance This box signposts key guidance that will aid the practitioner.	Sign Post

1. INTRODUCTION

1.1 This guide sets out the approach to be used by practitioners be they MOD staff, contractors or industry partners responsible for the assessment and management of land contamination and the associated liabilities across the defence estate as well as those involved in the development, purchasing, sale or lease of land affected by contamination. A base level of knowledge has been assumed and this guide is not intended to constitute a step by step manual, instead it sets out the process and approach including required MOD reporting formats. The level of detail has been tailored and targeted and key guidance and policy signposted. Key points of contact for advice and support are given in Annex A and consultation at an early stage is recommended.

1.2 This guide is structured to signpost relevant policy, legislation and guidance within the framework provided by the Safegrounds Key Principles (see Box 1.1) and Contaminated Land Report 11, Model Procedures for the Management of Land Contamination (CLR11).



1.3 The key guidance that will be referred to throughout this guide is detailed in Box 1.2.

Box 1.2 – KEY GUIDANCE

DEFRA Guidance 04/12: EPA 1990: Part 2A Contaminated Land Statutory Guidance, 2012. DEFRA

DECC Guidance on Radioactive Contaminated Land 2012

The Contaminated Land Statutory Guidance for Wales 2012, Welsh Government

EPA 1990: Part 2A Contaminated Land Statutory Guidance: Ed 2 2006, Scottish Government

Contaminated Land Report 11. Model Procedures for the Management of Land Contamination, 2004. DEFRA and EA.

R&D 66 Guidance on the Safe Development of Housing on Land Affected by Contamination, 2008 Volume 1 and 2. NHBC and EA

Towler, P et al Safegrounds LMG V2 W29 Good practice guidance for the management of contaminated land on nuclear licensed and defence sites CIRIA London 2009

BS 10175:2011+A1:2013. Investigation of potentially contaminated sites – Code of practice, 2011, BSI Standards Publication

Communicating understanding of contaminated land risks, 2010, SNIFFER

MOD JSP 418: Volume 2, Leaflet 2 Contaminated Land

DE (DIO) Design and Maintenance Guide 12 – site closure guide, Land and Property Policy: PI 6/2005.

Industry Guidance: Qualitative Risk Assessment for Land Contamination, including Radioactive Contamination, June 2012. Nuclear Industry Group for Land Quality

SEPA Guidance on monitoring for heterogeneous Radium-226 sources resulting from historic luminising or waste disposal sites. Draft document, August 2016.

2. MOD LQA POLICY, GUIDANCE, COMPLIANCE AND FUNDING RESPONSIBILITY

Introduction

2.1 MOD policy on the assessment and management of land contamination is detailed within: JSP 418 Vol.2 – Land Contamination. The following section provides additional clarification and guidance with respect to compliance with MOD policy.

2.2 Practitioners should be up to date with both current MOD policy and the contaminated land regulatory regime as it operates within the UK bearing in mind there are slight differences with and between the devolved administrations. Equally practitioners should be clear as to the role of the local authority versus that of the Environment Agency (EA), Scottish Environment Protection Agency (SEPA) and Northern Ireland Environment Agency (NIEA) as well as the basis on which a site is deemed to be a 'special site'. The latter is summarised in Box 2.1 for ease of reference.

Box 2.1 Role of the Regulators and 'Special Sites; that apply to MOD

The Local Authority is the enforcing authority for all sites which are not deemed 'special sites'. The EA and SEPA etc are the enforcing authority for 'special sites'. SEPA is the lead regulator with respect to radioactively contaminated land in Scotland.

The designation of a Special Site cannot take place until the land in question has been formally identified as Contaminated Land by the Local Authority and it meets one or more of the descriptions prescribed in the Regulations. The descriptions for Special Sites include: Any Contaminated Land either located at or is adjacent to current military, naval and air forces bases and other properties, including those of:

- Visiting forces;
- The Atomic Weapons Establishment;
- Certain lands at Greenwich Hospital;

• All land currently or formerly used for the manufacture, production, or disposal of chemical and biological weapons and related materials, regardless of current ownership;

- Land used in the manufacture of explosives; and
- Land which is contaminated land wholly or partly by the presence of radioactivity.
- The descriptions for Special Sites exclude:
 - o Off-base housing and Navy Army Air Force Institute (NAAFI) premises;
 - Property disposed of to civil ownership and occupation; and
 - Privately owned training areas and ranges which are used occasionally by the MOD.

Assessment and Management of Land Contamination

2.3 The drivers behind the assessment of land quality across the defence estate are:

• The secretary of State has a statutory duty to ensure that there are suitable and sufficient processes and procedures in place to both protect the health, safety and welfare of personnel, contractors and visitors on their establishments and the environment.

• To meet statutory requirements of planning controls and the environmental protection legislation to ensure land is suitable for use and substances present do not pose an unacceptable risk to humans or the wider environment.

• Failure to adequately assess and manage land contamination on the defence estate has the potential to impact on defence capability through reducing the availability of training areas,

limiting development of the estate, reducing disposal receipts and diversion of funding to meet statutory clean-up obligations.

2.4 MOD has committed to undertake a programme of LQAs to: "assess the land quality across the defence estate in order to provide a proper knowledge of the condition of the estate and ensure that it is 'a suitable for use' and not causing harm to the environment. Where it is identified that unacceptable risk is posed by presence of contamination, action must be taken to reduce and control the risks to an acceptable level".

2.5 An appropriate LQA is required to cover all land owned or occupied for defence purposes and for all property transfers. If it can be shown at an early stage that risk is low, then it may be unnecessary to proceed to the next phase.

2.6 Site Users are responsible for using land within agreed parameters or constraints. DIO is responsible for:

a. Managing an integrated and prioritised MOD estate wide LQA programme;

b. Ensuring that funding is in place for the LQA programme and any necessary remediation;

c. Coordinating/facilitating the compilation of the information required for the LQA as specified in this Management Guide;

d. Maintaining the catalogue and electronic library of LQA reports;

e. Maintaining establishment level records on known and suspected land contamination where DIO is responsible for the Infrastructure; and

f. Providing advice to Commanding Officer/Head of Establishment on the management of land contamination risks.

2.7 MOD will meet its statutory commitments and take voluntary action where a risk of significant harm to health and safety or the environment is confirmed and the MOD is the 'appropriate person' to bear the responsibility for remediation action. It is MOD policy to inform the appropriate Regulatory Authority if a risk of significant harm or significant environmental pollution is identified and agree with them the necessary remediation action.

2.8 The location together with details of the associated hazards and risks associated with land contamination identified by a LQA must be transferred to the establishment hazard register and where available the Land Condition File (LCF). This will ensure that land contamination is considered as part of the arrangements for notifying known site hazards to site users, Facility and Project Managers, contractors and visitors prior to commencement of their activities. Where significant land contamination risks have been identified, these should be regularly reviewed as part of the site Health and Safety and Environmental Management Systems (EMSs).

2.9 There are a number of methods for managing the risks associated with land contamination. These range from the removal of the contaminant, various physical, chemical and biological treatments or breaking the pollutant linkage by restricting access to the affected area. The choice of management response will be site-specific and depend upon the nature and extent of the contamination, the level of risk and the cost benefit. Where land contamination exists sites can still be suitable for MOD usage and may remain an asset if managed appropriately. Remediation may be a requirement for a change of use or development on a retained establishment.

Site Acquisition and Leasing

2.10 For the acquisition of land after 1990 it is likely that MOD will be deemed to have accepted financial liability for any necessary investigation and subsequent remediation of land contamination that pre-dates MOD ownership/occupation unless otherwise specified in the terms of the contract.

2.11 A reliable and robust LQA must therefore be undertaken to establish the land condition and potential health, environmental and liability risks prior to purchasing or leasing land. As a minimum such a LQA must comply with the requirements of this guide and current best practice.

2.12 Advice on specific LQA requirements is available through the contacts listed at Annex A. For advice on all other aspects of land acquisition and leasing as part of projects etc the local DIO Land Management Services (LMS) representatives must be consulted.

Divestment/Disposal

2.13 Known or suspected contaminated land can have a significant impact on potential use and disposal value. When deciding which establishments to release from MOD ownership, it is important to understand the nature and magnitude of the contaminated land liability. Hence, it is essential that the nature, extent and associated health, environmental and liability risks are adequately quantified and the LQA is sufficiently robust to provide for auditable and defensible decision making.

2.14 A site cannot be divested without:

- An Unexploded Ordnance Preliminary Risk Assessment (UXOPRA) and explosive ordnance clearance (EOC) if required. Reports should include details of any instrument search, intrusive investigation and clearance/disposal activities carried out on the site;
- An appropriate independent LQA, supported by a collateral warranty is required. The phases of assessment required will be depend upon the site situation; and
- A Closure Risk Assessment (CRA)

2.15 It is essential that a robust independent LQA is prepared as part of a site disposal to inform the defence against compensation claims arising from any post disposal contamination by the new or subsequent land owner(s). It will also enable MOD to take advantage of the mechanisms available under the UK Contaminated Land Regime for the transfer of the financial liability in respect of clean up to the purchaser. The LQA must be supported by a collateral warranty in order to provide the necessary assurance to a purchaser, their funders and insurers. Similarly site redevelopments under PFIs etc. will require robust independent LQAs supported by collateral warranties in order to establish the baseline land quality that can be relied upon by the PFI partner.

Box 2.2 – KEY GUIDANCE

MOD JSP 364 Joint Service EOD Manual

MOD JSP 403 Handbook of Defence Ranges Safety

MOD JSP 418: Volume 2, Leaflet 2 Contaminated Land

DE (DIO) Design and Maintenance Guide 12 – site closure guide, Land and Property Policy: PI 6/2005.

C681 – Unexploded Ordnance (UXO) A guide for the construction industry, 2009. CIRIA

2.16 Responsibility for organising and funding the LQA and any subsequent work for divestment and disposal, where required, currently falls to DIO LMS with delivery through DIO EPS EOLM.

2.17 Though written before the formation of DIO, the extant guidance for those DIO and site staff involved in the site closure process including the required outputs is detailed in: DE (DIO) Design and Maintenance Guide 12 – site closure guide, Land and Property Policy: PI 6/2005.

2.18 For disposal sites, remediation is generally confined to the removal of ordnance (where an unacceptable risk is present), or other defence specific contaminants such as chemical agents, radioactive and microbiological materials, where a civilian contractor might not have the relevant experience.

2.19 See Annex A for contact details to obtain additional guidance on the appropriateness of a particular level of LQA required to support a disposal.

Collateral Warranty

2.20 Third Parties such as purchasers, landlords, PFI partners and/or their funder(s) will seek reassurance that LQAs produced for MOD sites and landholdings, be it freehold or leasehold, are independent and reliable. Hence, it is usually the case that purchasers, PFI partners and their funders require internal MOD investigations to be checked and verified by independent specialists, often at MOD's expense. To overcome this and assist in maximising the sale receipt MOD policy is to commission independent LQAs supported by collateral warranties.

2.21 Box 2.4 provides a general overview of the form and function of collateral warranties together with the minimum requirement necessary to support a site disposal. Further guidance on the requirement for and form of collateral warranties is available from DIO LMS and DIO EPS EOLM. See Annex A for contact details.

Box 2.4 Collateral Warranties



Collateral warranties can be either 'agreements' or deeds and typically extend the duty of care of an author of a LQA to a third party, such as a purchaser or PFI partner, for a period of 12 years during which time they require the author to maintain a specified amount of Professional Indemnity insurance cover.

The exact form of the warranty, i.e. whether it is an agreement or a deed, the number of assignments possible and any associated costs will be dependent upon the circumstances, but for site disposal purposes the minimum requirement and conditions acceptable to DIO is:

'Provision of Collateral Warranties in the standard agreed form, at no additional cost, to the PFI Partner (where appropriate), first purchaser and/or tenant of the whole site or part thereof to a limit of two parts (or the number of parts where specified in the Invitation to Tender/Task Order Form), and to the first funder of those parties. Further collateral warranties should also, at the reasonable request of MOD, be provided in the standard agreed form to second purchasers and/or tenants and their funders of all or part of the site ("Secondary Warranties") at a reasonable fee per warranty not to exceed £1000. Should any party eligible to benefit from the Secondary Warranty require variations from the agreed standard form, the Consultant shall be entitled to levy additional fees and/or expenses to reflect the reasonable costs in negotiating such variations. The level of Professional Indemnity cover and form of the Collateral Warranty shall be agreed between the Consultant and the party eligible for the warranty, and will not exceed £5M in aggregate unless agreed otherwise.

Alienated Estate

2.22 DIO is responsible for the assessment and where necessary remediation of former MOD sites within the context of Part 2A of the Environmental Protection Act 1990. The lead element of DIO being: Land Management Services supported by DIO EPS EOLM.

Box 2.5 – MOD Position Statement – Alienated Estate

Alienated Land

1. In the case of land previously owned or occupied by the Ministry of Defence, the Ministry of Defence will look to the regulatory authority to act in accordance with the provisions of the Contaminated Land Regime as set out under Part 2A and the Statutory Guidance. Further the Ministry of Defence will look to the regulator to demonstrate that land is 'contaminated land' within the definition provided under Part 2A of the Environmental Protection Act and Statutory Guidance and the Ministry is an 'Appropriate Person'.

2. Where the Ministry of Defence is found to be an 'Appropriate Person', it will fulfil its legal obligation to meet its portion of the liability and carry out voluntary action including remediation where appropriate. In cases of two or more Appropriate Persons being identified by the regulator, the Ministry of Defence will work with the other Appropriate Person(s) and interested parties including the regulatory authorities to reach agreement on the management actions required and the necessary funding.

3. LQA AND MANAGEMENT PROCESS

Objectives

3.1 A robust assessment of land quality is essential to inform investment, development and divestment decisions and where necessary, identify appropriate remediation options and pollution prevention and control measures. The objective of a LQA is therefore, to quantify the contaminated land risks and the associated liability (health, environmental, legal and financial) in a logical and rational manner achieving both economy in the expenditure of resources and confidence in the end result in the process, such that the LQA provides the basis for defensible and auditable decision making. For this reason, a phased approach to this stage of the investigation process is recommended. As soon as sufficient information is obtained the investigation should cease.

Process

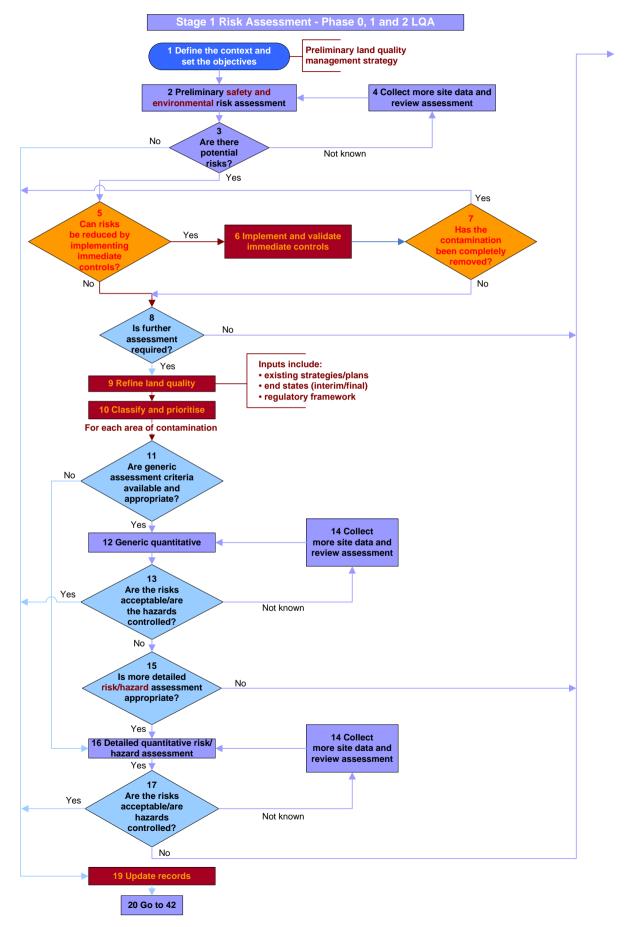
3.2 Figure 1, taken from the Safegrounds Land Management Guide, shows the process diagram for the assessment and management of land contamination adapted from CLR11 to take account of the Safegrounds key principles and aspects specific to radiological contamination on defence sites. This is a systematic process which follows three stages: 1. Risk Assessment, 2. Options Appraisal, and 3. Implementation of the Remediation (Management) Strategy.

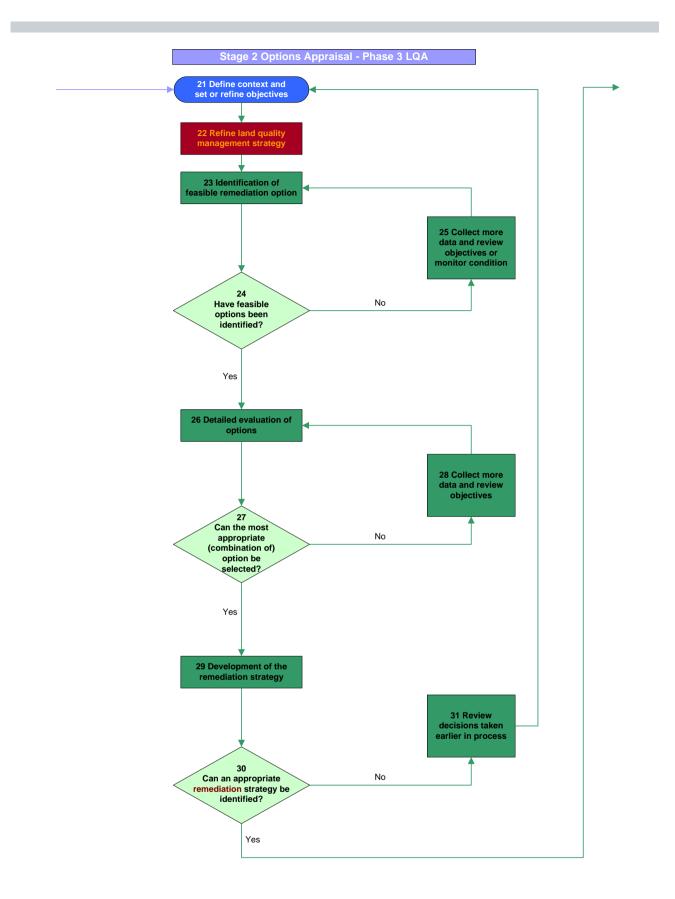
3.3 The systematic process outlined in Figure 1 can be best achieved by adopting a phased approach to the assessment and management of land contamination in line with the MOD Land Quality Management Strategy and site specific strategies. Within MOD this is achieved using the LQA process which is divided into the phases shown in Box 3.1, each of which has been superimposed onto Figure 1. These phases differ in terms of the terminology adopted by the recent BS 10175:2011+A1:2013 but are compatible and remain consistent with CLR11.

