Management of Residues Containing Naturally Occurring Radioactive Material from Uranium Production and Other Activities

Specific Safety Guide
No. SSG-60
Under the terms of Article III of its Statute, the IAEA is authorized to establish or adopt standards of safety for protection of health and minimization of danger to life and property, and to provide for the application of these standards.

The publications by means of which the IAEA establishes standards are issued in the IAEA Safety Standards Series. This series covers nuclear safety, radiation safety, transport safety and waste safety. The publication categories in the series are Safety Fundamentals, Safety Requirements and Safety Guides.

Information on the IAEA’s safety standards programme is available on the IAEA Internet site https://www.iaea.org/resources/safety-standards. The site provides the texts in English of published and draft safety standards. The texts of safety standards issued in Arabic, Chinese, French, Russian and Spanish, the IAEA Safety Glossary and a status report for safety standards under development are also available. For further information, please contact the IAEA at: Vienna International Centre, PO Box 100, 1400 Vienna, Austria.

All users of IAEA safety standards are invited to inform the IAEA of experience in their use (e.g. as a basis for national regulations, for safety reviews and for training courses) for the purpose of ensuring that they continue to meet users’ needs. Information may be provided via the IAEA Internet site or by post, as above, or by email to Official.Mail@iaea.org.

The IAEA provides for the application of the standards and, under the terms of Articles III and VIII.C of its Statute, makes available and fosters the exchange of information relating to peaceful nuclear activities and serves as an intermediary among its Member States for this purpose.

Reports on safety in nuclear activities are issued as Safety Reports, which provide practical examples and detailed methods that can be used in support of the safety standards. Other safety related IAEA publications are issued as Emergency Preparedness and Response publications, Radiological Assessment Reports, the International Nuclear Safety Group’s INSAG Reports, Technical Reports and TECDOCs. The IAEA also issues reports on radiological accidents, training manuals and practical manuals, and other special safety related publications.

Security related publications are issued in the IAEA Nuclear Security Series. The IAEA Nuclear Energy Series comprises informational publications to encourage and support research on, and the development and practical application of, nuclear energy for peaceful purposes. It includes reports and guides on the status of and advances in technology, and on experience, good practices and practical examples in the areas of nuclear power, the nuclear fuel cycle, radioactive waste management and decommissioning.
MANAGEMENT OF RESIDUES CONTAINING NATURALLY OCCURRING RADIOACTIVE MATERIAL FROM URANIUM PRODUCTION AND OTHER ACTIVITIES
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MANAGEMENT OF RESIDUES CONTAINING NATURALLY OCCURRING RADIOACTIVE MATERIAL FROM URANIUM PRODUCTION AND OTHER ACTIVITIES

SPECIFIC SAFETY GUIDE

INTERNATIONAL ATOMIC ENERGY AGENCY
VIENNA, 2021
The IAEA’s Statute authorizes it to “establish...standards of safety for protection of health and minimization of danger to life and property”. These are standards that the IAEA must apply to its own operations, and that States can apply through their national regulations.

The IAEA started its safety standards programme in 1958 and there have been many developments since. As Director General, I am committed to ensuring that the IAEA maintains and improves upon this integrated, comprehensive and consistent set of up to date, user friendly and fit for purpose safety standards of high quality. Their proper application in the use of nuclear science and technology should offer a high level of protection for people and the environment across the world and provide the confidence necessary to allow for the ongoing use of nuclear technology for the benefit of all.

Safety is a national responsibility underpinned by a number of international conventions. The IAEA safety standards form a basis for these legal instruments and serve as a global reference to help parties meet their obligations. While safety standards are not legally binding on Member States, they are widely applied. They have become an indispensable reference point and a common denominator for the vast majority of Member States that have adopted these standards for use in national regulations to enhance safety in nuclear power generation, research reactors and fuel cycle facilities as well as in nuclear applications in medicine, industry, agriculture and research.

The IAEA safety standards are based on the practical experience of its Member States and produced through international consensus. The involvement of the members of the Safety Standards Committees, the Nuclear Security Guidance Committee and the Commission on Safety Standards is particularly important, and I am grateful to all those who contribute their knowledge and expertise to this endeavour.

The IAEA also uses these safety standards when it assists Member States through its review missions and advisory services. This helps Member States in the application of the standards and enables valuable experience and insight to be shared. Feedback from these missions and services, and lessons identified from events and experience in the use and application of the safety standards, are taken into account during their periodic revision.