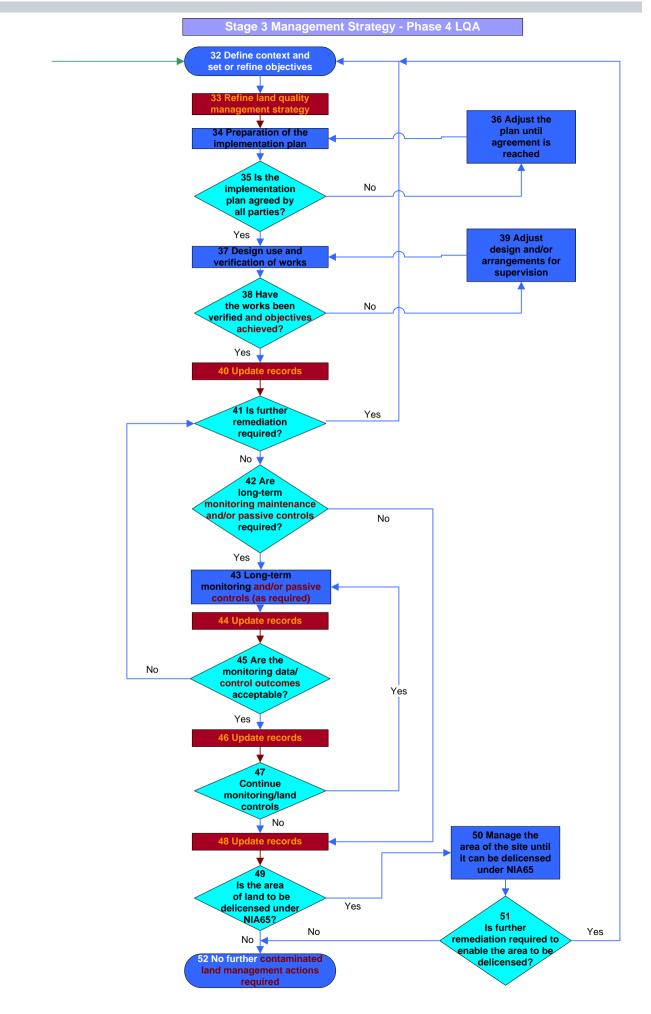
Box 3.1 MOD LQA Process Phases	í	
STAGE 1 - Risk Assessment		
PHASE 0 LQA - Preliminary Hazard Assessment;		
PHASE 1 LQA - Desk Study; and		
PHASE 2 LQA - Site Investigation (this may be phased).		
STAGE 2 - Options Appraisal		
PHASE 3 LQA - Management Option Appraisal/Decisions.		
STAGE 3 - Management Response		
 PHASE 4 LQA - Implementation of Management Option(s) (Management Response – this may also be phased and involve long term monitoring). 		

Figure 1 Generic Flow Diagram for Management of Contaminated Land

Note: The modifications to the CLR11 decision flow diagram for SAFEGROUNDS are highlighted in dark red boxes with tan lettering.







3.4 It will not be necessary to carry out every phase for each site. For instance the potential environmental, health and safety risks identified at Phase 0 may be sufficiently quantified not to warrant proceeding to Phase 1 and so on. The need for further investigation and remediation must be commensurate with the objectives of the LQA and the degree of confidence required in the decisions.

3.5 The decision on whether to progress a LQA to the next phase must be taken after consultation with key stakeholders, see Box 3.2, and advice from the relevant specialists detailed at Annex A, taking account of the situation and cost benefit. In the case of sites in disposal, alienated land and the MOD Estate-wide Phase 2 LQA Programme, DIO will take the lead with respect to stakeholder involvement.

Box 3.2 Stakeholder Involvement



Stakeholders are individuals or groups with a legitimate interest in the management of land contamination. They range from regulators and employees through to the Head of Establishment, Non-Governmental Organisations and local residents.

The level of involvement namely communication (including provision of information), consultation and participation in the decision making process will depend on the situation and context.

Responsibility for the final decisions on the management of land contamination falls within the jurisdiction of the DIO.

It is more effective to involve stakeholders throughout the planning and decision process/cycle rather than intermittently on individual issues.

Specific guidance on stakeholder involvement is available in: Safegrounds, Community Stakeholder Involvement. A report prepared within the Safegrounds project.

3.6 The cost of and time taken to complete each phase will be dependent upon the nature of the site activities, setting, accessibility and to an extent the size and complexity of the site/establishment in question.

3.7 Guidance on reporting formats for LQA Reports, Technical Notes and Land Quality Statements (LQS's) are presented at Annex B1 to B3. These are the MOD standard formats which are to be adopted and tailored to the site and situation.

Policy Process and Responsibility

3.8 Responsibility for procuring and funding an LQA rests with DIO EPS EOLM with the exception of those commissioned as part of a disposal, project or PFI etc. In the case of the latter technical advice and assistance is available from DIO EPS EOLM and the SME contacts given in Annex A. The TLB however, retains overall responsibility for the establishment, this includes all Health, Safety and Environmental issues in addition to being responsible for ensuring access to the site, the provision of key personnel and that the necessary documents etc are compiled and made available. The TLB is also responsible for compiling background information on the site such as current and historical practices and maintenance of the site hazard plan or where available, the LCF.

3.9 An LQA must only be undertaken by competent specialists, be they the in-house MOD specialists identified at Annex A or independent external vetted specialists from commercial consultancies under their management. All LQAs commissioned through commercial consultants must be reviewed and signed off by a SQEP with demonstrable professional experience and qualifications/accreditations.

3.10 When procuring work, whether via DIO EPS EOLM or an alternative procurement route it is good practice to issue a clear Statement of Requirement (SOR) to prospective suppliers outlining the requirements specific to the site. Details to be included within an SOR are detailed in Box 3.3. An SOR should be included alongside MOD policy guidance such as this document and the LQA Directive within the commissioning paperwork. Examples of SORs are available from DIO EPS EOLM, or alternatively DIO EPS EOLM can support by providing one.

Box 3.3 Statement of Requirement (SOR)	E)
The Statement of Requirement sets out clearly and succinctly:	
 background to and purpose of the LQA; 	
site location;	
 project objectives; 	
 constraints and considerations such as security; 	
 standards that apply; 	
 deliverables/reporting requirements; 	
deadlines;	
 proposed use for the site, i.e. continued military use or redevelopment to resid 	dential; and
 requirements for collateral warranties etc 	

Quality Assurance and Sign-off

3.11 All LQA reports commissioned through commercial consultants should be prepared, reviewed and authorised by Suitably Qualified and Experienced Persons (SQEP) with the necessary skills, knowledge and experience required for their particular role. Demonstrable evidence of competencies of key project personnel (including primary author, reviewer and authoriser) should be provided by the Consultant prior to the commission of their services.

3.12 For report authorisers, appropriate professional qualifications should be held by the individual, which should comprise Chartership with a relevant professional body as a minimum, together with additional qualifications relevant to the work being completed. Such qualifications may include accreditation with the Specialist in Land Condition Register (SiLC; see Box 3.4) or an equivalent.

Box 3.4 Specialist In Land Condition (SiLC)



What is a SiLC?

'A registered Specialist In Land Condition (SiLC) is a senior practitioner/professional able to demonstrate a broad awareness, knowledge and understanding of land condition issues, who can give impartial and professional advice in their field of expertise'.

Initially developed to support the use of Land Condition Records (LCR), SiLC is recognised as a much broader registration. The qualification is appropriate for experienced individuals involved in the assessment and management of land condition/contamination.

The registration is supported by a number of professional bodies including IEMA and CIWEM.

SiLC Vision Statement

To develop and maintain a high quality unifying qualification for assessment and remediation of Brownfield sites which fulfils the needs of public and private sectors and society as a whole.

Further Details are available from: www.SiLC.org.uk

3.13 For further advice on the LQA process or whether a site should be included on the MOD Estatewide LQA Programme contact DIO EPS EOLM - Contact details are enclosed in Annex A.

Ordnance, Munitions and Explosives

3.14 Ordnance, Munitions and Explosives should be considered where appropriate, in two parts:

a. Health and safety risk posed by kinetic effects in the event of UXO and munitions are triggered; and

b. Health and environmental risk posed by the leaching or deposition of explosives residues and metals from expended ordnance and munitions.

3.15 A UXO Preliminary Risk Assessment (UXOPRA) in accordance with CIRIA C681 addresses 3.10a. The UXOPRA will normally be addressed by DIO Explosive Ordnance Clearance Specialists, who can be contacted to advise on the potential presence of UXO, and the risk that UXO may present to those conducting LQA activities.

3.16 The LQA will cover the environmental risk of land contamination on a site from residues due to known firing or disposal of OME, as described in 3.10b.

3.17 Where appropriate, a UXOPRA can be appended to the LQA, as supporting evidence, given the LQA does not specifically assess the likelihood of encountering UXO nor the risk associated with this.

3.18 In the UK the explosive threat from UXO is primarily treated as a health and safety at work and a public safety issue. Following consultation with the EA, MOD's position statement on the assessment of significant contaminant linkages in relation to UXO is presented in Box 3.5.

Box 3.5 UXO Position Statement

In the UK the explosive threat from UXO is primarily treated as a health and safety at work and a public safety issue.

Defence Sites/MOD controlled Property:

Defence sites are subject to clearance operations to ensure they are safe and suitable for use. Clearance operations are conducted by trained personnel using the appropriate in-service equipment and in accordance with the accepted operating procedures at the time. Statements given following any of these operations cannot provide a 100% guarantee that all items have been recovered.

However, if ordnance is left undisturbed, it will under normal circumstances not pose an explosion threat. The accepted procedure is that if a suspicious object is found, the finder should contact either the Local Range Officer or Police who will contact one of the Service Explosive Ordnance Disposal (EOD) Teams. The Service EOD Teams will then assess the risk and deal with the immediate problem under Military Aid to the Civil Powers arrangements. They will also make an assessment on the need for further investigation/clearance work. Under normal circumstances UXO is not considered to pose a significant possibility of significant harm with regard to explosion at defence sites.

Alienated Sites/Former MOD Property:

MOD treats the explosive threat from UXOs as a health and safety issue.

If ordnance is left undisturbed, it will under normal circumstances not pose an explosion threat. The accepted procedure is that if a suspicious object is found, the finder should contact the Police who will contact one of the Service EOD Teams. The Service EOD Teams will then assess the risk and deal with the immediate problem under Military Aid to the Civil Powers arrangements. They will also make an assessment on the need for further investigation/clearance work. If there is the possibility that ordnance may be disturbed, MOD believes that it is usually possible to put in place suitable and sufficient risk management measures, including the provision of information to potentially affected parties on accepted procedures, to prevent significant harm from occurring.

3.19 Whilst there are no UK specific generic assessment criteria for explosives residues, values have been developed by BAe Systems (formally Royal Ordnance) and other organisations. However, these are to be used with care as they may not be applicable to the UK situation and are not necessarily compliant with UK policy and guidance.

Box 3.6 Explosives Residues

Si	gn	Pos	t	
	and and a			

- Environment Agency, Collation of toxicological data and development of guideline values for explosive substances, P5-036/01, 2002.
- Rudland, D J et al. Contaminated Land Risk Assessment, A Guide to Good Practice. CIRIA C552, London 2002.
- Royal Ordnance. Explosive Materials Determination of Toxicological Hazards and other Properties, 1990.
- LQM/CIEH Suitable 4 Use Levels for Human Health Risk Assessment, 2015

4. STAGE 1 - RISK ASSESSMENT

4.1 The process is tiered as shown in Figure 2 and starts with the identification and assessment of the potential site and situation specific hazards (contamination sources) culminating in a Preliminary Conceptual Site Model (CSM) and Preliminary (Qualitative) Risk Assessment (PRA) in the Phase 1 LQA which establish the potentially unacceptable risks. In doing so the potential sources, receptors and likely pollutant linkages are identified.

4.2 In the Phase 2 LQA the potentially unacceptable risks are estimated and further evaluated by testing and refining the Preliminary CSM using site specific data such as laboratory derived concentrations of contaminants in the soil. Risk estimation is concerned with assessing the likely magnitude and probability of harm that may result from an identified hazard (contaminant source) and which receptors will or are likely to be affected. Risk evaluation on the other hand is about deducing whether the risk is or has the potential to become unacceptable i.e. the focus is on identifying the 'significant contaminant linkage'.

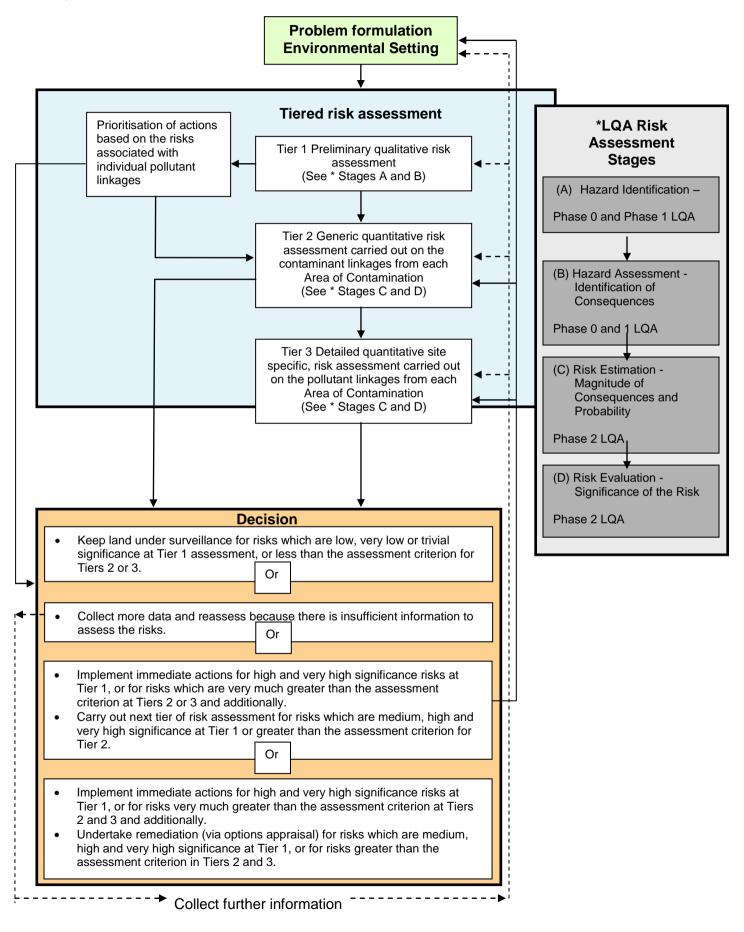
4.3 Hence, there are 2 parts to this stage:

Part 1A Hazard Identification and Assessment (Phase 0 Preliminary Hazard Assessment and Phase 1 Desk Study); and

Part1B Risk Estimation and Evaluation (Phase 2 Site Investigation)

Figure 2 Summary of Tiered Risk Assessment Process

Adapted from the NDA Direct Research Portfolio: Draft Practitioners' Guide TSG (10)0664 2010



4.4 Figure 3 shows the process starting with the Phase 0 LQA, progressing to the Phase 1 LQA and if required the Phase 2 LQA, onto other subsequent phases (Phase 3 and 4) should these be necessary.

4.5 The following sections and paragraphs are not intended to provide a definitive guide to risk assessment within the context of the LQA process, but outline best practice and identify current guidance. Risk assessments should not be undertaken in isolation and specialist support and advice should be sought. Appropriate contacts are listed in Annex A.

Box 4.1 Risk Assessment Good Practice Guidance

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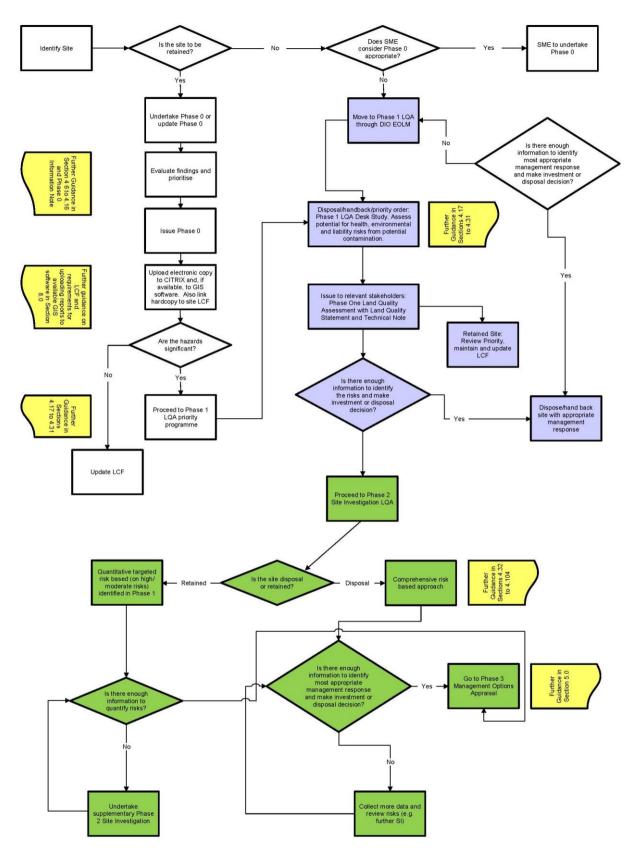
A useful overview of the Risk Assessment Process is provided by:

•Rudland, DJ et al. *Contaminated Land Risk Assessment, A Guide to Good Practice*. CIRIA C552, London 2002.

•R&D 66 Guidance on the Safe Development of Housing on Land Affected by Contamination, 2008 Volume 1 and 2. NHBC, CIEH and EA

•Green Leaves III Guidelines for Environmental Risk Assessment and Management. PB13670. November 2011

Figure 3 LQA Process Flow Chart Phase 0 to Phase 2



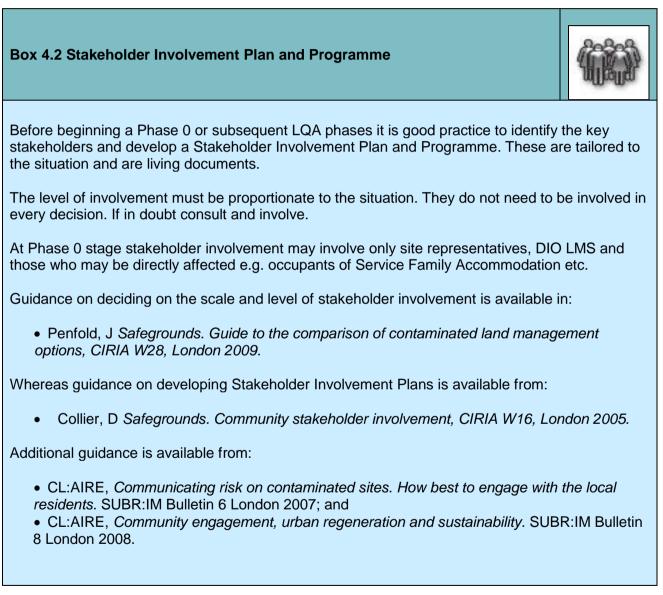
PART 1A HAZARD IDENTIFICATION AND ASSESSEMENT

PHASE 0 LQA – PRELIMINARY HAZARD ASSESSMENT

Introduction

4.6 As resources are finite it is essential that MOD identifies and targets the highest risk priorities i.e. where there is an immediate significant risk of significant harm or pollution or breach of legislation, carrying out a phased approach. The first step is completion of a Phase 0 LQA – Preliminary Hazard Assessment for MOD establishments and land holdings to identify potential environmental and health hazards to formulate a prioritised Phase 1 LQA Programme.