I believe the IAEA safety standards and their application make an invaluable contribution to ensuring a high level of safety in the use of nuclear technology.
I encourage all Member States to promote and apply these standards, and to work with the IAEA to uphold their quality now and in the future.
THE IAEA SAFETY STANDARDS

BACKGROUND

Radioactivity is a natural phenomenon and natural sources of radiation are features of the environment. Radiation and radioactive substances have many beneficial applications, ranging from power generation to uses in medicine, industry and agriculture. The radiation risks to workers and the public and to the environment that may arise from these applications have to be assessed and, if necessary, controlled.

Activities such as the medical uses of radiation, the operation of nuclear installations, the production, transport and use of radioactive material, and the management of radioactive waste must therefore be subject to standards of safety.

Regulating safety is a national responsibility. However, radiation risks may transcend national borders, and international cooperation serves to promote and enhance safety globally by exchanging experience and by improving capabilities to control hazards, to prevent accidents, to respond to emergencies and to mitigate any harmful consequences.

States have an obligation of diligence and duty of care, and are expected to fulfil their national and international undertakings and obligations.

International safety standards provide support for States in meeting their obligations under general principles of international law, such as those relating to environmental protection. International safety standards also promote and assure confidence in safety and facilitate international commerce and trade.

A global nuclear safety regime is in place and is being continuously improved. IAEA safety standards, which support the implementation of binding international instruments and national safety infrastructures, are a cornerstone of this global regime. The IAEA safety standards constitute a useful tool for contracting parties to assess their performance under these international conventions.

THE IAEA SAFETY STANDARDS

The status of the IAEA safety standards derives from the IAEA’s Statute, which authorizes the IAEA to establish or adopt, in consultation and, where appropriate, in collaboration with the competent organs of the United Nations and with the specialized agencies concerned, standards of safety for protection of health and minimization of danger to life and property, and to provide for their application.
With a view to ensuring the protection of people and the environment from harmful effects of ionizing radiation, the IAEA safety standards establish fundamental safety principles, requirements and measures to control the radiation exposure of people and the release of radioactive material to the environment, to restrict the likelihood of events that might lead to a loss of control over a nuclear reactor core, nuclear chain reaction, radioactive source or any other source of radiation, and to mitigate the consequences of such events if they were to occur. The standards apply to facilities and activities that give rise to radiation risks, including nuclear installations, the use of radiation and radioactive sources, the transport of radioactive material and the management of radioactive waste.

Safety measures and security measures have in common the aim of protecting human life and health and the environment. Safety measures and security measures must be designed and implemented in an integrated manner so that security measures do not compromise safety and safety measures do not compromise security.

The IAEA safety standards reflect an international consensus on what constitutes a high level of safety for protecting people and the environment from harmful effects of ionizing radiation. They are issued in the IAEA Safety Standards Series, which has three categories (see Fig. 1).

Safety Fundamentals

Safety Fundamentals present the fundamental safety objective and principles of protection and safety, and provide the basis for the safety requirements.

Safety Requirements

An integrated and consistent set of Safety Requirements establishes the requirements that must be met to ensure the protection of people and the environment, both now and in the future. The requirements are governed by the objective and principles of the Safety Fundamentals. If the requirements are not met, measures must be taken to reach or restore the required level of safety. The format and style of the requirements facilitate their use for the establishment, in a harmonized manner, of a national regulatory framework. Requirements, including numbered ‘overarching’ requirements, are expressed as ‘shall’ statements. Many requirements are not addressed to a specific party, the implication being that the appropriate parties are responsible for fulfilling them.

Safety Guides

Safety Guides provide recommendations and guidance on how to comply with the safety requirements, indicating an international consensus that it

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1 See also publications issued in the IAEA Nuclear Security Series.
is necessary to take the measures recommended (or equivalent alternative measures). The Safety Guides present international good practices, and increasingly they reflect best practices, to help users striving to achieve high levels of safety. The recommendations provided in Safety Guides are expressed as ‘should’ statements.

**APPLICATION OF THE IAEA SAFETY STANDARDS**

The principal users of safety standards in IAEA Member States are regulatory bodies and other relevant national authorities. The IAEA safety standards are also used by co-sponsoring organizations and by many organizations that design, construct and operate nuclear facilities, as well as organizations involved in the use of radiation and radioactive sources.

The IAEA safety standards are applicable, as relevant, throughout the entire lifetime of all facilities and activities — existing and new — utilized for peaceful purposes and to protective actions to reduce existing radiation risks. They can be

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**FIG. 1. The long term structure of the IAEA Safety Standards Series.**

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**Collection of Safety Guides**

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**Part 1. Governmental, Legal and Regulatory Framework for Safety**

**Part 2. Leadership and Management for Safety**

**Part 3. Radiation Protection and Safety of Radiation Sources**

**Part 4. Safety Assessment for Facilities and Activities**

**Part 5. Predisposal Management of Radioactive Waste**

**Part 6. Decommissioning and Termination of Activities**

**Part 7. Emergency Preparedness and Response**

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**FIG. 1.** The long term structure of the IAEA Safety Standards Series.
used by States as a reference for their national regulations in respect of facilities and activities.