4.7 This appraisal and prioritisation exercise will involve a stakeholder plan/programme, refer to Box4.2.



4.8 A Phase 0 LQA provides a summary of the likely land quality based on readily available desk based information by providing an initial review of potential contaminants and receptor sensitivities. Although the Phase 0 LQA does not consider pollutant linkages directly, the receptor sensitivity review and assessment considers the potential for a pathway to be present, thus identifying which sites are most likely to fall under the statutory definition of contaminated land.

Reporting

4.9 The Phase 0 LQA is an unrestricted document that carries an 'Official' marking, and as such will contain predominantly factual information together with the Outline CSM and Category/Priority Assessment of the overall land quality and suitability for redevelopment that sets the presence of any land contamination and pollution for controlled waters in context.

4.10 The Phase 0 LQA is usually reported using a standard 2 to 3 page template presented with an overview of the key findings and recommendations. Full details of the Generic MOD Phase 0 LQA Reporting Format and content is provided in Annex B1 with Section 6 of Annex B4 setting out the requirements for GIS compatible electronic copies. Please note that Annex B represents the standard default formats which are designed for retained and disposal sites alike, but may need to be tailored to the situation. For instance the requirements for a LQA undertaken in support of a planning application or voluntary inspection under Part 2A of the Environmental Protection Act 1990 can differ from the default format. Advice must therefore be sought from the relevant authority and key stakeholders at the earliest opportunity to avoid unnecessary expenditure and delay. In the case of disposals, retained sites, voluntary inspections and planning applications the technical authority is DIO and the relevant contacts for advice and guidance are detailed in Annex A.

Methodology

4.11 There are a number of published methodologies for accomplishing this, most notably the CLR6 Appendix 1 "Prioritisation and categorisation procedure for sites which may be contaminated".

4.12 DIO has developed the Phase 0 approach, 'Strategic Land Quality Appraisal and Prioritisation Methodology' taking account of CLR6 and the Source-Pathway-Receptor concept, also known as the pollutant linkage model. This approach provides a systematic and auditable methodology enabling sites to be screened and prioritised with production of a Phase 0 LQA Report.

4.13 The Phase 0 LQA Report provides a summary of the likely land quality based on the readily available information together with an initial identification of potential hazards and the likely risks. As such it allows the relative significance of the site in terms of the potential for significant harm to be determined together with the need or otherwise for further assessment. In this way it enables a prioritised programme of Phase 1 LQAs to be developed and assists in identifying the immediate and longer term need for institutional controls to mitigate potential risks. As such it allows DIO to take a holistic view of the land quality across their sites and develop a coherent, defensible and prioritised management programme that will target resources where there is greatest need. This will minimise the risk of both regulatory action and nugatory work and allow effective budgeting.

4.14 Whilst the methodology has been developed to assist the MOD in identifying hazards, risks and liabilities with respect to land contamination on the retained estate it is insufficiently detailed to allow a determination of a site's status under the Part 2A Contaminated Land Regime. It does however, enable MOD to identify those sites which are most likely to fall within the statutory definition of 'contaminated land' and will assist local authorities in discharging their statutory obligations in terms of inspecting the land in their area and minimise the risk of sites being inappropriately designated as 'Contaminated Land'.

4.15 The detailed methodology is available as a standalone document from DIO EPS EOLM who is also able to advise you on its application or else undertake the assessment and prioritisation for you.

4.16 Subsequent phases of LQA will be required if the potential for SPOSH is identified. Figure 3 provides a simple guide through the Phase 0 process.

PHASE 1 LQA – DESK STUDY AND SITE RECONNAISANCE

Introduction

4.17 Where the Phase 0 appraisal and prioritisation identifies the need for further assessment or it is considered that notable risks exist on-site without the need for a Phase 0 LQA, the next step is the Phase 1 LQA which is to be undertaken as part of a prioritised programme or as instructed by DIO / MOD. This phase involves a site reconnaissance visit, interviews with key staff and a more detailed review of factual data concerning the site history, geology, hydrology, hydrogeology, regulatory issues, planning and site operations etc to validate and refine the findings of the Phase 0 LQA or expand upon any existing information held on file for the site. This is done via the creation of a Preliminary CSM and the undertaking of a PRA to better establish the associated potential health, environmental and liability risks from the identified sources. In conclusion the Phase 1 will identify the need for and scope of any further work be it additional desk based research, intrusive investigations which could inform subsequent Phase 2 LQA works or the use of institutional controls to manage and mitigate the risk.

Box 4.4 Phase 1 LQA Top Tips



Always treat each site and therefore each Phase 1 LQA as unique as there will be combinations of factors that are specific to each one.

Establish clear, relevant objectives that reflect the site situation and context such as whether the LQA is in aid of a disposal or voluntary inspection, as this will avoid confusion, misunderstanding and potentially inappropriate conclusions/recommendations.

Always include a 6 figure National Grid Reference as MOD sites often have several names reflecting the fact that they have been occupied by different Services and undergone changes in use. Where appropriate the relevant Spec 005 codes should also be considered.

Consult DIO EOLM.

Scope the Phase 1 LQA appropriately. Under or inappropriately scoped Phase 1 LQAs cost time and money.

All assumptions, caveats and limitations should be clearly stated so there are no misunderstandings.

Seek advice from MOD SMEs as soon as possible.

Reporting

4.18 The Phase 1 LQA will normally contain three elements:

- LQS;
- Land Quality Assessment Report; and
- Technical Note.

4.19 The LQS is a 2 to 3 page document which takes the place of the Executive Summary within the LQA report. It provides a non-technical summary of the land quality based on the available information and site reconnaissance and includes the potential risks to human health and the environment, including controlled waters (groundwater and surface water). It also identifies the available historical records, details of known environmental pollution and previous investigations and remediation that have taken place and provides an indication of the suitability for use (current/redevelopment). The

LQS must be written in such a way that it can be used as a freestanding document. It should not reference specific sections of the LQA report or figures and must not contain any recommendations.

4.20 The LQA Report is an unrestricted document that carries an 'Official' marking, and as such will contain predominantly factual information together with the Preliminary CSM, PRA, an assessment of the overall land quality and suitability for use (current/redevelopment) that sets the presence of any land contamination and pollution of controlled waters in context i.e. is it localised, limited in extent and confined to shallow soil horizons etc.

4.21 The Technical Note is a restricted document that carries an 'Official Sensitive-Commercial' protective marking. The note sets the conclusions of the LQA report into a regulatory context, providing an assessment of the liabilities associated with any known or potential contamination including a view on whether the site is likely to be determined 'Contaminated Land', an evaluation of the management options, cost estimates for each option and a defensible recommendation that is consistent with current best practice and affords Value For Money (VFM). The management options and costs necessary to address the immediate and longer term risks and liabilities may include, but not limited to Phase 2 LQA (site investigation or monitoring). For each option, including the recommended option uncertainty and justification should be provided.

4.22 Full details of the Generic MOD Phase 1 LQA Reporting Format and content is provided in Annex B2 respectively with explanatory notes detailed in Annex B4. Please note that Annex B represents the standard default formats which are designed for retained and disposal sites alike, but may need to be tailored to the situation. For instance the requirements for a LQA undertaken in support of a planning application or voluntary inspection under Part 2A of the Environmental Protection Act 1990 can differ from the default format. Advice must therefore be sought from the relevant authority and key stakeholders at the earliest opportunity to avoid unnecessary expenditure and delay. In the case of disposals, retained sites, voluntary inspections and planning applications the technical authority is DIO and the relevant contacts for advice and guidance are detailed in Annex A.

Box 4.5 LQA Content Guidance – Phase 1



Guidance on the content of Phase 1 LQAs depending upon the situation and purpose are is provided by:

• Guidance for the Safe Development of Housing on Land Affected by Contamination. NHBC and EA R&D 66, 2008.

• CLR2. Guidance on preliminary site inspection of contaminated land. Report by AERC Ltd. DoE 1994.

• CLR11. Model Procedures for the Management of Land Contamination. EA and DEFRA 2004.

• EA Technical Report P5-042/TR/01. Land Contamination: Technical Guidance on Special Sites: MOD Land.

• CIRIA C552. Contaminated Land Risk Assessment. A Guide to Good Practice. Report by CIRIA. January 2001.

Methodology

4.23 Current best practice is set out in CLR11 and useful guidance on the scope, approach and content including the Preliminary CSM and the PRA is available in R&D 66, CIRIA C552 and generic technical guidance on conceptual models can also be found in the EAs R&D publication NC/99/38/2 (refer Appendix 1). Additional guidance is provided in Annexes D and E respectively to this guide.

4.24 For MOD sites the Phase 1 LQA will include a site reconnaissance / walkover visit which must be tailored and involve as a minimum, an appraisal detailed below. It is essential that the appropriate stakeholders are consulted and involved to ensure the scope is fit for purpose:

- Site infrastructure, drainage and services;
- Site operating procedures both past and present; and
- Potential environmental issues/incidents requiring attention and / or investigation.

Box 4.6 Stakeholder Involvement



At the Phase 1 stage the level of stakeholder involvement may not vary from that of the Phase 0.

However, depending upon the situation there may be a need to involved the regulators either directly in the planning and scoping of the LQA or else by keeping them informed of what is intended and why.

The Stakeholder Involvement Plan and Programme should be amended to reflect any changes.

4.25 Where the site reconnaissance involves the inspection of fuel infrastructure and entering confined spaces etc then full adherence to the relevant health and safety legislation and regulations is required together with the appropriate MOD Safety Rules and Procedures (SRPs). The latter are managed and enforced by DIO through the Senior Authorising Authorities located within the DIO SEE Team in conjunction with DIO / MOD business partners and facilities management personnel. Contact details are presented in Annex A.

4.26 The review and assessment of factual data should involve the collation and appraisal of all available desk based information relating to the site that is deemed to be appropriate together with anecdotal evidence collected from site staff. This data combined with the site reconnaissance should be used to refine the Outline CSM (from Phase 0, where undertaken) into the Preliminary CSM where all the potential pollutant linkages are identified and qualitatively assessed, with the objective of establishing the potential health, environmental, infrastructure and liability risks, likelihood of pollutant linkage, potential consequence of the pollutant linkage and likely significance. The outcome is the PRA which should be presented both as a summary narrative and as a summary table, refer Annex E.

Immediate Management Action

4.27 If the PRA or the site reconnaissance identifies immediate risks then action must be taken to mitigate and manage those risks in advance of any Phase 2 LQA work.

4.28 The objectives should be to protect human health and the environment by:

• Stopping the situation getting worse by preventing the spread of contamination or further pollution of controlled waters;

• Controlling exposure through the use of access restrictions etc; and

• Implementing monitoring regimes if required pending the outcome of further assessment and a decision on the long term management approach

4.29 The nature of the risk will dictate the need for and level of regulator involvement, MOD policy is quite clear: the regulators are to be made aware of pollution incidents etc immediately. Whilst it is good practice to develop an Immediate or Interim Management Plan in situations such as this, this should not be at the expense of delaying the necessary action of contacting the regulator.

PART 1B RISK ESTIMATION AND EVALUATION

PHASE 2 LQA – SITE INVESTIGATION

Introduction

4.30 The findings and evaluation of the Phase 1 LQA Desk Study will determine the need and scope for a Phase 2 LQA intrusive and/or non-intrusive site investigation using geophysical and other such techniques.

4.31 The aim is to provide a reliable assessment of land quality and in doing so confirm the presence and quantify the nature and extent of contamination setting out the level of uncertainty (inherent in the sampling, sampling preparation and analysis), assess the significance in terms of the risks and associated environmental, health and financial liability, provide an appraisal of the management options together with a reliable order of cost estimate (OCE) and make recommendations on how to manage the risk and liability cost effectively. The options looked at will include, but not be limited to: institutional controls / management procedures, remediation and further investigation and/or monitoring. These may be tackled in a single stage or in a number of targeted phases that may be spread over a number of months or financial years.

4.32 Where the Phase 2 LQA is being undertaken for disposal purposes (divestment) then if known consideration must be given to the potential future land uses, the associated risks and costs of making the land suitable for use and potential for contamination to be caused as a result of demolition.



Reporting

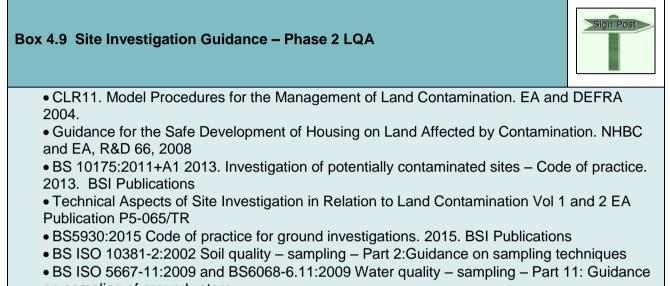
4.33 The Generic MOD Phase 2 LQA Reporting Format is set out in Annex B3 with explanatory notes detailed in Annex B4. As with Phase 1 LQAs the generic format represents the standard default designed for retained and disposal sites alike, but may need to be tailored to the situation. The requirements for a LQA undertaken in support of a planning application or voluntary inspection under Part 2A of the Environmental Protection Act 1990 can differ from the default format. Advice must be sought from the relevant authority and key stakeholders at the earliest opportunity to avoid unnecessary expenditure and delay. In the case of disposals, retained sites, voluntary inspections and planning applications the technical authority is DIO and the relevant contacts for advice and guidance are detailed in Annex A. Any site or project specific requirements should be set out in the commissioning paperwork and SOR as defined in Box 3.3.

Methodology

4.34 The Phase 2 LQA is intended to estimate and evaluate the potential risks identified by the Phase 1 LQA through testing and refining the Preliminary CSM using site specific data such as laboratory derived concentrations of contaminants in the soil.

4.35 The key steps are:

- Define / set clear objectives;
- Develop an investigation strategy;
- Scope the investigation
- Sampling, field testing / monitoring and analysis;
- Quantitative Risk Assessment;
- Evaluation and Conclusions; and
- Reporting.



- on sampling of groundwaters
- NHBC standards, 2016

Objectives

4.36 The objectives must be appropriate and achievable and the site investigation must reflect both the objectives and adequately test the Preliminary CSM thereby enabling it to be refined and a robust Quantitative Risk Assessment undertaken. The latter is key to identifying the associated risks and liabilities together with the measures necessary to manage and mitigate them. To achieve this, the investigation may if required comprise more than one phase with each subsequent phase being informed by the preceding ones.

Planning

4.37 When planning the site investigation care must be taken to minimise the environmental impact and risk of making the situation worse through, for instance, cross contamination and short circuiting contamination with a borehole and creating a preferential pathway to an aquifer. To this end it is good practice to develop a Sampling Strategy supported by a Sampling and Analysis Plan as described in CLR 4. This sets out the locations of sampling points, the spacing/density, describes what they are be they boreholes, trial pits or window sample locations etc. and explains why samples are being collected in that location and by that method. As part of this the plan should explain where and why targeted and/or non-targeted sampling is to be undertaken. The plan should also set out the sampling protocols and any gas, groundwater or surface water monitoring together with an estimate of the degree of uncertainty. Guidance is provided by BS 10175:2011+A1:2013.

Box 4.10 Sampling Density and Uncertainty



The number and density of samples i.e. the spacing is dictated by the Preliminary CSM and the LQA objective. There is no set rule of thumb.

Even so, regardless of how many sampling points there are, the number of samples, number and range of chemical analyses there will always be a level of uncertainty inherent in any site investigation.

The site investigation design must be appropriate, proportional and keep uncertainty to a minimum. It is useful to quantify and document the level of uncertainty so that this can be factored into the risk assessment and overall evaluation of the sampling data.

Guidance on this is provided by: R&D 66 and BS 10175:2011+A1:2013

Sampling and Analysis

4.38 Soil, water and soil gas/vapour samples must be representative and collected and stored in such a way as to avoid cross contamination or compromising the sample integrity. The collection of soil samples is to be undertaken in accordance with BS10381. In the case of surface and groundwater samples these must be sampled in accordance with BS6068/BS ISO 5667. For the collection of soil gas and vapours, references for their collection are detailed in Box 4.11.

4.39 All collected samples should be both stored and preserved in accordance with the requirements specified by the laboratory undertaking the chemical analysis.

Box 4.11 Guidance on the collection and assessment of soil gas and vapours (volatiles)	Į
 CIRIA C665 Assessing risks posed by hazardous ground gases to buildings, 2007 BSI BS8485 Code of practice for the characterisation and remediation of ground gas in Brownfield developments, 2007 CIRIA C682 The VOCs Handbook. Investigating, assessing and managing risks from inhalation of VOCs at land affected by contamination, 2009 NHBC Guidance on Evaluation of Development Proposals on sites where methane and carbon dioxide are present, 2007 BS 10175:2011+A1:2013. <i>Investigation of potentially contaminated sites</i> – Code of practice. 2011. BSI Publications BS8576:2013 Guidance on investigations for ground gas – Permanent gases and Volatile Organic Compounds (VOCs) 	

4.40 An appropriate Quality Assurance Regime should be put in place involving the use of blank samples including trip and equipment blanks together with duplicate samples to provide a check on the accuracy and precision of the sampling and analyses at a suitably justified ratio based upon the scope of the works proposed.

4.41 All hand/trial pits boreholes and window sample holes must be logged in accordance with BS5930:2015 ensuring odours and visual evidence of contamination and water ingress are recorded.

4.42 As a minimum the chemical analysis of samples must be carried out by UKAS accredited laboratories. Where possible the analyses should be undertaken using the EAs Monitoring Certification Scheme (MCERTS) as this provides the greatest assurance of both quality and reliability.

Health and Safety

4.43 Health and safety are paramount when designing and undertaking a site investigation. Fundamental requirements include: adequate health and safety risk assessment and the employment of suitable and sufficient safe systems of work. In addition all MOD regulations and instructions concerning safe working procedures shall be rigorously adhered to, and advice should be sought from a health and safety SQEP.

4.44 To inform the health and safety risk assessments a Statement of Known Hazards (SoKH) is to be requested from site in advance of all penetrative ground investigation works. This should detail known hazards that may affect the proposed works on site including (but not limited to) service locations, site activities and potential contamination. In addition to this, existing available information, such as previous LQAs should be used where available.

4.45 Prior to works commencing, all health and safety paperwork including method statements and risk assessments should be undertaken, finalised and provided to the MOD / DIO representatives in good time in order that there is sufficient time for them to provide comment prior to the start of any works on-site. These documents should be dynamic, and updated throughout the LQA works, as and when site conditions change and/or hazards identified.

4.46 Construction (Design and Management) Regulations 2015 (CDM 2015) apply to all construction works on site, which for LQAs applies to Phases 2, 3 and 4. On commissioning of the project, roles of the contractor and designer are to be assigned. There will be a requirement for the production of a health and safety file, to be retained on record on site. A template of the required health and safety file is included in Annex B5 and should be supplied to the MOD /DIO upon completion of the project.