The IAEA’s Statute makes the safety standards binding on the IAEA in relation to its own operations and also on States in relation to IAEA assisted operations.

The IAEA safety standards also form the basis for the IAEA’s safety review services, and they are used by the IAEA in support of competence building, including the development of educational curricula and training courses.

International conventions contain requirements similar to those in the IAEA safety standards and make them binding on contracting parties. The IAEA safety standards, supplemented by international conventions, industry standards and detailed national requirements, establish a consistent basis for protecting people and the environment. There will also be some special aspects of safety that need to be assessed at the national level. For example, many of the IAEA safety standards, in particular those addressing aspects of safety in planning or design, are intended to apply primarily to new facilities and activities. The requirements established in the IAEA safety standards might not be fully met at some existing facilities that were built to earlier standards. The way in which IAEA safety standards are to be applied to such facilities is a decision for individual States.

The scientific considerations underlying the IAEA safety standards provide an objective basis for decisions concerning safety; however, decision makers must also make informed judgements and must determine how best to balance the benefits of an action or an activity against the associated radiation risks and any other detrimental impacts to which it gives rise.

DEVELOPMENT PROCESS FOR THE IAEA SAFETY STANDARDS

The preparation and review of the safety standards involves the IAEA Secretariat and five Safety Standards Committees, for emergency preparedness and response (EPReSC) (as of 2016), nuclear safety (NUSSC), radiation safety (RASSC), the safety of radioactive waste (WASSC) and the safe transport of radioactive material (TRANSSC), and a Commission on Safety Standards (CSS) which oversees the IAEA safety standards programme (see Fig. 2).

All IAEA Member States may nominate experts for the Safety Standards Committees and may provide comments on draft standards. The membership of the Commission on Safety Standards is appointed by the Director General and includes senior governmental officials having responsibility for establishing national standards.

A management system has been established for the processes of planning, developing, reviewing, revising and establishing the IAEA safety standards.
It articulates the mandate of the IAEA, the vision for the future application of the safety standards, policies and strategies, and corresponding functions and responsibilities.

INTERACTION WITH OTHER INTERNATIONAL ORGANIZATIONS

The findings of the United Nations Scientific Committee on the Effects of Atomic Radiation (UNSCEAR) and the recommendations of international expert bodies, notably the International Commission on Radiological Protection (ICRP), are taken into account in developing the IAEA safety standards. Some safety standards are developed in cooperation with other bodies in the United Nations system or other specialized agencies, including the Food and Agriculture Organization of the United Nations, the United Nations Environment Programme, the International Labour Organization, the OECD Nuclear Energy Agency, the Pan American Health Organization and the World Health Organization.
INTERPRETATION OF THE TEXT

Safety related terms are to be understood as defined in the IAEA Safety Glossary (see https://www.iaea.org/resources/safety-standards/safety-glossary). Otherwise, words are used with the spellings and meanings assigned to them in the latest edition of The Concise Oxford Dictionary. For Safety Guides, the English version of the text is the authoritative version.

The background and context of each standard in the IAEA Safety Standards Series and its objective, scope and structure are explained in Section 1, Introduction, of each publication.

Material for which there is no appropriate place in the body text (e.g. material that is subsidiary to or separate from the body text, is included in support of statements in the body text, or describes methods of calculation, procedures or limits and conditions) may be presented in appendices or annexes.

An appendix, if included, is considered to form an integral part of the safety standard. Material in an appendix has the same status as the body text, and the IAEA assumes authorship of it. Annexes and footnotes to the main text, if included, are used to provide practical examples or additional information or explanation. Annexes and footnotes are not integral parts of the main text. Annex material published by the IAEA is not necessarily issued under its authorship; material under other authorship may be presented in annexes to the safety standards. Extraneous material presented in annexes is excerpted and adapted as necessary to be generally useful.
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1. INTRODUCTION

BACKGROUND

1.1. Radionuclides of natural origin are ubiquitous in the environment and in some geological formations have become sufficiently concentrated to be exploited for the purpose of uranium production. Uranium production, including mining, processing and management of radioactive residues, as either primary or secondary minerals, has long been recognized as needing regulatory control. However, significant concentrations of radionuclides of natural origin also occur in facilities and activities involving the processing of other minerals. These natural radionuclides can be present in the raw materials and in the residues from the processing of those other minerals.

1.2. Radioactive material is defined as material designated in national law or by a regulatory body as being subject to regulatory control because of its radioactivity [1]. Naturally occurring radioactive material (NORM) is defined as radioactive material containing no significant amounts of radionuclides other than naturally occurring radionuclides; the exact definition of ‘significant amounts’ would be a regulatory decision [1]. Material in which the activity concentrations of the naturally occurring radionuclides have been changed by a process is also considered NORM [1]. A NORM residue is defined as material that remains from a process and comprises or is contaminated by NORM. NORM waste is defined as NORM for which no further use is foreseen [1]. For the purpose of this Safety Guide, NORM residues and NORM waste can be in solid or liquid form and might emit radioactive gases. The term ‘NORM activity’ is used in this Safety Guide to describe those facilities and activities that involve management of NORM residues.

1.3. NORM residues can have a radiological impact on workers, the public and the environment. The fundamental safety objective established in IAEA Safety Standards Series No. SF-1, Fundamental Safety Principles [2], is “to protect people and the environment from harmful effects of ionizing radiation.” Consequently, a governmental, legal and regulatory framework, as described in IAEA Safety Standards Series No. GSR Part 1 (Rev. 1), Governmental, Legal and Regulatory Framework for Safety [3], for control of NORM residues might be necessary.

1.4. Uranium production activities have typically been subject to regulatory control, generally as part of the nuclear fuel cycle. Unlike uranium production,
the residues arising from other NORM activities (which may have been recycled, used in other applications or disposed of as waste) have not always been subject to appropriate regulatory control in the past, even though they might have contained radionuclides at levels that would now raise radiation safety concerns.

1.5. NORM residues, particularly those generated in mining and mineral processing, differ from radioactive residues generated at, for example, nuclear power plants or medical facilities. Such NORM residues can be generated in very large volumes but tend to contain radionuclides at relatively low activity concentrations. This has important implications for the management of NORM residues, including siting and engineering options. In some cases, NORM residues contain radionuclides at higher activity concentrations, but normally in smaller volumes1.

1.6. Various IAEA Safety Standards Series publications have some relevance to NORM and to NORM residues, including the following:

(a) IAEA Safety Standards Series No. GSR Part 1 (Rev. 1), Governmental, Legal and Regulatory Framework for Safety [3];
(b) IAEA Safety Standards Series No. GSR Part 3, Radiation Protection and Safety of Radiation Sources: International Basic Safety Standards [4];
(c) IAEA Safety Standards Series No. GSR Part 4 (Rev. 1), Safety Assessment for Facilities and Activities [5];
(d) IAEA Safety Standards Series No. GSR Part 5, Predisposal Management of Radioactive Waste [6];
(e) IAEA Safety Standards Series No. GSR Part 6, Decommissioning of Facilities [7];
(f) IAEA Safety Standards Series No. SSR-5, Disposal of Radioactive Waste [8];
(g) IAEA Safety Standards Series No. SSR-6 (Rev. 1), Regulations for the Safe Transport of Radioactive Material, 2018 Edition [9];
(h) IAEA Safety Standards Series No. GSR Part 7, Preparedness and Response for a Nuclear or Radiological Emergency [10];
(j) IAEA Safety Standards Series No. GSG-7, Occupational Radiation Protection [12];

1 The volume of NORM residues can range from less than one cubic metre up to millions of cubic metres.
IAEA Safety Standards Series No. GSG-9, Regulatory Control of Radioactive Discharges to the Environment [13];

IAEA Safety Standards Series No. GSG-15, Remediation Strategy and Process for Areas Affected by Past Activities or Events [14];

IAEA Safety Standards Series No. SSG-32, Protection of the Public against Exposure Indoors due to Radon and Other Natural Sources of Radiation [15];


1.7. A number of Safety Reports containing practical information on NORM residues from specific industries (work involving minerals and raw materials, oil and gas, zircon and zirconia, rare earth processing, titanium dioxide and related industries, and the phosphate industry) have also been published [17–22]. Further publications that are relevant to the management of NORM residues are listed in the bibliography.


1.9. The terms used in this Safety Guide are to be understood as defined and explained in the IAEA Safety Glossary [1].

OBJECTIVE

1.10. The objective of this Safety Guide is to provide recommendations to regulatory bodies, operating organizations, technical support organizations and other interested parties on approaches for the safe management of NORM residues arising from uranium production and other NORM activities, in accordance with a graded approach. These recommendations are aimed at meeting the relevant requirements established in GSR Part 3 [4] for the protection of people and the environment, both now and in the future.