4.47 Should the project exceed 30 days, 500 person days and have more than 20 workers on the project at any one time, then the Principal Contractor will need to notify the Health and Safety Executive (HSE), and will undertake HSE notification via form F10 on behalf of the MOD / DIO.

Waste Management

4.48 It is essential that waste arising from intrusive investigations is minimised and managed appropriately. Guidance on sustainable waste management is provided at Annex C.

Risk Estimation

4.49 This is concerned with assessing the likely magnitude and probability of harm that may result from an identified hazard (contaminant source) and which receptors will or are likely to be affected. It therefore involves the refining of the Preliminary CSM and moves from qualitative to quantitative risk assessment.

Updated CSM

4.50 The Preliminary CSM must be updated once all the chemical analyses and site specific data has been derived to confirm or exclude/discount the potential pollutant linkages identified by the Phase 1 LQA. Only then should the process of risk estimation by means of Quantitative Risk Assessment (QRA) begin. Guidance on the presentation and construction of the CSM is provided at Annex D.

Quantitative Risk Assessment – Tier 2 and 3

4.51 There are two types of QRA: Tier 2 - Generic QRA (GQRA) and Tier 3 - Detailed QRA (DQRA). Ordinarily you should start with a GQRA whereby the determined contaminant concentrations are compared to appropriate and justifiable Generic Assessment Criteria (GAC) such as the S4ULs, C4SLs, UK CLEA Soil Guideline Values (SGVs), UK Drinking Water Standards (DWS's) and

Environmental Quality Standards (EQS) before potentially moving to DQRA. As GACs take into account a degree of uncertainty they are inherently conservative and as such an exceedance should be considered an indication that further works may be required. Where measured concentrations of contaminants fall below them it can be concluded that there is no significant risk providing the guidelines have been applied correctly. It is therefore essential to understand the limitations of the application of the various GACs, in particular the exposure scenarios to which they relate. In the case of the CLEA SGVs they are not suitable for situations where the CSM does not match the CLEA exposure scenarios.

4.52 The 2012 revision to the statutory guidance for England and Wales introduced a framework comprising four land/ human health risk categories with which to determine whether non-radioactive land contamination presents a significant possibility of significant harm.

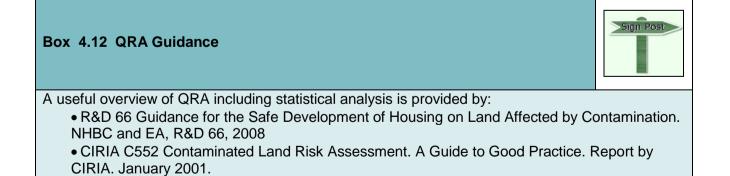
- Category 1 : Human Health: unacceptably high probability that significant harm will occur if no action is taken;
- Category 2: Human health : the risks posed by contamination are sufficient for the land to be deemed to meet the legal test for posing a significant possibility of significant harm;
- Category 3: Human health: whilst the risks posed by contamination may not be low, the legal test with respect to the land posing a significant possibility of harm is not met; and
 - Category 4: Human Health: there is considered to be negligible risk to human health.

4.53 Category 1 and 2 will constitute statutory contaminated land whereas Category 3 and 4 will not.

4.54 The level of conservatism within for example the 2009 SGVs or the more recent C4SLs is such that they would be considered or in the case of the C4SLs have been developed, to fall within Category 4 i.e. they represent concentrations of contaminants below which the risk to human health is negligible and the land is 'very unlikely to pose a significant risk of significant harm'.

4.55 Where GACs are either not available or deemed inappropriate or for that matter there is sufficient knowledge to indicate that GACs will be exceeded it is good practice to move immediately to a DQRA to derive, in the case of human health, Site Specific Assessment Criteria (SSAC) from published toxicity and exposure data or in the case of controlled waters, to derive remedial target concentrations. The aim is to identify those pollutant linkages that are significant in terms of posing an unacceptable level of risk. These are often referred to as Relevant Pollutant Linkages (RPL).

4.56 Guidance on the use of GACs and other tools for risk assessment is provided in DEFRA Guidance 04/12 and CL:AIRE bulletin CSB 10 dated March 2012.



4.57 Guidance on the presentation of the risk assessment is provided at Annex E.

GQRA - Human Health

4.58 There are various sets of screening criteria that can be applied within a human health risk assessment, which will have been derived using the EA CLEA model or similar, either using the existing data set for standard land use scenarios or manipulation to create new land uses. Examples

of published sets of screening criteria include the recently published CIEH/LQM S4ULs and C4SLs. Some guidance on the key differences between some of the published UK GAC sets is provided in CL:AIRE bulletin CSB 10, although this does not include S4ULs and C4SLs but the previous edition of CIEH/LQM GACs.

4.59 Any GACs selected for use within a Phase 2 LQA must be appropriate and justified within the settings of the works undertaken and must take in to consideration both known current and potential future end uses.

4.60 It is worth noting that when screening criteria are updated and replaced, potential exceedances or risks highlighted within historical reports may require comparison to more recent GACs in order to assess whether the original findings are still relevant in the current regulatory setting. For example, the older LQAs commonly refer to the 'Dutch Values' which do not typically fit with the current UK legislative and policy regime.

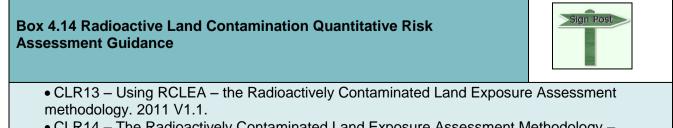
4.61 Justification should also be provided for any nonstandard, non-published input parameters, e.g. soil type and assumptions made within the human health risk assessment particularly when deriving site specific values.

4.62 Care needs to be taken when assessing the risk posed by Petroleum Hydrocarbons. Where the GAC exceeds the theoretical soil saturation limit the EA recommend that modelling other than 3-phase partitioning should be used to derive SSAC/screening values or else the saturation limits should be used. However, reliable theoretical saturation limits are difficult to derive owing to the multiplicity of variables and the model assumptions for vapour formation break down where free phase is or is likely to be present. This means that the model will over predict how much vapour can be formed and the calculated screening values will, as a result, be overly conservative potentially resulting in unnecessary remediation. In view of this and the fact that the prediction of theoretical saturation limits using a model is not sufficiently accurate the only way of being sure free phase is present is to look at the soil and to use gas standpipes to assess the presence and nature of any associated vapour hazard.

Box 4.13 Petroleum Hydrocarbons Quantitative Risk Assessment Guidance	Sign Post
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• Environment Agency Science Report P5-080/TR3: UK approach for evaluating the human health risks from petroleum hydrocarbons in soils 2005.

4.63 The preferred EA/DEFRA approach to GQRA of radioactive contaminants is to us the Radioactively Contaminated Land Exposure Assessment Methodology (RCLEA) to generate GACs known as RSGVs (Radioactivity in Soil Guideline Values). RCLEA is aligned to Part 2A and as such compliments the CLEA Model. However, there are other approaches and early consultation with the regulator is essential.



• CLR14 – The Radioactively Contaminated Land Exposure Assessment Methodology – Technical report 2011 V1.2

- CLR15 The RCLEA Software application. 2011 V1.3
- NRPB Vol 9, No2 Radiological Protection Objectives for Land Contaminated with Radionuclides. 1998.
- NRPB Methodology for estimating the Doses to members of the Public from the Future Use of Land Previously Contaminated with Radioactivity.
- Guidance on Characterisation and Remediation of Radioactively Contaminated Land. EA 2002.
- Smith G. Safegrounds. Review and commentary on site end-points and radioactively contaminated land management. CIRIA W20. 2005.

GQRA – Controlled Waters

4.64 For initial assessment purposed at a GQRA level, test results (which may include leaching testing and/or direct analysis of water; whichever technique(s) is most appropriate and justifiable) should be compared to water quality standards. These may comprise DWSs as an initial screen, or published EQS where available; taking care to ensure any parameter that can affect the EQS such as pH or water hardness etc is included in the Sampling and Analysis Plan. As EQS were derived primarily to assess surface water bodies care should be taken in the application of these GAC to groundwater. Alternatively Water Screening Values (WSV) can be derived using commercial software. For example, Atkins has derived commercially available WSVs for groundwater containing VOCs, specifically to address the risk posed by vapour inhalation.

4.65 International GACs may be used at this stage but as with soil GACs their use must be justified as they may not be applicable to the UK situation.

DQRA – Human Health

4.66 Following identification of exceedances of GACs at the GQRA stage, the next step is to use a DQRA whereby site specific screening criteria are derived. It should be noted whilst exceeding a SGV or other GAC does not itself constitute an unacceptable risk to human health or for that matter a SPOSH, it is advisable to assess further, where appropriate.

4.67 Care must be taken to ensure the SSACs are robust. Ideally the DQRA should incorporate the relevant bioavailability and bio-accessibility data. However, there is much uncertainty inherent in the available bio-accessibility data.

DQRA – Controlled Waters

4.68 Where the GQRA findings indicate a potentially significant risk to Controlled Waters, further assessment should be undertaken to refine understanding of the likely contaminant migration mechanisms. This should form part of a DQRA, whereby site-specific information (which may for example include laboratory test results, permeability assessments, source zone delineation etc.) may be combined with literature values (which may for example include contaminant physico-chemical parameters etc.) and utilised in bespoke modelling software to model the fate and transport of the key chemicals of concern. Commercially available modelling packages include the EA's Remedial Targets Methodology (RTM) worksheets and ConSim, and other software packages may be utilised where fully justified and agreed prior to use. The use of modelling software is to understand whether the contaminant concentrations identified by the LQA works pose an unacceptable risk to the receptor of concern, and thereby whether any remedial action is required to mitigate the identified risks. Modelling tools such as the RTM or ConSim allow the user to derive site-specific remedial targets or Clean-Up Goals (CUGs), which can be used during any necessary remediation for validation purposes.

4.69 As part of a Controlled Waters DQRA, an assessment of the sensitivity of the key parameters utilised within the model should typically be undertaken via sensitivity analysis. The parameters included within the sensitivity analysis, and the variances in the values applied from the initial

modelling, should be fully justified by the Consultant. Copies of all model output sheets (including sensitivity analysis) should be included within the report appendices.

GQRA and DQRA – Vapours and Gas

4.70 The risk posed by bulk ground/soil gas should be assessed by means of deriving Gas Screening Values (GSV) and establishing the risk and need or otherwise for mitigation measures.

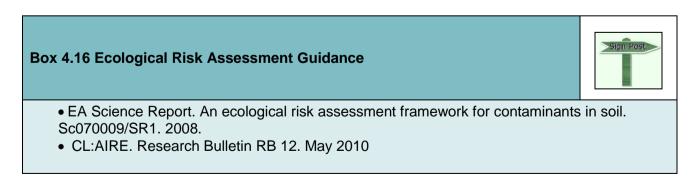
4.71 Soil vapour assessment for volatile organic compounds should include development of soil vapour screening values, drawing upon site-specific information such as soil physical characteristics (e.g. particle size distribution) and where appropriate, consider development specific information (e.g. building dimensions and construction).

Box 4.15 Vapour and Gas Risk Assessment Guidance CIRIA, Investigation & Assessment of Volatiles at Brownfield Sites 2008 CIRIA, C665, Assessing risks posed by hazardous ground gases to building, 2007 CIRIA, C659, Assessing risks posed by hazardous ground gases in buildings, 2006 NHBC, Guidance on Evaluation of Development Proposals on sites where methane and carbon dioxide are present, 2007 Wilson et al, Ground Gas Handbook 2008. The Local Authority Guide to Ground Gas 2008 British Standards Institution (2013): BS 8576 Guidance on investigations for ground gas – Permanent gases and Volatile Organic Compounds British Standards Institution (2007): BS8485:2007, Code of Practice for the Characterization and Remediation from Ground Gas in Affected Developments The VOC handbook. Investigation, Assessing and Managing Risks from inhalation of VOCs

GQRA and DQRA – Ecological

4.72 This must consider impact on protected species, areas of natural and ecological importance such as Sites of Special Scientific Interest (SSSIs), trees, hedgerows etc. Care will need to be taken to ensure an appropriate assessment. Merely using EQSs for instance to assess the ecological risk within the aquatic environment may not be enough.

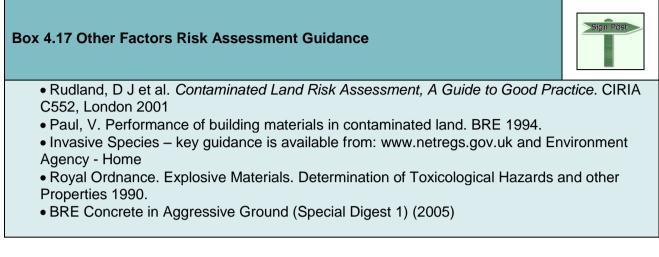
at land affected by Contamination. Baker et.al CIRIA C682, 2009



GQRA and DQRA – Other Factors

4.73 Other factors you may wish to consider include: the potential presence of PFAS/PFOS or their associated compounds, the effect of sulphate on concrete and hydrocarbons on the integrity of plastic pipes, and invasive species such as Japanese Knotweed however, these should not be the sole consideration for any investigation without reasonable justification.

4.74 As part of the risk assessment the potential for future land contamination and pollution of controlled waters occurring as a result of the demolition of existing buildings should be considered unless otherwise specified. This is particularly important for Phase 2 LQAs being prepared in support of site disposal or redevelopment.



Statistical Analysis

4.75 As part of the assessment of the laboratory test data acquired from the site investigation, the use of statistical analysis may be considered by the risk assessor. It should be noted that the use of statistical assessments will not always be appropriate, and it is incumbent on the Consultant to understand where and when the application of statistical tools is both justified and relevant. In accordance with the CL:AIRE Guidance on Comparing Soil Contamination Data with a Critical Concentration, the following points should be considered as part of the dataset review: a) the dataset should be sufficiently robust to represent a meaningful dataset for statistical analysis; b) the dataset should comprise unbiased sample data which are representative of the soil of interest and have been collected using a sampling density appropriate for the area and depth of soil of interest, likely degree of heterogeneity of the soil type and the nature of the risk assessment; and c) adopt the appropriate type of statistical distribution for the dataset being considered (i.e. normal or non-normal distribution). The Consultant should be able to provide justification of the appropriateness of the statistical distribution being adopted, for example through the utilisation of tools such as probability plots or the Shapiro Wilks test (detailed more fully in the appendices to the CL:AIRE guidance document).

4.76 When considering statistical methods, any outliers or anomalous readings within the dataset should be fully investigated prior to undertaking any assessments. Outliers should only be excluded from a dataset when they are obviously and demonstrably the result of an error that can be identified and explained, or clearly indicate that more than one soil population exists within the dataset and can be justified/explained by the conceptual site model.

Risk Evaluation

Acute vs. Chronic

4.77 This is where the need for options to mitigate and manage unacceptable acute (short term) and chronic (long term) health risks as well as significant environmental and safety risks are determined.

4.78 As there are no UK GACs and other guideline values to assess acute risks from exposure to soil contamination it may be appropriate to use a combination of available occupational exposure limits for vapours and dusts or else derive an SSAC using a one-off high soil ingestion rate and maximum concentrations. Alternatively it may be more appropriate to remediate rather than attempt to derive an SSAC.

4.79 Risk evaluation should therefore, be done at the GQRA stage as well as the DQRA stage.

Management Options

4.80 In terms of identifying the management options consideration should be given to: source removal, pathway disruption and receptor protection as well as the cost benefit of undertaking further site investigation, data collection and risk assessment to reduce the level of uncertainty.

4.81 It may be more cost effective and politically expedient to remediate following the GQRA, compared with the cost and time involved in collecting further data to inform the DQRA and reduce the level of uncertainty. Investing in a DQRA can significantly reduce the scope of any remediation or discount the need for it completely. It is essential that all the factors including stakeholder expectations, technical feasibility, political acceptability, budget availability and cost are considered when assessing the need for a risk management response.

4.82 Given the limited budget available for investigative and potential remedial works across the MOD estate, consideration should be given to whether identified risks could be adequately managed on-site without undertaking further works. The 'do nothing' and 'do minimum' options and the potential implications of implementing these should be considered as well as the optimum 'recommended' option.

4.83 For LQAs in support of site disposal consideration should also be given to the most likely future uses of the land and the remediation necessary to ensure the land will be suitable for use.

Financial Appraisal

4.84 A reliable OCE must be provided for each mitigation and management option which should include whole life costing in order that the options can be compared and a recommended option identified. This information should be restricted to the Technical Note.

4.85 When costing remedial options, please follow guidance provided within DIO Technical Bulletin 99/19, including a financial risk analysis in accordance with DIO Technical Bulletin 99/21 as outlined in Box 4.18. These DIO Technical Bulletins are provided in Annex G.

4.86 This approach includes considering both an "Average Risk Estimate" and "Maximum Likely Risk Estimate". Full details of the risks and their individual contribution to the risk element should be identified in the Technical Note only.

Box 4.18 Order of Cost Estimate (OCE) for Remediation Options

A reliable OCE is required for each remedial option and shall be carried out in accordance with DIO Technical Bulletin 99/19, entitled Order of Cost Estimates, dated July 1999. Costs are to be current quarter price levels without inflation and the relevant cost index and its source should be quoted. VAT and fees should be itemised clearly.

The OCE should include a Financial Risk Analysis, carried out and presented in accordance with DIO Technical Bulletin 99/21, entitled Estimating using Risk Analysis, dated July 1999.

Part 2A Risk Assessment Guidance

4.87 All LQAs should include an assessment of whether the site meets the criteria for designation under Part 2A. Ordinarily this assessment would be presented in the Technical Note as part of the assessment of liability within the 'regulatory context'. The following guidance is intended to assist with this assessment as part of the overall risk evaluation following changes to the statutory guidance for England and Wales introduced in April 2012.

Assessing the Significance of Contaminant/Pollutant Linkages

4.88 Within the Technical Note the assessment of potential liability within the regulatory context must include an assessment of whether the identified contaminant/pollutant linkages are:

- Resulting in significant harm being caused to the receptor in the contaminant/pollutant linkage;
- Present a significant possibility of significant harm being caused to that receptor;
- Are resulting in the significant pollution of the controlled waters which constitute the receptor, or;
- Have significant possibility to result in such pollution.

Assessing Significant Harm – Non Radioactive Contaminants

4.89 The revised Statutory Guidance on significant harm is set out within Section 4.1 of DEFRA Circular 04/12 for England and Welsh Government Guidance Document WG15450 respectively which replaces paragraphs A22 to A26 and Table A of Chapter A, Annex 3 of DEFRA Circular for England 01/2006 for England (Appendix 1) and paragraphs 1.22 to 1.26, 2.44 and Table A of Chapter 1 the NAW Guidance for Wales (Appendix 1).

4.90 The Statutory Guidance for Scotland on significant harm is set out Scottish Executive Paper SE/2006/44 Annex 2 paragraphs 2.4 to 2.6 and Table A of Chapter A to Annex 3.

4.91 When considering whether "significant harm is being caused" the statutory Guidance requires an appropriate scientific and technical assessment of all the available evidence, before a judgement is to be made on "the balance of probabilities".

Assessing Significant Harm – Radioactive Contaminants

4.92 The Statutory Guidance on harm is set out within Section 4a of the Radioactively Contaminated Land Statutory Guidance published by DECC in April 2012.

Assessing Significant Pollution and Significant Possibility of Harm – Non Radioactive Contaminants

4.93 The Statutory Guidance on significant possibility of significant harm is set out within Section 4.2 of DEFRA Circular 04/12 for England and Welsh Government Guidance Document WG15450 respectively which replaces paragraphs A27 to A34 and Table B of Chapter A, Annex 3 of DEFRA Circular 01/2006 for England (Appendix 1) and paragraphs 1.27 to 1.34 and 2.45 to 2.49 and Table B of Chapter 1 of the NAW Guidance for Wales (Appendix 1).

4.94 The Statutory Guidance for Scotland on significant possibility of significant harm is set out Scottish Executive Paper SE/2006/44 paragraphs A28 to A37 and Table B, Annex 3.

4.95 When considering whether "there is a significant possibility of significant harm being caused" the Statutory Guidance requires that a scientific and technical assessment of the risks arising from the pollutant linkage be made using relevant, appropriate, authoritative and scientifically based guidance. A significant risk of harm is considered to exist if the assessment indicates that the pollution linkage in question meets the conditions set out in the relevant Statutory Guidance for England, Wales and Scotland, and that there are no suitable and sufficient risk management arrangements already in place to prevent the harm in question. In considering whether there is a significant risk of significant harm, the Statutory Guidance advises that only the current use of land should be considered and furthermore, that account should be taken of any evidence that the current use will cease in the near future.

4.96 The 2012 revision to the Statutory Guidance for England and Wales introduced a framework comprising 4 land human health risk categories known as the Category 1-4 Approach with which to determine whether land contamination presents a significant possibility of significant harm.

- Category 1 : Human Health: unacceptably high probability that significant harm will occur if no action is taken
- Category 2: Human health : the risks posed by contamination are sufficient for the land to be deemed to meet the legal test for posing a significant possibility of significant harm
- Category 3: Human health: whilst the risks posed by contamination may not be low, the legal test with respect to the land posing a significant possibility of harm is not met.
- Category 4: Human Health: there is considered to be negligible risk to human health.

4.97 Category 1 and 2 will constitute statutory contaminated land whereas Category 3 and 4 will not.

4.98 The level of conservatism within the 2009 SGVs is such that DEFRA consider them to fall within category 4 i.e. they represent concentrations of contaminants below which the risk to human health is negligible and the land is 'very unlikely to pose a significant risk of significant harm'¹.

4.99 The revised Statutory Guidance for England and Wales also includes guidance on background or 'normal' levels of contaminants in Section 3.

Assessing Significant Pollution and Significant Possibility of Harm – Radioactive Contaminants

4.100The Statutory Guidance on the significant possibility of harm is set out within Section 4b of the Radioactively Contaminated Land Statutory Guidance published by DECC in April 2012.

Assessing Significant Pollution and Significant Possibility of Pollution of Controlled Waters

4.101 The Statutory Guidance on the interpretation of the occurrence or likelihood of pollution of controlled waters is set out within Section 4.4 paragraphs 4.34 to 4.46 of DEFRA Circular 04/12 for England and Welsh Government Guidance Document WG15450 respectively and paragraphs A38 to A42 of Part 4 to Chapter A of Annex 3, and paragraphs B50 and 52 of Part 4 of Chapter B to Annex 3 of Scottish Executive Paper SE/2006/44 for Scotland.

Policy, Process and Responsibility

4.102The Regulatory Authority must be informed if a risk of significant harm is identified. In the instance that notification is required, MOD's relevant subject matter experts (see Annex A) should be notified in order to advise on appropriate consultation and any required future action.

¹ DEFRA Circular 04/12 – EPA 1990 Part 2A Contaminated Land Statutory Guidance. 2012

5. STAGE 2 OPTIONS APPRAISAL

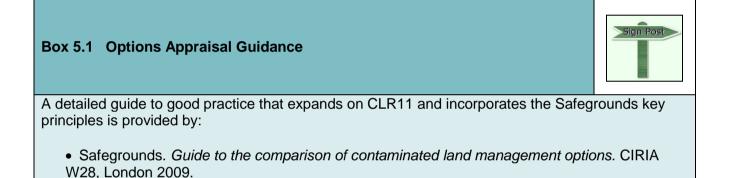
PHASE 3 LQA – MANAGEMENT OPTION APPRAISAL / DECISIONS

Introduction

5.1 Where the QRA undertaken as part of the Phase 2 LQA confirms that there is an unacceptable risk or risks posed by the presence of contamination then action must be taken to reduce (mitigate) or control (manage) those risks. This will involve some form of management response that focuses on: pathway disruption (breaking), source removal and receptor protection and can involve everything from the use of institutional controls such as fencing and standing orders to remediation or even changing the use of the land to a less sensitive one.

5.2 In general there will be more than one option to reduce or control the unacceptable risks. Therefore, to identify the optimum option or combination of options requires an Options Appraisal. According to CLR11 there are in essence three key stages to an options appraisal:

- Identification of feasible management options;
- Detailed evaluation to identify optimum option to address the RPLs; and
- Production of a Management Strategy.



Reporting

5.3 There is no default generic format for an options appraisal. However, the format must be clear, concise and reflect the needs of stakeholders.

Methodology

- 5.4 The process must:
 - Be systematic, structured and transparent;
 - Involve relevant stakeholders (this is an integral component and the extent of involvement will be specific to the situation);
 - Involve a level of detail commensurate with the nature and extent of the contamination issue come risk;
 - Consider a comprehensive range of options; and
 - Have clearly documented outputs.
- 5.5 The key steps comprise:
 - Definition of management objectives, assumptions and constraints as part of establishing the scope and context and setting out a Problem Statement;
 - Identification of options;
 - Definition of evaluation criteria (practicality, effectiveness, durability, time and VFM etc) ensuring needs of/factors important to stakeholders are reflected;

- Assessment of options against evaluation criteria. This may need to be tiered; and
- Identification of preferred option or combination of options informed by stakeholder views and technical constraints.

5.6 The options comparison process is set out below in Figure 4 within the context of the Safegrounds and CLR11 processes:

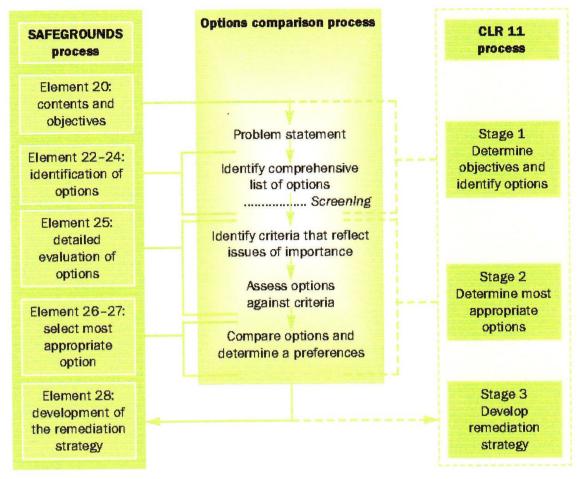


Figure 4 Options Comparison Process (Taken from CIRIA W28, 2009)

Definition of Management Objectives

5.7 Good practice is to define the strategic management objective, which will be reflected in the Management Strategy, and associated specific objectives that will inform the options comparison upfront in a Problem Statement which could include the need to address or avoid a statutory notice.

- 5.8 In R&D 66 the specific objectives are grouped as follows:
 - Contamination related these set out the required end point and are related to the CSM and risk assessment. They can be qualitative or quantitative;
 - Engineering related these relate to the modification etc of the ground conditions; and
 - Management related these range from programme requirements through to long term monitoring
- 5.9 Further information and examples are presented in R&D66.

5.10 The key constraints should be documented in the statement such as: time, cost, policy, need for licenses etc. and conditions set by regulators and other stakeholders etc. Also the fate of the land / site may be a key constraint, particularly if it is going for disposal or is to be redeveloped.

5.11 The critical assumptions should also be recorded, namely those that address the principal areas of uncertainty such as the extent of the contamination, access and the presence or otherwise of UXO etc. These will of course be site and situation specific.

Identification of Options

5.12 These must be practical and address the RPLs such that the unacceptable risks associated with the land contamination and / or controlled water pollution is addressed. The options should be distinct and range from the 'do minimum' to the 'maximum possible'. Whilst CLR11 focuses on the identification of options to address individual RPLs the more holistic approach advocated by CIRIA W28 is preferable as this should produce a more integrated cost effective solution.

5.13 It may be appropriate to sift the options at this stage in order to whittle the number down to a manageable size. The criteria may include: technical feasibility, acceptability to stakeholders and time available. Whatever set of criteria is selected it must be defensible and ideally should reflect the needs of the stakeholders.

Assessment Criteria

5.14 These must support the objectives and the interests of the stakeholders. Ideally they must avoid bias and reflect factors such as: policy, practicality and economic considerations. The latter should not focus solely on capital cost, but the through life costs and sustainability.

Assess Options

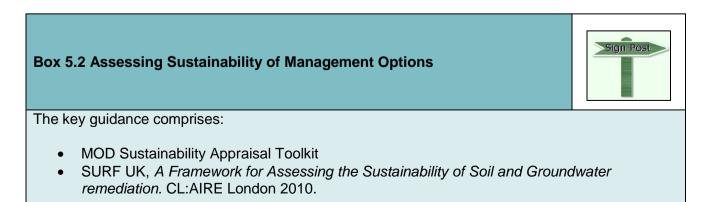
5.15 The assessment must be objective, auditable, structured and focused on the pros and cons of each option within the context of the objectives and associated constraints and needs / interests of stakeholders in order to identify a preferred option / combination of options.

5.16 To undertake this assessment it is good practice to use a scoring matrix based on either:

- 'Relative' scores are given in relative terms i.e. rank options worst to best etc; or
- 'Absolute' options are scored according to a scale that defines performance according to a number of categories from low to high.

5.17 For complex sites it may be necessary to undertake a tiered assessment to narrow down the range of options which may need to be combined to deliver the required risk reduction and management.

5.18 SURF UK has in partnership with CL:AIRE developed a framework for assessing the sustainability of remediation options. This should be used to identify the relative sustainability of the available remediation options as part of the investment appraisal of each option. Where the options do not involve remediation or only in part then it may be appropriate to undertake a Sustainability Appraisal.



Development of Management Response Strategy

5.19 Once the preferred management option is identified the Management Strategy referred to as the 'remediation strategy' must be developed. This strategy must be capable of either managing or mitigating the unacceptable risks associated with the identified RPLs. It is essential that the decision process leading to the strategy is documented, sets out the considerations, assumptions and priorities and reflects the interests of the stakeholders.

5.20 The type of response will be dependent upon the level of risk and the nature of the hazard. In the case of remediation, the strategy, clearance levels and end point need to be agreed with the appropriate regulatory authority. The strategy must address the practicalities associated with the site and situation in question such as how the:

- Strategy will be implemented;
- Site will be zoned; and
- Success of the strategy will be measured in terms of the strategic and site specific objectives being met.
- 5.21 The strategy must also take account of:
 - Precedents for other sites;
 - Departmental policy, objectives and commitments;
 - Needs of stakeholders and whether they will be adequately addressed; and
 - Sustainability.

5.22 Finally the strategy must be justified and represent VFM within the constraints that apply. In the words of CLR11 the strategy should be 'acceptable on cost-benefit grounds'.

Box 5.3 Justification and Optimisation/ Reasonableness of Remediation

If the site has been designated as 'contaminated land' under Part 2A then Justification and Optimisation studies will be required to justify the need for and optimum form of the necessary remediation.

The justification element can be accommodated as an extra step within the Options Appraisal process outlined previously.

The optimisation element should be integral to the process anyway so no additional step is required.

5.23 Effective stakeholder involvement is essential if this phase of the LQA is to be a success. Specialist technical advice is available from the SMEs listed in Annex A. DIO can advise on the technical aspects and application of remediation techniques and both develop and implement the management strategy on behalf of a TLB or project. Equally, DIO can advise on the need or otherwise for remediation in the immediate to long term and whether institutional controls are the most appropriate response.

Box 5.4 Stakeholder Involvement



At the Phase 3 stage the level of stakeholder involvement will typically focus on the following:

- Identification, assessment and comparison of management options
- Identification of preferred management option or options
- · Decision on which option or options to proceed with

The Stakeholder Involvement Plan and Programme should be amended to reflect any changes.

Policy

5.24 Current MOD Policy is to undertake remediation where there is a significant (unacceptable) risk to health, safety and the environment taking account of the current or intended use. In the case of sites in disposal, it is generally confined to defence specific contaminants, such as chemical agents, where it is unreasonable to expect a civilian contractor to be able to undertake the work. Otherwise, sites are sold in an un-remediated condition with a view to the purchaser undertaking the remediation necessary to make the site suitable for its intended use. In the case of radiological contamination it is MOD policy to require independent verification of purchaser-managed remediation.

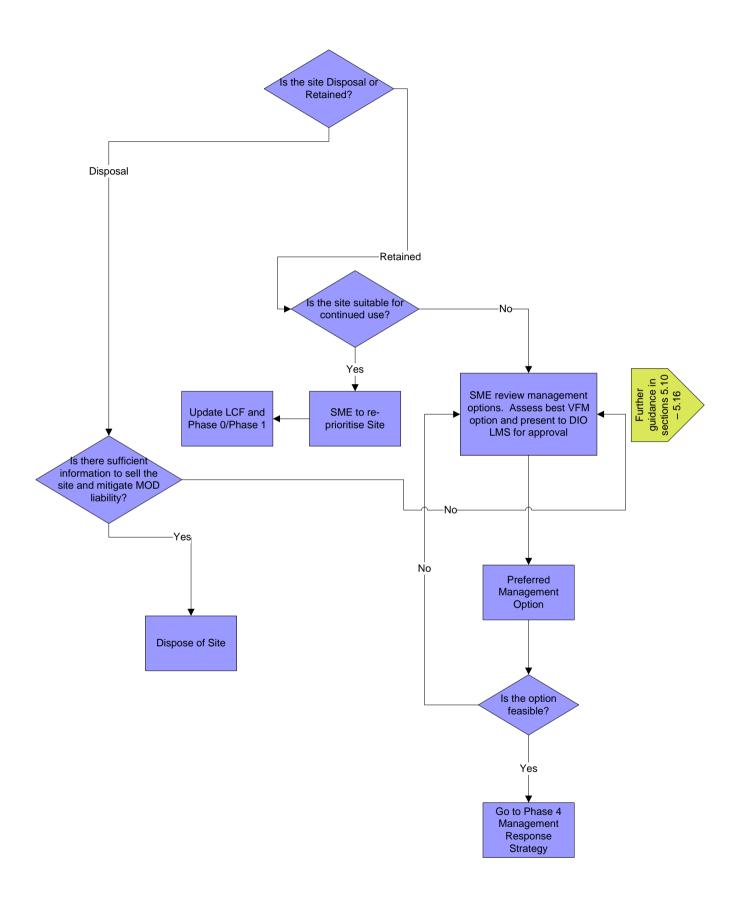
5.25 Further clarification of MOD Policy on remediation is available in the Design and Maintenance Guide 12, Site Closure Guide, Land and Property Policy: PI 6/2005.

Waste Management

5.26 It is essential that waste arising from any management response, particularly as part of a remediation scheme, is minimised and managed appropriately. Guidance on sustainable waste management is provided at Annex C.

5.27 The Phase 3 LQA Process is summarised in Figure 5.





6. STAGE 3 MANAGEMENT RESPONSE

PHASE 4 LQA – IMPLEMENTATION OF MANAGEMENT OPTION(S) (THIS MAY ALSO BE PHASED AND INVOLVE LONG TERM MONITORING)

Implementation

6.1 The starting point is the development of the Implementation Plan which will set out and define:

- The strategic and site specific objectives;
- Programme;
- How stakeholder interests/needs including those of the regulator will be met;
- How the inherent uncertainties such as the actual extent of contamination will be managed;
- Level of supervision;
- How regulatory compliance will be achieved;
- Financial management process;
- How the work will be documented and validated;
- End points and critical success factors; and
- Scope and duration of long term management and monitoring requirements.

6.2 The plan must also take into account of commercial, contractual and policy considerations such as tendering and letting of contracts. For remediation projects a re-measurement contract may be more appropriate than a fixed price as this provides for more equitable risk sharing and avoids contractors having to front load the risk associated with the uncertainty over ground conditions etc and reflect this in their bid.

6.3 In addition the plan must address the financial aspects of the strategy. It may therefore be useful to establish a Financial Management Plan detailing milestone payments etc.

Monitoring and Maintenance

6.4 Where the management strategy involves on-going monitoring such as groundwater monitoring to check that the situation does not deteriorate and/or the maintenance of physical access restrictions such as fencing then it is good practice to develop a Monitoring and Maintenance Plan. Similarly if the management strategy involves some form of remediation that has a monitoring and/or maintenance requirement such as the use of a capping layer then this too should be covered by such a plan.

6.5 Typically a Monitoring and Maintenance Plan comprises:

- Scope of work;
- Technical specification;
- Locations, frequency and duration of monitoring activities;
- Analytical suite with limits of detection etc;
- Evaluation criteria such as EQSs in the case of groundwater;
- Reporting schedule; and
- Contingency plan in case monitoring indicates the remediation has or is failing or the situation is deteriorating such that remediation may be required.

6.6 According to CLR11 these form the 'lines of evidence' necessary to demonstrate success.

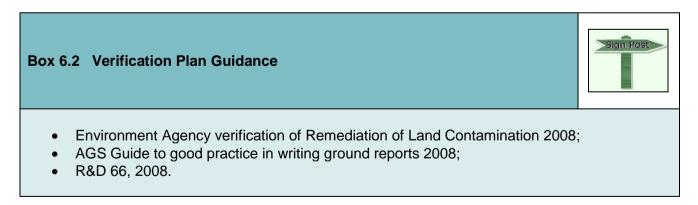
Verification

6.7 Upon completion of the Implementation Plan a Verification Plan should be developed that sets out the activities and data necessary to demonstrate that the objectives set out in the Management Strategy have been achieved. This is particularly important where remediation will be undertaken. The plan must therefore also address:

- Planning conditions;
- Licensing and permitting requirements; and
- Specific condition imposed by regulators such as control of dust etc.
- 6.8 The Verification Plan should confirm/verify:
 - The nature and extent of the residual contamination and/or pollution;
 - That imported materials and those destined for re-use on site are suitable for use;
 - Whether the management strategy is succeeding, i.e. meeting the objectives;
 - That the management strategy is not causing land contamination or pollution of controlled water;
 - Compliance with planning conditions, licences, permits and consents



6.9 Ultimately this should be reported within a Verification Report which according to CLR11 should provide 'a complete record of all remediation activities on site and the data collected as identified in the verification plan to support compliance with agreed remediation management objectives and criteria. It also includes a description of the work (as-built drawings) and details of any unexpected conditions (e.g. contamination) found during remediation and how they were dealt with'.