1.11. This Safety Guide addresses the management of the radiological hazards and risks associated with various types of NORM residue. It addresses radioactive residues arising from uranium production and from other NORM activities that generate very large quantities of NORM residues, such as tailings from mining and mineral processing. This Safety Guide also addresses activities that generate comparatively small volumes of NORM residues such as sludge and scales. Though the fundamental principles of managing these hazards and risks are similar, the options for the management of this broad range of materials are necessarily quite different.

1.12. This Safety Guide covers the entire lifetime of a NORM residue management facility, including siting, construction, operation, decommissioning, closure, post-closure and a period of institutional control, as appropriate to the facility. A NORM residue management facility can be a facility for the processing, storage and/or long term management of NORM residues, including the permanent disposal of NORM waste.

1.13. This Safety Guide identifies organizational and regulatory requirements (including for exemption and clearance and for reuse and recycling). It includes recommendations on the conduct of screening assessments and, where necessary, safety assessments for facilities and activities involving NORM residues, including those facilities and activities for which a formal safety case is appropriate (e.g. the management of uranium production tailings).

1.14. This Safety Guide provides recommendations for regulatory bodies to determine which facilities and activities carrying out NORM residue management should be considered for regulatory control.

1.15. This Safety Guide is principally directed towards the management of NORM residues as a planned exposure situation (i.e. including the generation, reuse and recycling, long term management, and disposal of residues). It also applies to residues arising from the decommissioning of NORM facilities.

1.16. This Safety Guide does not address the remediation of areas contaminated by residual radioactive materials arising from past practices. The requirements for the remediation of such areas are established in GSR Part 3 [4], and further recommendations are provided in GSG-15 [14].
1.17. This Safety Guide is intended to address new facilities; however, it is also relevant to the review and upgrade of existing facilities. It might not be practical to apply all the recommendations to existing facilities; in such cases, the regulatory body should decide to what extent these recommendations should be applied. In accordance with national policies, appropriate steps should be taken to review existing facilities and, where reasonably practicable, to upgrade the provisions for protection and safety in accordance with the recommendations provided in this Safety Guide.

1.18. The radionuclides contained in NORM residues are not the only potential hazard. The chemical constituents within many NORM residues are also capable of causing harm to people and the environment, and it might be necessary to apply controls through environmental regulations or occupational health and safety regulations. These chemical constituents include heavy metals, inorganic elements (e.g. arsenic), acids and various organic compounds. The potential for such substances to cause harmful effects needs to be considered when planning the management of NORM residues. Although outside the scope of this Safety Guide, there is a particular need for regulatory bodies to take account of non-radiological hazards, which in many cases represent the primary risk to people and the environment. Achieving a consistent regulatory and integrated approach to protect against these different hazards is a challenge for regulatory bodies.

STRUCTURE

1.19. Section 2 provides an overview of NORM activities and NORM residues. Recommendations on the governmental, legal and regulatory framework for the safe management of NORM residues are provided in Section 3, and recommendations on the protection of people and the environment are provided in Section 4. Recommendations on the regulatory control process are provided in Section 5, while Section 6 provides recommendations on strategies for NORM residue management. Section 7 provides recommendations on the development of a safety case and supporting safety assessment. Section 8 addresses the full lifetime of facilities for the long term management of NORM residues, from siting through to long term institutional controls.

1.20. Three appendices and four supporting annexes complete the publication. Appendix I provides information on special considerations for managing residues from uranium production. Appendix II recommends a residue management plan for uranium production. Recommendations for a closure plan for a tailings management facility at a uranium production site are provided in Appendix III.
Annex I provides examples of NORM residues to be assessed for possible regulatory control. Annex II provides information on sampling NORM residues and determining radionuclide activity concentrations. An example of the application of the graded approach in the management of NORM residues is provided in Annex III. Annex IV provides information on the reuse and recycling of NORM residues. In addition, the bibliography provides a list of publications that are relevant to the management of NORM residues.

2. OVERVIEW OF NORM ACTIVITIES AND NORM RESIDUES


"Exposure due to natural sources is, in general, considered an existing exposure situation and is subject to the requirements [for existing exposure situations]. However, the relevant requirements…for planned exposure situations apply to:

(a) Exposure due to material in any practice…where the activity concentration in the material of any radionuclide in the uranium decay chain or the thorium decay chain is greater than 1 Bq/g or the activity concentration of 40K is greater than 10 Bq/g;
(b) Public exposure due to discharges or due to the management of radioactive waste arising from a practice involving material as specified in (a) above".3

2.2. The requirement stated in para. 2.1 does not apply to NORM residues in fertilizers, soil amendments or construction materials (or components of such) or to NORM residues that exist as residual radioactive material in the environment. In all such cases, the requirements for existing exposure situations apply, irrespective of the activity concentrations (see para. 5.1 of GSR Part 3 [4]). However, in terms of planning an activity for recycling NORM residues (including recycling residues into construction materials), the optimum protection strategy might include treating this as a planned exposure situation.