6.10 In addition to the verification report, monitoring reports will need to be produced separately at the required intervals. Once the monitoring confirms that the management strategy or the remediation component has complied with the objectives/compliance criteria then the verification report can be finalised.

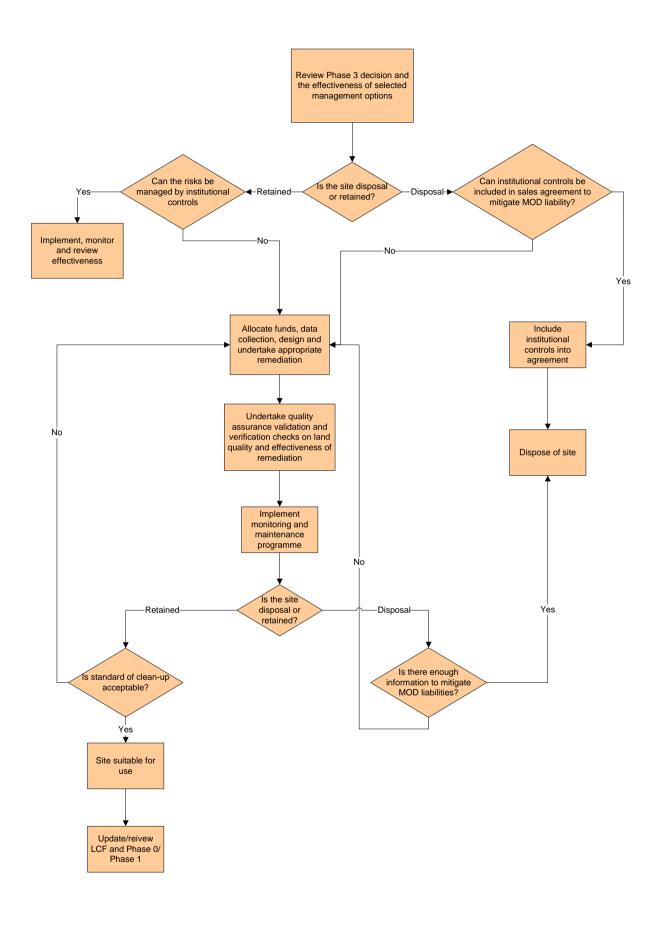
6.11 Management responses include: do nothing, the use of institutional controls, such as standing orders and the permit to dig system, as well as remediation. However, remediation will be of reduced value if not properly documented. Quality control must therefore, be managed throughout the design and implementation of a remediation programme.

6.12 A post remediation validation survey must therefore be carried out together with a post project review and the lessons learnt circulated.

Policy, Process and Responsibility

6.13 The Phase 4 Management Response Process Map, providing guidance on the management process, is presented as Figure 6.

Figure 6 Phase 4 Management Strategy Flow Chart



7. LQA RECORD RETENTION

7.1 Once a LQA report has been completed for site, electronic and hard copies are sent to site for their records. The LQA report or relevant information from the LQA can be then passed to any contractors coming to work on site via the SoKH system to inform them of any land quality issues for consideration within the contractors risk assessments and method statements.

7.2 Electronic copies of all LQA Reports, Technical Notes and associated documentation such as Remediation Validation Reports will also be uploaded onto the DIO LQA Database managed by the DIO EOLM team. Details of the electronic format required to achieve this is presented in Annex B4.

7.3 Upon completion of LQA or remediation works, each TLB is responsible for ensuring the relevant LQA information is kept up to date for land and property retained by MOD. It is recommended for any proposed developments on site or following pollution incidents, DIO EOLM are contacted to advise whether an LQA update is required, See Box 7.1 below.

Box 7.1 LQA Shelf Life and Decision Making



A LQA provides a snap-shot in time, therefore the risk assessment and management measures detailed within the LQA should be reviewed whenever there is a significant change to site activity, change in relevant legislation, change of statutory guidance and/or a pollution incident occurs.

Where significant land contamination risks have been identified, these should be kept under review through the site Health & Safety and EMSs.

Where an investment decision is to be made based on a pre-existing LQA the parameters and assumptions within the LQA should be checked to ensure they are still valid. LQAs must be tailored to the situation and undertaken by competent persons.

7.4 In addition, where available, site LCFs and site based EMS's can be used to identify a need to update an LQA. The EMS provides an effective vehicle to identify when an update is required and the LCF provides a structured means of retaining and summarising land quality information and ensuring known areas of contamination and potential hazards are identified. It should be noted though that the LCF is not a substitute for an LQA.

7.5 For further information on LCFs please refer to guidance documents detailed in Box 7.2.

Box 7.2 – LAND CONDITION FILES (LCFs)	í					
 Guidance on the use and form of the LCF can be found in: DIO Information Note IN 03/09 LCF; and Safegrounds: 'Good practice guidance for land quality records management for nuclear licensed and defence sites, CIRIA W21 London 2007. 						

ANNEX A

ANNEX A

SUBJECT MATTER POINTS OF CONTACT – LQA

1. Policy

FMC Cap Infra Policy Team

Tel: 0121 311 3648

2. LQA Process, Practice, Procedures, Risk Assessment and Reporting

DIO EPS EOLM

Tel: 0121 311 3618

3. UXOPRA

DIO EPS EOLM

Tel: 01225 847230

ANNEX B1

GENERIC MOD PHASE 0 LQA FORMAT

LQA REPORT STANDARD FRAMEWORK – OFFICIAL

Phase 0 Land Quality Assessment (LQA)	Project No.
Purpose of Phase 0 LQA	
Site Description	
Site Name	
Site Address	
Parcel Name (s)	
National Grid Reference (NGR)	
Size (Hectares)	
Site location plan	Site layout plan
Previous Reports	
Site History	
Onsite	Adjoining Land (<500m)
Buildings and Infrastructure	
Buildings	
Infrastructure	
Services	
Ancient Monument	
Geology	
Drift / Bedrock	
Receptors - Controlled Waters Groundwater	
Aquifer Classification	1
Groundwater flow direction	
Source Protection Zone (SPZ)	
Abstractions	
Leaching Potential	
Sensitivity	
Surface water	
Distance and direction from site	
Classification	

Abstractions								
Flow Direction	n							
Sensitivity								
Receptors –	Human Health							
Location (ons	ite or offsite)							
Activities and	Activity Patterns	;						
Sensitive Sub	populations							
Sensitivity								
Receptors –	Ecosystems							
Distance and	direction from si	te						
Designation								
Sensitivity								
Receptors –	Livestock, Prop	perty, Flo	ora &Faui	na & Othe	r			
Outline Cond	ceptual Site Mod	del (CSN	I) and Ca	tegory As	sessn	nent		
Potential	Receptor	Pathw	/ay	Sensitivit	y	Potential of		Category
Source						Occurrence		Assessment
Onsite	T	-				T		r
Adjoining La	nd (<500m)				T		-	
Land Quality Statement								
Author								
Reviewer								
Date								

ANNEX B2

GENERIC MOD PHASE 1 LQA FORMAT

LQA REPORT STANDARD FRAMEWORK - OFFICIAL

Land Quality Statement (takes place of Executive Summary)

Introduction Site description and History Site Sensitivity Potential Sources of Contamination Summary of Potential Risks Overall Land Quality and Suitability for Redevelopment

1. Introduction

- 1.1 Terms of Reference
- 1.2 Objectives
- 1.3 Methodology (including sources of information)
- 1.4 Structure of this Report

2. Site Description and Setting

- 2.1 Site Location
- 2.2 Site Layout
- 2.3 Site Activities
- 2.4 Site History
- 2.5 Site Constraints
- 2.6 Surrounding Area

3. Summary of Previous Reports

4. Environmental Setting and Site Sensitivity

- 4.1 Geology
- 4.2 Hydrogeology
- 4.3 Hydrology
- 4.4 Site Sensitivity

5. Preliminary Conceptual Site Model

6. Summary of Potential Contaminant Sources

- 6.1 Current Potential Sources (onsite/offsite)
- 6.2 Historic (onsite/offsite)
- 6.3 Discounted (onsite/offsite)
- 6.4 Potential Pathways
- 6.5 Potential Receptors

7. Preliminary Risk Assessment

- 7.1 Methodology
- 7.2 Risk Assessment Table

7.3 Risk Summary (To include headings such as: Current and Future Site Users, Construction Users, Groundwater etc.)

8. Overall Land Quality and Suitability for Use (Current/Redevelopment)

9. References

Figures (this is not an exhaustive list, site specific figures may be required) Figure 1: Site Location Figure 2: Site Layout

Figure 3: Areas of Potential Concern Figure 4: Preliminary Conceptual Site Model

Appendices (to include but not limited to): Searches information – local authority, DSTL, UXO, BGS Database Search, e.g. Envirocheck or similar Aerial photography Relevant site walkover photographs providing overview of potential sources of contamination and general site characterisation, where appropriate these should be referenced in text and on figures

TECHNICAL NOTE – OFFICIAL SENSITIVE – COMMERCIAL (see explanatory notes in annex B4)

- 1. Background
- 2. Summary of Findings
- 3. Preliminary Risk Assessment (with MOD liability classifications)
- 4. Liability Assessment
- 5. Options Appraisal
- 6. Conclusion and Recommendation(s)
- 7. References

Figures (this is not an exhaustive list, site specific figures may be required)

- a. Site Location Plan
- b. Site Layout Plan

c. (If applicable) Proposed exploratory locations discussed for a management option (overlain on APC plan)

Appendices

a. Summary of LQA (LQA data capture proforma*)

*To be included as spreadsheet on final disc but copy provided as an appendix. Please see Annex B4, at end of Section 7 in practitioners guide for further information on how to complete this section.

- b. Risk Assessment Table including liability classifications
- c. Cost Breakdown for Option 1 and 2
- d. Rough Order of Cost Estimate for Option 3

ANNEX B3

GENERIC MOD PHASE 2 LQA FORMAT

LQA REPORT STANDARD FRAMEWORK - OFFICIAL

Land Quality Statement (takes place of Executive Summary)

Introduction Site description and History Site Sensitivity Sources of Contamination Summary of Risks Overall Land Quality and Suitability for Redevelopment

1. Introduction

- 1.1 Terms of Reference
- 1.2 Objectives
- 1.3 Methodology (including sources of information)
- 1.4 Structure of this Report

2. Site Description and Setting

- 2.1 Site Location
- 2.2 Site Layout
- 2.3 Site Activities and Operations
- 2.4 Surrounding Area

3. Summary of Phase 1 LQA

- 3.1 Site History
- 3.2 Environmental Setting
- 3.3 Site Sensitivity
- 3.4 Environmental Condition of the Site
- 3.5 Sources of Potential Contamination
- 3.6 Preliminary Risk Assessment

4. Site Investigation

- 4.1 Objective
- 4.2 Scope of Work and Rationale
- 4.3 Methodology

5. Findings of Site Investigation

6. Quantitative Risk Assessment

- 6.1 GQRA
- 6.2 DQRA

7. Updated CSM and Risk Assessment

8. Overall Land Quality and Suitability for Use (Current/Redevelopment)

9. References

Figures (this is not an exhaustive list, site specific figures may be required)

Figure 1: Site Location

Figure 2: Site Layout

Figure 3: Exploratory Hole Location (showing areas of potential concern)

Figure 4: Areas of Visual/Olfactory contamination

Figure 5: Groundwater Contours (presenting indicative flow direction)

Figure 6: Locations of Exceedances

- Soils
- Waters

Figure 7: Updated Conceptual Site Model

Appendices (this is not an exhaustive list, site specific appendices may be required): Photographs - Photographic record of site investigation Exploratory Hole Logs Monitoring Data Laboratory Certificates Screening of Laboratory Data Risk Assessment Methodology & Outputs (if applicable)

TECHNICAL NOTE – OFFICIAL SENSITIVE – COMMERCIAL (see explanatory notes in Annex B4)

- 1. Background
- 2. Summary of Findings
- 3. Tier 2 Risk Assessment (with MOD liability classifications)
- 4. Liability Assessment
- 5. Options Appraisal
- 6. Conclusion and Recommendation(s)
- 7. References

Figures (this is not an exhaustive list, site specific figures may be required)

- a. Site Location Plan
- b. Site Layout Plan

c. (If applicable) Proposed exploratory locations discussed for a management option (overlain on APC plan)

Appendices

a. Summary of LQA (LQA data capture proforma*)

*To be included as spreadsheet on final disc but copy provided as an appendix. Please see Annex B4, at end of Section 7 in practitioners guide for further information on how to complete this section.

- b. Risk Assessment Table including liability classifications
- c. Cost Breakdown for Option 1 and 2
- d. Rough Order of Cost Estimate for Option 3

ANNEX B4

EXPLANATORY NOTES FOR GENERIC PHASE 1 AND 2 LQA REPORT AND TECHNICAL NOTE FORMATS

1. For Phase 1 and 2 LQAs

LQS

This is a non-technical summary (**2 to 3 pages maximum**) of the land condition together with its suitability for re-use. The LQS must present the presence, nature and extent of known contamination and pollution of controlled waters in context establishing whether the majority of the land is likely to be free of contamination. For instance, where present, is contamination localised and limited in extent, associated with and restricted to fill materials within made ground and so on? The effects of the known contamination etc on the development potential must also be addressed particularly where the LQA is in support of a site disposal.

The LQS must not include reference to recommendations for further work and all risks must be presented in context. For instance in a Phase 1 LQA the 'Overall Land Quality and Suitability for Redevelopment' section could be phrased along the lines of:

'Overall based on the information available, the majority of the site is unlikely to have been contaminated as a result of historical and/or current activities. Where present land contamination is likely to be limited in extent, comprise...... and be confined to fill materials within areas of made ground. If present contamination will pose a low to negligible risk to health and the wider environment providing the areas affected remain undisturbed.

The site is currently suitable for use as.....and is likely to be suitable for,,,, subject to the necessary investigation and assessment.

TECHNICAL NOTE (OFFICAL SENSITIVE-COMMERCIAL)

Content

a. **Background**: Brief details of the terms of reference, aims and objectives, methodology and constraints and structure of the report.

b. **Interpretative information**: Must include, but not necessarily be limited to: quantitative risk assessment including MOD liability classification and comments, liability assessment including regulatory context, management options appraisal, recommended option/s and conclusions. The liability assessment should include a brief summary of relevant legislation followed by description of LQA findings in a regulatory context. As a minimum this should include the likelihood of the classification of the site as contaminated land under Part 2A.

c. **Management Options Appraisal**: Options included should demonstrate how to manage and mitigate the potential environmental health and liability risks identified and consider the future of the site. A detailed summary should be provided for each of the proposed options, including an associated limit of liability, with a breakdown of costs and fees in appendix, as well as any drawings sufficient to explain the option(s) to those unfamiliar with the site. A very brief comparison of advantages and disadvantages of each option considered should be included. Ordinarily 3 key options will be considered inclusive of the 'do nothing/do minimum', and if to support a disposal it should also be considered whether the site can be sold with knowledge and if so reference the appropriate exclusion test. This section should not indicate the preferred management option.

d. **Conclusion and Recommendation**: Giving clear-cut recommendations, the option preferred and the operational and economic advantages (including timescales) that justify the recommendation(s).

e. **Summary of LQA**, also to be completed electronically.

Ordinarily the Technical Note will not be released to third parties outside of MOD.

2. Phase 1 LQA Report

a. **Introduction**: Briefly set out the terms of reference under the Client commission, together with the: objectives of the LQA phase, methodology and structure of the report. This should include any constraints.

b. **Site Description and Setting**: Set out details of the site, or sites, giving location(s), layout and MOD unit(s) establishing zones where necessary and including brief details of the surrounding area. Include description of the proposed future use where applicable. Where key aspects can be shown more clearly in the drawn form then drawings and/or diagrams should be included. There should be no straying into a discussion of potential sources of contamination. Site Activities section should include details of the site walkover. Site History section should provide detailed description of the information obtained from the various research sources and split into separate subsections where necessary.

c. **Summary of Previous Reports and Sources of Information**. This should set out the sources and nature of the information reviewed/considered including interviews with site personnel. Where previous LQA and/or other desk study and investigation reports etc exist then their scope, purpose, limitations and pertinent findings should be summarised

d. **Environmental Setting**. This should summarise the geology, hydrogeology, and hydrology based on available information before presenting a summary assessment of the site sensitivity as follows:

 Box x.x Example Groundwater sensitivity summary
 Image: Comparison of the sensitivity: Moderate

 Groundwater sensitivity: Moderate
 The site is underlain by Secondary A and Secondary B aquifers. There are licensed groundwater abstractions within 1km though the site does not lie within a groundwater SPZ.

e. **Potential Sources of Contamination**: Details of the potential sources and the basis on which they have been identified should be described under relevant subsections with sources grouped accordingly. A summary table should be provided with each potential source provided a unique reference number (URN) which relates to its location on a Areas of Potential Concern plan (Annex B6).

f. **Preliminary Conceptual Site Model**. The potential contaminant sources, pathways and receptors need to be identified together with the necessary rationale and summarised pictorially as per Annex D to this guide. The potential sources should summarised in a table with each potential source provided a unique reference number (URN) which relates to its location on a Potential Sources of Contamination/Areas of Potential Concern plan and carried forward to subsequent risk assessment table and CSM.

Box x.x Example Potential Contaminant Sources Summary



Potential Source Number (carried forward to RA and CSM)	Potential Source	Associated Feature	Potential Source Location (shown on figure xx)
1	Vehicle Maintenance Areas – potential for leaks or spillages associated with	Historical aircraft maintenance hanger	1.1
	current and historic maintenance activities	Historical MT section	1.2
	(hydrocarbons, antifreeze, metals, solvents and acids)	Current central servicing workshop	1.3
2	Historic Fuel Storage – potential for leaks or spillages associated with historic fuel/oil storage (hydrocarbons)	Redundant BFI1	2
3	Current Fuel/Oil Storage – potential for leaks or spillages associated with	Current main POL area	3.1
current fuel/oil storage (hydrocarbons)		Current MTFI	3.2
etc.	etc.	etc.	etc.

g. **Preliminary Qualitative Risk Assessment**. To comprise text and risk assessment table as per Annex E to this guide.

h. **Overall Land Quality and Suitability for Use**: This should reflect the summary presented in the LQA and be of a form that is easily understood by the layperson. Summarising whether the findings of the LQA indicate the site is suitable for current use or future proposed use, whichever is most suitable for the objective of the LQA.

3. Phase 2 LQA Report

a. **Introduction**: As per Phase 1 briefly set out the terms of reference under the Client commission, together with the: objectives of the LQA phase, methodology and structure of the report. This should include any constraints.

b. Site Description and Setting: Set out brief details as per Phase 1 LQA.

c. **Summary of Phase 1 LQA**: provide brief summary and over view including historical use of the site.

d. **Site investigation and Site Investigation Findings**. Detail: objective, investigation methodology, rationale and findings (including ground conditions). The sampling and analysis strategies must be consistent with accepted best practice, documented and justified. The

sampling methodologies must also be consistent with best practice, documented and justified together with the QA and QC measures taken. All assumptions must be clearly set out.

e. **Quantitative Risk Assessment.** The approach and assessment criteria must be documented and justified and the findings summarised both in the text and as a summary table (refer Annex E).

f. **Updated CSM and Risk Assessment**. This must be updated in light of the Phase 2 LQA findings.

g. **Overall Land Quality and Suitability for Use**. This should reflect the summary presented in the LQA and be of a form that is easily understood by the layperson. Summarising whether the findings of the LQA indicate the site is suitable for current use or future proposed use, whichever is most suitable for the objective of the LQA.

4. LQA REPORT AND TECHNICAL NOTE COVER SHEET

This should contain:

a. Title "DIO" centred at the top of the sheet and the DIO and MOD logos should be shown.

b. Title as shown on the Statement of Requirement in the centre of the sheet, with the Project No. immediately below.

c. Name and address of the relevant TLB office in the bottom left hand corner of the sheet.

d. The legend "Prepared by (name) for the MOD, TLB (e.g. DIO), under commission (contract number)" in the bottom right hand corner of the sheet along with the month and year in which the Study was produced.

e. Marked areas for 'prepared by' and 'authorised by' including SiLC or appropriate professional logo and registration number

f. The final report must be signed by the PM and reviewer.

g. In order to be compliant with the Data Protection Act, all reports must, with the exception of the cover sheet, have the names and initials etc. of individuals removed from the main body of the text. Other than the cover sheet there is to be no personal information included in the report.

4. SIZE

This should always be A4 vertical format, but may contain folded A3 or larger sized sheets in clear A4 pocket inserts.

5. MAPS AND PLANS

Maps should indicate the site location and where scale permits should also show the site boundary. Site Plans should include a scale bar, a north arrow and a key.

6. GUIDANCE NOTES FOR THE COMPLETION OF LQA RETURNS

a. The excel spreadsheet (compatible with Microsoft Office Version 2007) will be saved on a CD and submitted with the final reports to the Task Officer. A data shape (polygon) containing tagged spatial information should accompany the LQA Data Capture Proforma excel spreadsheet shown below.

b. The spreadsheet fields will be filled out as follows:

- (1) Site DPR Ref.#: this is only to be filled out if known
- (2) Site Name: provide current name and aliases
- (3) Area: provide area in hectares
- (4) Grid Reference: provide 8 figure grid ref. for the centre of the site
- (5) LQA Priority: insert the priority number as follows:

Priority 1: Land identified for disposal or subject to rationalisation or where significant change in land use is envisaged.

Priority 2a: Land in sensitive area and with known or suspected contamination Priority 2b: Known threat; site in sensitive area such as major aquifer Priority 2c: Strongly suspected threat or possible threat from e.g. radioactive substances, dioxins, CW materials

Priority 2d: No known evidence if threat, i.e. all other sites

- (6) Current LQA Phase: state whether it is
 0 Prioritisation; 1 Desk Study; 2 Site Investigation; 3 Assessing need to remediate; 4 Remediation
- (7) Overall Land Quality: please insert appropriate number

1 No known or potential sources of contamination

2 Majority of the site is unlikely to be contaminated. A number of localised sources of contamination are or may be evident.

3 Majority of the site is or is likely to be contaminated.

(8) Approximate area of contamination: Please estimate area likely to be affected in m²

(9) Liability Class: This should be presented as the risk assessment table within the Technical Note.

7. ELECTRONIC FORMATTING OF LQA REPORT AND TECHNICAL NOTE

a. The format of the LQA Report and Technical Note is to conform to the standard as outlined in annexes B and amended where appropriate in agreement with the task officer.

b. Electronic copies of the LQA Report and Technical Note should typically be submitted as two separate pdfs, with each pdf containing the text, figures and appendices taking full account of the necessary security protocols. The technical note cannot be sent over email due to its security classification, unless at the discretion of the task officer.

c. CD1 should contain the finalised LQA report only. CD2 should contain the finalised LQA report, Technical Note, excel spreadsheet and data shape file.

d. The CD spine should clearly denote the Project Number, Site Name, Phase of Works, Consultant Company and Date e.g. '12345 RAF Banner Phase 1 WatCon Ltd June 2005'. The front of the CD should denote the Site Name, Phase of Works, LQA and Technical Note, Security Classification, Project Number, Report Status, Date of Issue, and the DIO and the Consultant Company Logos. The CD itself should be similarly marked as the front cover.

LQA DATA CAPTURE PROFORMA

1. DIO LQA Ref	2. Report Date (Month/Year)	3. Site Name	4. Investigation Phase	5. Report Type	6. Volume No.	7. Author	8. Polygon Area (Ha)	9. Eastings (6 Fig)	10. Northings (6 Fig)	11. Country	12. Priority Rating	13. Polygon Supplied	14. Total spend to date
----------------------	-----------------------------------	--------------------	------------------------------	----------------------	---------------------	--------------	----------------------------	---------------------------	-----------------------------	----------------	---------------------------	----------------------------	----------------------------------

 DIO LQA Ref: 5 numbers identifying project number Report Date: Month/Year e.g. 09/2007 	12. Priority Rating	Comment assuming likely or high likelihood of pollutant linkage occurrence	Action required		
 3. Site Name: e.g. RAF Hythe (Site 4), 4. Investigation Phase: LQA Phase 0 LQA Phase 1 LQA Phase 2 LQA Phase 3 LQA Phase 5 BLANK 	Priority 1	Site probably not suitable for present use and/or environmental setting. Contaminants probably or certainly present and probably have an unacceptable impact on identified sensitive receptors.	Urgent action required in the short term		
5. Report Type:6. Volume No:Technical NoteVol 1LQA ReportVol 2Regulator ReportVol 3OtherVol 4	Priority 2	Site may not be suitable for present use and/or environmental setting. Contaminants probably or likely to be present, and may have an impact on identified sensitive receptors	Urgent action required in the short term		
 Author: Company Name of Consultant Polygon Area: the area (ha) of the work covered in the report Eastings: centre of the site (6 figures) Northings: centre of the site (6 figures) Country where site is located, please insert country code in accordance with <u>ISO 3166</u>. (E.g. England is GB-ENG). 	Priority 3	Site considered likely to be suitable for present use and/or environmental setting. Contaminants may be present but unlikely to impact sensitive receptors identified.	No immediate action needed while site remains in present use and remains undisturbed. Management options to prevent land contamination may need to be implemented in order to reduce the risk of land contamination		
12. Send an electronic GIS polygon of the site area with attached spatial data to be incorporated into a GIS system.13. Total spend to date rounded up to the nearest pound.	Priority 4	Site considered suitable for present use and/or environmental setting. Contaminants may be present but very unlikely to have an unacceptable impact on key targets.	No action needed while site remains undisturbed. In general management options to prevent land contamination are likely to be sufficient, although a review of preventative measures should be undertaken periodically.		

ANNEX B5

ANNEX B5

EXPLANATORY NOTES FOR CDM HEALTH AND SAFETY FILE

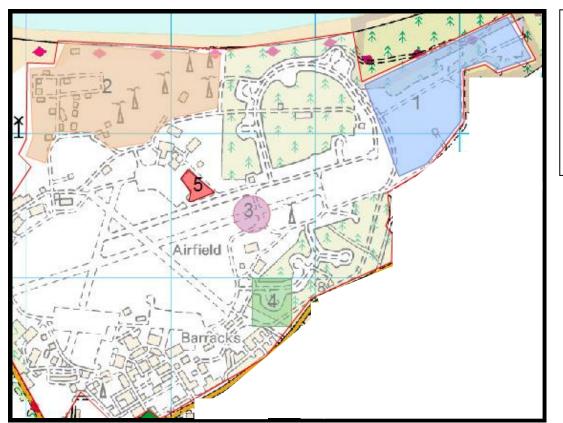
See Appendix 4 The health and safety file of the CDM Regulations 2015 document. Please include as a minimum the following outlined below:

- Project Contacts (Client, Principal Designer, Principal Contractor, Other Contractors)
- Description of Works
- Significant Hazards/Risks Encountered During the Works (include information on how these were mitigated)
- Significant Hazards/Risks Remaining Following the Works
- Hazardous Materials Used During the Works
- Equipment Used During the Works / left in-situ (How was this removed/dismantled)
- Significant Services (nature, location and marking)
- Information and As-built Drawings (e.g. Exploratory location plan)
- Attachments / Reference Material (e.g. additional LQAs, UXO reports, Hazard Plans)

ANNEX B6

Example Figure: Areas of Potential Concern

Figure should use site plan as base with areas of potential concern colour coded and numbered, described in a legend detailing the former/current site use and potential contaminant.



LEGEND

- 1 Potential for Radiological Material
- 2 Potential for Ordnance/Explosive Residues
- 3 Known area of historic fuel spill
- 4 Known area of historic tipping
- 5 Former BFI (potential for hydrocarbons)

ANNEX C

SUSTAINABLE WASTE MANAGEMENT

Introduction

1 Apart from the cost associated with the disposal of waste arising from LQAs and in particular remediation works, MOD is committed to reducing the amount of waste sent to landfill and following the more sustainable approach set out by SURFUK. It is essential that every effort is made to reduce the amount of waste produced in the first instance and where possible reuse materials.

2 To achieve this with regard to remediation and subsequent work in aide of site development it is essential that the waste management hierarchy is embedded and followed namely:

- prevention
- preparing for reuse
- recycling
- recovery
- disposal.

3 CL:AIRE has derived a framework for managing the movement of site-won soils, which is detailed within their guidance document titled 'Definition of Waste: Industry Code of Practice' (presently v2). The Code of Practice (CoP) supports the use of materials in accordance with the waste hierarchy, such that waste is minimised, recovered and reused, subject to satisfying the underpinning principles of the CoP. The aim of the framework is for the utilisation of chemically and geotechnically suitable soils in a sustainable manner.

4 The CoP sets out good practice for establishing on a site by site basis whether excavated materials are 'waste' and/or treated excavated waste can cease to be waste for a particular use. The guidance documents detailed in Box C4 will provide an understanding of the issues surrounding the re-use of soils on a particular site:

Box C4	Key Supporting Guidance
•	"Industry Profiles" (Department of the Environment) "Verification of remediation of land contamination" (EA, 2010) BS 10175:2011+A1:2013 "Investigation of potentially contaminated sites – Code of practice
•	SURF UK, A Framework for Assessing the Sustainability of Soil and Groundwater Remediation, 2010. CL:AIRE Definition of Waste: Development Industry Code of Practice (v2, 2011)

5 It must be noted that the CoP is voluntary and applies to England and Wales only at this time, but it has the support of the EA building on their guidance document: "Definition of waste: Developing greenfield and brownfield sites" (2006). Hence, the EA will take account of the CoP when deciding whether excavated materials should be regulated as waste. If materials are dealt with in accordance with this CoP they will take the view that those materials are unlikely to be waste if they are used for the purpose of land development

6 The CoP covers excavated materials namely: top soil and sub-soil, underlying geology; soil and mineral based dredgings; ground based infrastructure that is capable of reuse within earthworks projects, e.g. road base, concrete floors; made ground; source segregated aggregate material from demolition work, crushed brick and concrete to be reused on the site of production within earthworks projects or as sub-base or drainage materials; and stockpiled excavated materials.

7 The CoP also applies to uncontaminated and contaminated material for use on the site from which it has been excavated, either without treatment or after on-site treatment as part of the development of that land (i.e. re-use on site of origin Scenario). It also applies to such material for use following treatment at a Hub Site (Cluster Project Scenario) or without treatment at another development site subject to the material meeting the requirements set out in the CoP (Direct Transfer Scenario).

8 However the CoP specifically excludes certain material such as soils contaminated with invasive plant species with the exception of those soils that are used on the site of production in accordance with relevant guidance such as the Japanese Knotweed Code of Practice.

9 A key component of the CoP is the Materials Management Plan (MMP). It is essential that this is both adequate and based on an appropriate risk assessment that underpins the Remediation Strategy or Design Statement. As such the MMP must ensure that using the materials in the specified manner will prevent harm to human health and environmental damage. Ultimately this will need to be demonstrated in a Verification Report.

10 The key enabler is the Qualified Person (QP) who must review the relevant documents and provide a Declaration prior to the use of materials in line with the MMP. A list of authorised QPs is provided on the CL:AIRE webpage. The Declaration is initially submitted to the managing body (CL:AIRE) who will then formally submit the Declaration to the EA. When the Declaration is provided to the EA demonstrating that the materials are to be dealt with in accordance with the MMP, the EA will take the view that the materials are not waste. If the materials were not used in accordance with the MMP and underpinning risk assessment, are found not to be 'suitable for use', are to be used in 'excessive quantities' or could cause harm to human health or the environment, then the EA will view the materials as being waste and subject to regulation. The EA will undertake annual auditing of the MMP/QP submissions to ensure the scheme is being managed in the agreed manner.

11 If the material is waste an Environmental Permit will be required to lawfully deposit or re-use it unless the material is "uncontaminated soil and other naturally occurring material excavated in the course of construction activities where it is certain that the material will be used for the purposes of construction in its natural state on the site from which it was excavated", which is excluded from waste regulation by the Waste Framework Directive (2008).

12 Reference must also be made to the requirements under the relevant waste legislation and regulations such as: The Waste (England and Wales) Regulations 2011 which specifically require confirmation that the waste management hierarchy has been applied when transferring waste.

ANNEX D

CONCEPTUAL SITE MODEL (CSM)

Construction, Refinement and Representation

The CSM must present the source-pathway-receptor relationships clearly as pictorial/schematic (2D or 3D) supported by an appropriate written justification.

The starting point is the Outline CSM prepared at the Phase 0 LQA stage. This will normally form the basis of the Preliminary CSM prepared at the Phase 1 LQA stage which will be progressively refined through the various tiers of risk assessment as the LQA proceeds through Phase 2.

The level of detail will vary from CSM to CSM depending upon the site conditions such as the complexity of the geology and nature and likely extent of the contaminants.

Where appropriate to do so contaminants and pathways, be they potential or otherwise, can be grouped together rather than treated individually. This will help keep the model as manageable and simple as possible.

Box D1 CSM GUIDANCE

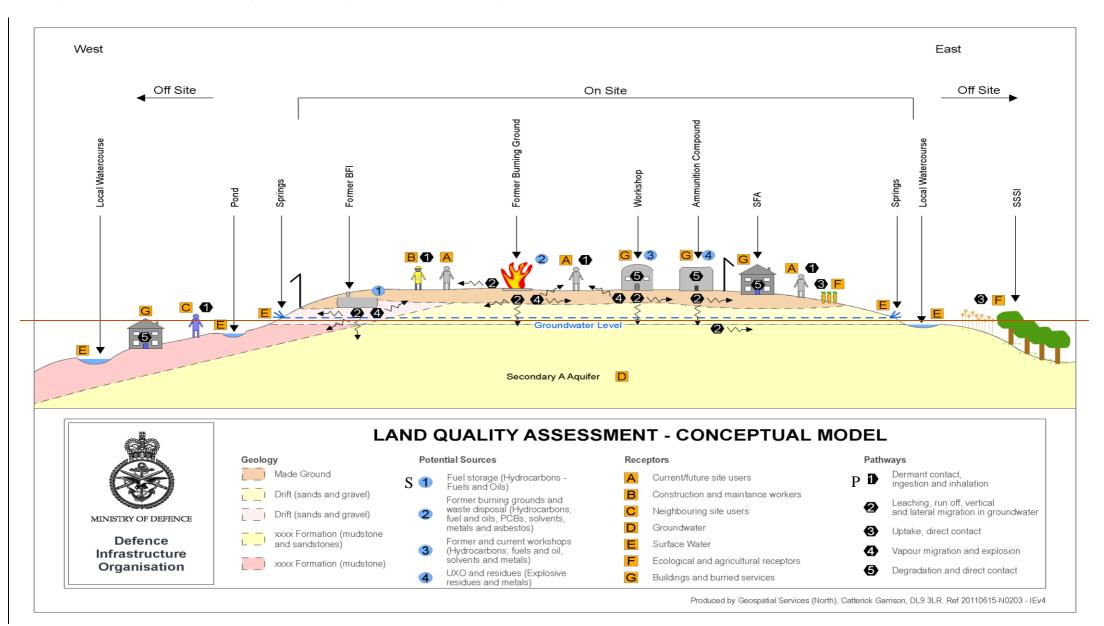
Guidance is available in:

- Environment Agency. *Guidance on good practice for the development of Conceptual Models and the Selection and Application of Mathematical Models of Contaminant Transport Processes in the Subsurface.* National Groundwater and Contaminated Land centre Report NC/99/38/2, 2001; and
- <u>R&D 66</u> Guidance on the Safe Development of Housing on Land Affected by Contamination, 2008 Volume 1 and 2. NHBC and EA.

Schematic Representation

In order to present the CSM clearly and to set the identified pollutant linkages in context in terms of the area of land being assessed it is necessary to produce a site plan illustrating areas of potential concern. A schematic cross-section representation will be required where the distribution and possible migration of contaminants in the sub-surface needs to be illustrated. An example of a CSM schematic representation is shown overleaf by means of illustration.

Example - Detailed Schematic Representation (Phase 1 LQA onwards)



Refinement and Written Justification

From the construction of the Outline CSM through the development of the Preliminary CSM and subsequent refinement it is essential that the inclusion, omission and discounting of plausible contaminants, pathways and receptors are documented and justified with reference to the relevant evidence together with any changes. This provides the necessary audit trail and allows those reviewing the model to understand the thought process and rationale.

Assessing viable/plausible source-pathway-receptor linkages requires the application of both common sense and general scientific knowledge about the nature of a particular contaminant, including how it may move or be transported, the circumstances of the land in question (e.g. geology, hydrogeology etc) and the behaviour of certain receptor types on the site (primarily applicable to humans and other living organisms).

Only 'plausible' sources, pathways and receptors should be considered. By 'plausible' we mean 'more likely than not'.

CSM Considerations:

The following list is not exhaustive and should be taken as a guide.

Site Environmental Context

- Site boundary, layout and topography;
- Geology/ground conditions;
- Hydrogeology; aquifer designation, distance from source protection zone/flow direction, etc;
- Hydrology; surface water drainage;
- Ecology; and
- Land-Use: Identification of current land uses.

Contaminant (Source) Information

- Contaminant location, potential and known;
- Contaminant Types: Identification of contaminants of concern based upon site history;
- Contaminant Properties: Physical properties of contaminants such as solubility, density, viscosity etc;
- Contaminant Form: Solid phase (particles), sorbed phase (bound to soil), free phase (NAPL's), vapour phase (in soil & air) and dissolved phase (in groundwater & pore water);
- Contaminant Distribution: Point sources (pipes and tanks), diffuse sources (stack emissions and land-spreading), possible lateral extent, concentration and depth profiles. Complicating Effects: De-commissioning, redevelopment and partial remediation;
- Potential hazard posed by contaminants phytotoxicity etc;
- Cause of the contamination;
- Likely mobility of contaminants and factors that affect this; and
- Uncertainties and assumptions.