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3 The criteria in para. 3.4(a) of GSR Part 3 [4] represent (in order of magnitude terms) the upper bounds of the activity concentrations in normal soil.
2.3. In addition to uranium production, other NORM activities also generate residues that might be of regulatory concern. This includes the following industry sectors:\footnote{The list is not exhaustive. NORM residues that might be of regulatory concern can also arise from other sectors, such as the potash industry, geothermal energy use, use of deep water with a high mineral content, limestone processing and shale gas production.} [17]:

1. Extraction of rare earth elements;
2. Production and use of thorium and its compounds;
3. Production of tantalum, niobium and ferro-niobium;
4. Mining of ores other than uranium ore;
5. Production of oil and gas;
6. Manufacture of titanium dioxide pigments;
7. The phosphate industry;
8. The zircon and zirconia industries;
9. Production of tin, copper, aluminium, zinc, lead, and iron and steel;
10. Combustion of coal;

2.4. Table 1 provides a general overview of the NORM residues arising from uranium production and other industrial activities. Annex I provides more details of the typical characteristics of the NORM residues that might be of regulatory concern. Residues of different origins can vary significantly with respect to their radiological, chemical and physical characteristics. The information in Table 1 and Annex I includes the majority of industry sectors and NORM residues that need to be considered; however, NORM residues might also occur in other industrial activities that are yet to be identified.

2.5. Of the different residues generated by NORM activities, those residues that are generated in bulk amounts (of the order of millions of tonnes) represent the greatest challenge in terms of safe management. Although such residues contain radionuclides at relatively low activity concentrations, they are generated in very large volumes and contain long lived radionuclides and (often) other hazardous substances, such as heavy metals. Such bulk residues include waste rock from uranium mining, mineral process tailings, phosphogypsum, red mud from alumina processing, and fly ash.
<table>
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<tr>
<th>Industrial activity</th>
<th>Bulk amounts of residues</th>
<th>Moderate to small amounts of residues</th>
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<td>Tailings</td>
<td>Waste rock</td>
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<td>Conventional uranium production</td>
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<td>Industrial activity</td>
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<td>Manufacture of titanium dioxide pigments</td>
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<td>Phosphate and potash industries</td>
<td>Phosphogypsum</td>
<td>Thermal production</td>
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<td>Zircon and zirconia industries</td>
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<td>Production of tin, copper, aluminium, zinc, lead, iron and steel</td>
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<td>Industrial activity</td>
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<td>Tailings</td>
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<td>Combustion of coal</td>
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<td>Water treatment and geothermal energy use</td>
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</tbody>
</table>
2.6. Some residues might be of a relatively small volume but have a relatively high activity concentration, for example the following:

(a) Scales and sludge that accumulate in pipes or process vessels in oil and gas production, coal production with radium-rich inflow water, geothermal energy use and rare earth production;
(b) Anode slimes from electrowinning processes;
(c) Precipitated smelting dusts;
(d) Rare earth extraction residues (e.g. thorium hydroxide);
(e) Residues from decontamination processes;
(f) Contaminated equipment and process filters.

2.7. Plant and equipment, such as pipes, valves, process vessels, pumps and machinery, used for the handling or processing of material containing NORM can become contaminated with NORM residues, which can be a concern during the operation, and particularly during the decommissioning, of relevant facilities. These residues are often associated with scrap metals, which also require appropriate management (see Annex IV).

2.8. Liquid residues of various origin are also generated, in some cases in large volumes, including the following:

(a) Process water;
(b) Leaching fluids;
(c) Rainfall runoff (from the process plant area, residue management area, and residue and ore stockpiles);
(d) Seepage from process tailings, stockpiles and waste rock management areas;
(e) Mine water (e.g. groundwater that has entered open pits or underground mines).
3. GOVERNMENTAL, LEGAL AND REGULATORY FRAMEWORK FOR SAFETY

RESPONSIBILITIES OF THE GOVERNMENT

3.1. Requirement 1 of GSR Part 1 (Rev. 1) [3] states:

“The government shall establish a national policy and strategy for safety, the implementation of which shall be subject to a graded approach in accordance with national circumstances and with the radiation risks associated with facilities and activities, to achieve the fundamental safety objective and to apply the fundamental safety principles established in the Safety Fundamentals.”

3.2. For the safe management of NORM residues, the government should establish a policy and strategy that is appropriate to the national situation. The policy and strategy should acknowledge existing governmental, legal and regulatory frameworks; promote a graded approach to regulation; identify further industries that might need oversight; and coordinate the overall approach to the management of NORM residues. The policy and strategy should reflect, and be consistent with, the principles as set out in SF-1 [2] and the recommendations provided in Sections 4–8 of this Safety Guide.

3.3. The policy and strategy for the management of NORM residues should be consistent with the national policy and strategy for the development of activities that generate NORM residues. Together, these policies and strategies should address controls on the generation of NORM residues and encourage the reuse and recycling of NORM residues, where it is safe and appropriate to do so. Reuse and recycling of NORM residues are described further in Section 6 (i.e. as options for residue management), and more information on the application of these options is given in Annex IV.

3.4. The policy and strategy for the management of NORM residues should also take into account the national policies and strategies for safety, for management of non-radioactive waste and for radioactive waste management. States may choose to integrate key elements of the strategy for NORM residue management into their national policy, legal framework and regulatory instruments. In such cases, a separate national strategy for NORM residue management might not be necessary.
3.5. The government should consider the need for, and the extent of, public involvement and coordination among relevant governmental organizations during the development and implementation of the policy and strategy, including the establishment of a system for regulatory control. Increasing consultation with the public is a feature of the authorization process in many States; however, the responsibility for regulatory decisions remains with the regulatory body.

3.6. To enable oversight of NORM activities, the government should first identify which industries within the State process NORM and/or generate NORM residues. The government should then identify the regulatory body, or other authorities appropriate to these industries, to oversee NORM activities. If there are multiple activities or industries, there might be more than one regulatory body or authority involved.

3.7. In accordance with Requirements 3 and 4 of GSR Part 1 (Rev. 1) [3], the government is required to establish and maintain a regulatory body that is effectively independent and has the authority and sufficient resources (staff and financial) to properly oversee the safety of facilities and activities. For regulatory bodies that historically have not been involved in regulating radiation sources, this is likely to involve cooperation with other agencies or organizations with relevant radiation protection expertise.

3.8. The government should coordinate the establishment of an appropriate national inventory of significant NORM residues arising from new and existing NORM activities. Where possible, residues identified from past practices (i.e. those that need to be considered as part of the national strategy for residue management) should also be included in the inventory.

3.9. The government should establish legislation that allows the regulatory body to maintain effective oversight of NORM activities, where such legislation does not already exist. Such legislation should address the relevant requirements of GSR Part 3 [4] and should include provision for the authorization of facilities and activities and for the establishment of financial resources by the operating organization, where these are required. Financial resources are explained in more detail in Section 5.
3.10. For activities such as uranium production, effective legislation will do the following:

(a) Establish requirements and/or safety criteria for the management of residues, including for long term safety and the disposal of mill tailings and other residues as waste when no further use is foreseen;
(b) Prohibit the generation or storage of residues and waste unless these activities have been licensed by the regulatory body;
(c) Enable the regulatory body to specify conditions to be attached to licences;
(d) Make the failure to comply with licence conditions an offence subject to enforcement action;
(e) Require information and any associated fees to be provided with the licence application;
(f) Require that the operating organization prepare plans for the management of residues and waste;
(g) Require financial resources for the purposes of decommissioning, remediation, closure and institutional controls, as relevant;
(h) Require regulatory approval for significant changes to operations;
(i) Require regulatory approval before any licence is relinquished or is transferred to another party;
(j) Grant the regulatory body access to the facility to undertake inspections and measurements, as necessary.

For other NORM activities, the legislation and the regulatory effort should be commensurate with the risks and should take into account existing legislation and systems of control. In some cases, existing regulations for workplace health and safety and for environmental protection may already provide adequate protection against radiation; hence, further legislation specific to radiation protection might not be necessary for such NORM activities.

3.11. Given the range of industries concerned, it is possible that several different parts of government will have responsibilities relating to NORM activities, and it is likely that several pieces of legislation will apply. For effective and efficient regulation, it is important that responsibilities be defined and formally coordinated through instruments such as administrative agreements or memorandums of understanding between different agencies. This coordination can be achieved by one regulatory body acting for the government to coordinate regulatory oversight across multiple industries. More commonly, there will be multiple regulatory bodies. In the case of multiple regulatory bodies, it should be ensured that regulatory requirements and any authorization conditions are suitably aligned.
RESPONSIBILITIES OF THE REGULATORY BODY

3.12. For planned exposure situations involving NORM, as stated in Requirement 12 of GSR Part 3 [4], “The government or the regulatory body shall establish dose limits for occupational exposure and public exposure, and [operating organizations] shall apply these limits.”