Receptor Information

- **Humans:** Site workers, visitors and whether children or adults as this influences the exposure averaging;
- Areas (based upon receptor behaviour);
- **Ecosystems:** Habitat and ecosystem descriptions, species composition, temporal trends & animal and plant distributions;
- **Property (Flora & Fauna):** Identification of crops, domestic produce, livestock, owned or domesticated animals, wild animals subject to shooting or fishing rights and protected species;
- **Property (Buildings):** Buildings (including constituent material types) and ancient monuments; and
- **Controlled Waters:** Identification of coastal waters, inland freshwaters, ponds, lakes, rivers, watercourses and groundwater.

Pathway Information

- **Direct Exposure:** Direct with the contaminant (dermal, plant roots, building materials, etc.), direct ingestion and inhalation of vapours or dust in air; and
- Indirect Exposure: Ingestion of contaminated foods, migration into controlled waters.

Source-Pathway-Receptor Linkages

- Identify plausible linkages and place in context (i.e. identified on other sites);
- Consider existing risk management measures; and
- Consider relevant data indicating the presence of absence of pathways

ANNEX E

PRELIMINARY QUALITATIVE RISK ASSESSMENT (PRA)

The PRA is underpinned by the Preliminary CSM which identifies the hazards (source of contamination) and sets out the potential pollutant linkages with a view to identifying the nature and magnitude of the potential risks to receptors.

This requires consideration of the probability or likelihood of the linkage occurring and the severity/significance of the potential consequence taking into account the nature of the pollutant linkage and the potential severity of the hazard coupled with the sensitivity of the receptor within the context of the current and/or envisaged land use.

Ultimately, the absence of a pollutant linkage means there is no risk. That said a view will need to be taken on whether there is sufficient data to provide the necessary confidence.

A classification of: consequence/severity, probability/likelihood and risk together with definitions are presented in the various tables within this annex. These tables were developed initially in 1995 in line with the DoE Guide to Risk Assessment and Risk Management for Environmental Protection published that year and has been updated into the DEFRA 2011 Guide to Risk Assessment and Risk Management Green Leaves III_to take account of the following guidance:

- CIRIA C552 Contaminated Land Risk Assessment, A Guide to Good Practice, 2001
- DEFRA Statutory Guidance on Contaminated Land, 2006
- EA R&D Publication 66, Guidance for the Safe Development of Housing on Land Affected by Contamination, 2008

Most recently R&D 66 introduced the consideration of the 'immediacy of hazards'.

The tables provide a logical and consistent framework for assessing the potential risk by defining the categories of consequence severity, probability/likelihood of occurrence and levels of risk also referred to as 'risk terms' which follows current best practice.

The first step is to establish the consequence/severity (Table 4) and probability/likelihood (Table 5) before combining/comparing them to establish the risk category or term (Table 6). The resultant risk class is defined in Table 7.

It is worth noting that the classification of the consequence (severity) does not take account of the probability (likelihood) of that consequence being realised. Hence a 'severe' consequence refers to acute (short term) risk and a 'medium' consequence refers to chronic (long term) risk as would be the case of carcinogens and asbestos etc. Both can be classed as SPOSH and ultimately result in death. Therefore, only those contaminants that pose an acute risk to human health i.e. exposure duration of less than 24 hours should be classed as severe. Similarly contaminants that result in temporary health impacts that are non fatal should be classed as 'minor' in consequence. Care must therefore be taken and due consideration given to acute versus chronic risks otherwise the severity may be over estimated.

There is also a need to classify the liability (Table 8) to inform management decisions and the priority/urgency with which action is required.

When complete the results should be summarised in Table 3. Table 9 provides guidance on classifying the consequence/severity for sites affected by radioactively contaminated land. Though, it must be remembered that this table was prepared for nuclear licensed sites and so must be used with care.

TABLE 3 LQA RISK ASSESSMENT SUMMARY

Area / Building (Source URN No.)	Potential Contaminant	Potential Receptor	Potential Pathway to Receptor	Associated Hazard	Potential Consequence of Hazard-Receptor Link	Likelihood of Hazard- Receptor Linkage	Potential Significance	Comments / Justification
		(specific to contaminant)	(specific to contaminant)	(specific to contaminant)	e.g. Severe Moderate Mild Negligible	e.g. Certain Almost Certain Likely Possible Unlikely Nil Chance	Risk : Very High Risk High Risk Moderate Risk Low Risk Negligible/Negligible Risk No Potential Risk For use in Technical Note only: Liability Classification: A B C D E F	

Classifications are defined on the accompanying Table

TABLE 4 CLASSIFICATION OF CONSEQUENCES

TABLE 5 CLASSIFICATION OF PROBABILITY

(Only applies if there is a possibility of a pollutant linkage being present)

		[
Classification	Definition		Classification	Definition	Probability
Severe	 Acute risks to human health. Short-term risk of pollution of controlled waters or significant impact on controlled waters e.g. large scale pollution or very high levels of contamination equivalent to EA category 1 pollution incident including persistent and/or extensive effects on water quality; leading to closure of a potable abstraction point; major impact on operational effectiveness and/or amenity value or major damage to agriculture or commerce. Catastrophic damage to buildings or property (e.g. explosion causing building collapse). Ecological system effects – Immediate risks of major damage which is likely to result in: irreversible substantial adverse changes in the functioning of the ecosystem or harm to a species of special interest that endangers the long-term maintenance of the population. 		High likelihood	There is a pollutant linkage and an event is High Likelihood to occur in the short term, and is almost inevitable over the long term OR there is evidence at the receptor of harm or pollution.	>95% Likelihood of Consequence Occurring
Medium	 Chronic risks to human health. Pollution of sensitive water resources (e.g. leaching of contaminants into controlled waters) that is the equivalent of an EA Category 2 pollution incident including significant effect on water quality; notification required to abstractors; reduction in amenity value or significant damage to site operations, agriculture or commerce. Ecological system effects – Immediate risks of significant damage which may result in substantial adverse changes to the ecosystem's functioning or harm to a species of special interest that may endanger the long-term maintenance of the population. Significant damage to buildings, structures and services (e.g. damage rendering a building unsafe to occupy, such as foundation damage). 		Likely	There is a pollutant linkage and it is probable than an event will occur. It is not inevitable, but possible in the short term and likely over the long term.	50 – 95% Likelihood of Consequence Occurring
Mild	 Non-permanent health effects to human health (exposure unlikely to lead to 'significant' harm). Pollution of controlled waters or non-sensitive water resources (e.g. pollution of non-classified groundwater) that is equivalent to an EA Category 3 pollution incident or short lived effect on water quality; marginal effect on operational capability, amenity value, agriculture or commerce. Minor damage to buildings, structures and services (e.g. damage rendering a building unsafe to occupy, such as foundation damage). Ecological systems effects – Minor or short term damage which is unlikely to result in substantial adverse changes to the ecosystem's functioning or harm to a species of special interest that may endanger the long-term maintenance of the population Substantial damage to non-sensitive environments (unprotected ecosystems e.g. crops). 		Low likelihood	There is a pollutant linkage and circumstances are possible under which an event could occur. It is by no means certain that even over a longer period such an event would take place, and less likely in the short term.	5 – 49% Likelihood of Consequence Occurring
Minor / Negligible	 No measurable effects on human health including non-permanent health effects to human health that are easily prevented by appropriate use of PPE etc. Minor pollution of controlled waters including non-sensitive water resources with no discernable effect on water quality or ecosystems. Minor damage to non-sensitive environments (unprotected ecosystems e.g. crops). Easily repairable effects of damage to buildings, structures, services or the environment (e.g. discoloration of concrete, loss of plants in a landscaping scheme). 		Unlikely	There is a pollutant linkage and it is improbable that an event would occur even in the very long term.	<5% Likelihood of Consequence Occurring

These tables do not indicate direct correlation between the classification systems shown.

TABLE 6 CLASSIFICATION OF RISK (SIGNIFICANCE)

		Consequence					
Probability (likelihood)		Severe	Medium	Mild	Minor		
lid of	High Likelihood	Very High Risk	High Risk	Moderate Risk	Moderate/Low Risk		
elil	Likely	High Risk	Moderate Risk	Moderate/Low Risk	Low Risk		
r ≣	Low likelihood	Moderate Risk	Moderate/Low Risk	Low Risk	Negligible Risk		
	Unlikely	Moderate/Low Risk	Low Risk	Negligible Risk	Negligible Risk		

TABLE 7 DEFINITIONS OF CLASSIFIED RISKS/RISK TERMS

Classification	Definition
Very High Risk	Severe harm to a receptor may already be occurring OR a high likelihood that severe harm will arise to a receptor, unless immediate remedial works / mitigation measures are undertaken. Realisation of that risk is likely to present a substantial liability to MOD
High Risk	Harm is likely to arise to a receptor, and is likely to be severe, unless appropriate remedial actions / mitigation measures are undertaken. Remedial works may be required in the short term, but likely to be required over the long term. Realisation of that risk is likely to present a substantial liability to MOD
Moderate Risk	Possible that harm could arise to a receptor, but low likelihood that such harm would be severe. Harm is likely to be mild. Some remedial works may be required in the long term. Realisation of that risk is unlikely to present a substantial liability to MOD, but further work may be required to determine whether this is the case
Moderate/Low Risk	Possible that harm could arise to a receptor, but where a combination of likelihood and consequence results in a risk that is above low, but is not of sufficient concern to be classified as mild. It can be driven by cases where there is an acute risk which carries a severe consequence, but where the exposure is unlikely. Such harm would at worse normally be mild. Unlikely to present a substantial liability to MOD. Limited further investigation may be required to clarify the risk and liability. If necessary remediation works likely to be limited in extent.
Low Risk	Possible that harm could arise to a receptor. Such harm would at worse normally be mild.
Negligible Risk	Low likelihood that harm could arise to a receptor. Such harm unlikely to be any worse than mild. No liability.
No Potential Risk	There is no potential risk where no pollutant linkage has been established. No liability.

TABLE 8 LIABILITY CLASSIFICATIONS

Classification	Definition
A	Immediate risk of severe harm to human health. Requires the potential presence of significantly elevated concentrations of contaminants with high acute toxicity, sufficient to present the significant possibility of significant harm to human health, following short-term exposure.
В	Immediate risk of severe harm to the natural or built environment. E.g. a large fuel spill (or the imminent risk of such an event), the risk of explosion causing building collapse, or the possibility of irreversible adverse changes to a protected ecosystem.

A1	Health hazard to workforce during demolition or construction works. The potential for health hazards to workers involved in demolition or construction projects on site, arising from the potential presence of contaminants.
B1	Risks to the natural and built environment during demolition or construction works.

с	Large remediation liability. Remedial works will be required; large financial liability.
D	Minor remediation liability. <i>Remedial works will be required; minor financial liability.</i>

E	No remediation required. Potential contaminant source identified, but no risks ² under current site conditions, due to the lack of a pathway and/or a receptor. NB future changes to land use and/or receptor behaviour may change the liability classification.
F	No effect on re-use option or site value. Potential contamination sources may be present or have been identified, however, site is suitable for all potential end-uses, although contaminant concentrations may exceed natural background concentrations.

More than one liability classification letter may be used if appropriate, e.g. A, D.

 $^{^{2}}$ Risks under current site use deemed sufficiently low that remedial works are not considered to be necessary at this time.

TABLE 9 DESCRIPTORS FOR 'POTENTIAL SEVERITY OF CONSEQUENCE' – RADIOACTIVELY CONTAMINATED LAND

The summary table below provides guidance specific to assessing the severity and risk classification for radioactively contaminated land associated with nuclear licensed sites and so should be used with care.

	Inconsequential	Negligible	Mild	Moderate	Severe
Radiation Dose to Public	Not Distinguishable from Negligible.	Less than 0.01 mSv y ⁻¹ , if exposure occurs. This level corresponds to a risk of death of 10^6 y ⁻¹ , as defined by the BSS Direction 2000, and is not subject to any regulatory controls. May be demonstrated using GRACs for scenarios applicable to site use for 0.01 mSv y ⁻¹	Of order 0.01 - 0.1 mSv y ⁻¹ , if exposure occurs. May be demonstrate using GRACs for scenarios applicable to site use for 0.1 mSv y ⁻¹	Of order 0.1 - 1 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the legal limit for effective dose in a calendar year for any member of the public from sources of ionising radiation originating from a nuclear licensed site. This range is of a similar order to the dose constraint of 0.3 mSv y ⁻¹ in EA Briefing Note 3 (2006) and recommended by HPA (Documents of NRPB 9(2), 1988) for development of land under Planning legislation. May be demonstrated using GRACs for scenarios applicable to site use for 1 mSv y ⁻¹ .	Of order > 1 mSv y '1, if exposure occurs. This level of dose exceeds the legal limit for effective dose in a calendar year for members of the public from sources of ionising radiation originating from a nuclear licensed site. This is of a similar order to the 3 mSv y' ¹ criterion for determination of 'radioactive contamination land' not on a nuclear licensed site under Part 2A. May be demonstrated using the Part 2A criteria for 'radioactive contaminated land' or GRACs for scenarios applicable to site use of order 1 mSv y' ¹ .
Radiation Dose to on-Site 'General Employees'	Less than 0.01 mSv y ⁻¹ , if exposure occurs. This level corresponds to a risk of 10^{-6} y ⁻¹ , as defined by the BSS Directive 2000, and is not subject to any regulatory controls.	Of order 0.01 - 0.1 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the Basic Safety Objective for "other employees" working on nuclear licensed site - Target 1 in HSE SAPs.	Of order 0.1 - 2 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the Basic Safety Level target for "other employees" working on nuclear licensed site - Target 1 in HSE SAPs.	Of order 2 - 10 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the Basic Safety Level target for average effective dose in a calendar year to defined groups of "employees working with ionising radiation" on nuclear licensed site - Target 2 in HSE SAPs.	Of order > 10 mSv y ⁻¹ , if exposure occurs. This exceeds the Basic Safety Level target for average effective dose in a calendar year to defined groups of "employees working with ionising radiation" on a nuclear licensed site - Target 2 in HSE SAPs.
Radiation Dose to on-Site 'Employees Working with Ionising Radiation'	Less than 0.1 mSv y ⁻¹ , if exposure occurs. This level corresponds to the Basic Safety Objective for 'other employees' working on a nuclear licensed site – Target 1 in HSE SAPs.	Of order 0.1 - 1 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the Basic Safety Objective for "employees working with ionising radiation" on nuclear licensed site - Target 1 in HSE SAPs.	Of order 1 - 10 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the Basic Safety Level target for average effective dose in a calendar year to defined groups of "employees working with ionising radiation" on nuclear licensed site - Target 2 in HSE SAPs.	Of order 10 - 20 mSv y ⁻¹ , if exposure occurs. The upper level corresponds to the Basic Safety Legal Level Limit for average effective dose in a calendar year to "employees working with ionising radiation" on nuclear licensed site - Target 2 in HSE SAPs.	Of order > 20 mSv y ⁻¹ , if exposure occurs. This level exceeds the Basic Safety Level Legal Limit for average effective dose in a calendar year to "employees working with ionising radiation" on nuclear licensed site - Target 2 in HSE SAPs.

TABLE 9 (CONTINUED) DESCRIPTORS FOR 'POTENTIAL SEVERITY OF CONSEQUENCE' (CONTINUED)

	Inconsequential	Negligible	Mild	Moderate	Severe	
Harm to Humans (Health Risks from Non- Radioactive	No acceptable risk to human health May be demonstrated by being much less than GACs (SGVs,	Minimal risk to human health, if exposure occurs, and no perceptible nuisance.	Minimal risk to human health, if exposure occurs, and no perceptible nuisance. (e.g. odour from VOC).	Non-permanent (reversible) health effects to humans, if	"Significant Harm" as defined for Part 2A is certain, if exposure occurs (Death, disease, serious injury, genetic	
Contamination)	LQA/CIEH etc.) or indistinguishable from background.	May be demonstrated by non- exceedance of GACs (SGVs, LQM/CIEH etc).	Could be compatible with some minor exceedances of GACs (SGVs, LQM/CIEH etc).	exposure occurs.	mutation, birth defects or impairment of reproductive functions).	
Harm to Flora and Fauna (Rad and/or Non-Rad)	No significant changes to population densities in the environment or in any ecosystem.	Some change to population densities but with no negative effects on the function of the ecosystem.	A change to population densities of non-sensitive species.	Irreversible adverse change in ecosystem functioning, or danger to population of a species of special interest, for a designated site.	Widespread extinctions of one or more species.	
	No noticeable effect on crop	Minor effect on a crop yield.	Noticeable effect on crop yield. Reversible impairment to the	Substantial diminution (<20%) of crop yield. Death, serious	Agricultural land taken out of production.	
Harm to Property (Rad and/or Non-Rad)	yield, no harm to domestic animals or damage to other property.	No noticeable harm to health of domestic animals. domestic animals or damage to other property. health of domestic animals. Minor damage to other property.	disease to domestic animals. Repairable damage to other property.	Serious physical damage to other property rendering it unusable for intended purpose. Any part of a building becomes		
Harm to Buildings (Rad and/or Non-Rad)	Not distinguishable from Negligible.	No noticeable or actual harm to buildings or structures.	Easily repairable effects of damage to buildings or structures.	Damage to sensitive buildings or structures.	unusable for its intended purpose, or significant impairment of a scheduled monument.	
Pollution of the Water	Concentrations of contaminants measured are	Concentrations at least ~ 10 times less than the most	Concentrations at, or just	Concentrations slightly above the most applicable water quality standard.	Concentrations well above the most applicable water quality	
Environment (Rad and/or Non-Rad)	marginally above background levels or indistinguishable from background.	restrictive potentially relevant water quality standard.	below the most applicable water quality standard.	Concentrations in the water environment that result in dose rates >400 μ Gy h ⁻¹ to aquatic organisms or >40 μ Gy h ⁻¹ to terrestrial organisms.	standard.	
Regulatory Infringement	No regulatory infringement.	No regulatory infringement.	For example, approaching the discharge limit of the authorisation.	For example, minor unauthorised discharge of radioactivity from the site occurs, of small environmental consequence.	For example, unauthorised discharge of radioactivity from the site occurs, of direct environmental consequence.	

Table taken from NDA Direct Research Portfolio Report TSG(20)0664.

ANNEX F

ANNEX F

TABLE 10 CONTAMINANT/POLLUTANT LINKAGE EVALUATION TEMPLATE

Risk Assessment	Comment assuming Likely or High Likelihood of Pollutant Linkage Occurrence	Action Required
Category 1	Site probably not suitable for present use and / or environmental setting. Contaminants probably or certainly present and probably have an unacceptable impact on identified sensitive receptors.	Urgent action required in the short term.
Category 2	Site may not be suitable for present use and / or environmental setting. Contaminants probably or likely to be present, and may have an impact on identified sensitive receptors.	Action may be needed in the short term to medium term.
Category 3	Site considered likely to be suitable for present use and / or environmental setting. Contaminants may be present but unlikely to impact sensitive receptors identified.	No immediate action needed while site remains in present use and remains undisturbed. Management options to prevent land contamination may need to be implemented in order to reduce the hazard of land contamination.
Category 4	Site considered suitable for present use and / or environmental setting. Contaminants may be present but very unlikely to have an unacceptable impact on key targets.	No action needed while site remains in present use and remains undisturbed. In general management options to prevent land contamination are likely to be sufficient, although a review of preventative measures should be undertaken periodically.