3.13. In addition, as stated in para. 3.22(c) of GSR Part 3 [4] (footnote omitted):

“The government or the regulatory body: ...Shall establish or approve constraints on dose and on risk, as appropriate, or shall establish or approve a process for establishing such constraints, to be used in the optimization of protection and safety.”

3.14. The regulatory body is also required to oversee compliance with conditions specified in licences and to review and assess the results of inspection and enforcement activities, as appropriate, in accordance with Requirements 25, 27 and 31 of GSR Part 1 (Rev. 1) [3].

3.15. The regulatory body should establish regulations or guides for the exemption of practices and sources, for the clearance of material and for the release of sites from regulatory control, and should establish end state criteria for such sites. The regulatory body should oversee the implementation of the operating organization’s plans for decommissioning, for the management of NORM residues and waste and, where appropriate, for closure (including any institutional controls or long term monitoring) to verify that progress to meet the end state criteria is being made.

3.16. The regulatory body is also responsible for establishing guidance on the implementation of regulatory requirements and on the authorization process (see paras 2.5(9) and 4.34 of GSR Part 1 (Rev. 1) [3]). The regulatory body should also establish guidance on regulatory review and assessment, operational oversight, and oversight of the closure or decommissioning of a facility. The process for making regulatory decisions should be transparent, independent and justifiable, such that if a decision is challenged the regulatory body can explain how it was reached.

3.17. Through the implementation of regulatory criteria that are based on the established national policy, strategy and legislation, the regulatory body should identify those facilities or processes that require formal regulatory control and those for which guidance on best practice is more appropriate. A key role of the regulatory body is to identify which facilities and activities involving NORM or NORM residues are likely to be subject to the requirements of legislation and to
provide guidance to industry on the scope and application of regulations. This will then lead to the notification and assessment processes described in Section 5 of this Safety Guide. Uranium production facilities are likely to already be familiar with the regulatory framework, but for other NORM activities this guidance will be important.

3.18. The regulatory body should consider an outreach programme to communicate with operating organizations involved with NORM residues, to make these operating organizations aware of the potential need for regulation and radiation protection. The outreach programme should also encourage the sharing of data between the operating organization and the regulatory body and should include communications with workers and, where appropriate, the public.

3.19. The regulatory body should encourage the reuse and recycling of NORM residues in accordance with national laws and regulations, as appropriate (i.e. rather than these residues being managed as waste), where relevant safety criteria can be met and residues can be cleared from further regulatory control.

3.20. The regulatory body should ensure that it maintains the necessary technical expertise to evaluate processes and activities that generate and manage NORM residues.

3.21. The regulatory body should ensure that the operating organization keeps relevant records concerning any facility that generates, handles, processes or stores NORM residues, in particular where residues are held for long term management. The regulatory body should ensure that the operating organization provides it with access to the facility and to safety related information, in accordance with para. 2.13 of GSR Part 1 (Rev. 1) [3] and para. 2.45 of GSR Part 3 [4].

3.22. The regulatory body should assess the need for inspection, audit and periodic reassessment of the inventories of NORM residues and of environmental monitoring data.

RESPONSIBILITIES OF THE OPERATING ORGANIZATION

3.23. The operating organization is responsible for all aspects of safety of the NORM activity, including protection of workers, the public and the environment against any hazards associated with NORM residues throughout the lifetime of the NORM facility or activity, including decommissioning or closure.
3.24. The operating organization is required to notify the regulatory body of an intention to undertake a NORM activity, in accordance with Requirement 7 of GSR Part 3 [4]. As such, the operating organization is required to inform the regulatory body of any circumstances or changes that might increase occupational exposures or public exposures, in accordance with para. 3.14 of GSR Part 3 [4].

3.25. The operating organization is required to provide the regulatory body with access to the facility and to safety related information, in accordance with para. 2.13 of GSR Part 1 (Rev. 1) [3] and para. 2.45 of GSR Part 3 [4].

3.26. In accordance with Requirement 6 of GSR Part 1 (Rev. 1) [3], the operating organization is required to comply with all legal and regulatory requirements; for some facilities and activities that require a licence (see Section 5), these requirements will include collecting baseline data prior to site development and preparing a safety case and supporting safety assessment (see Section 7) associated with siting, design, construction, commissioning, operation, decommissioning or closure.

3.27. The operating organization is responsible for developing a plan for the management of NORM residues.

3.28. The operating organization is responsible for establishing and implementing an appropriate management system that incorporates radiation protection requirements to a degree that is commensurate with the complexity and risk of the facilities and activities relating to NORM residues. The management system should meet the requirements of IAEA Safety Standards Series No. GSR Part 2, Leadership and Management for Safety [23].

3.29. By means of design measures, procedures and processes, the operating organization should identify and implement measures to minimize the amounts of NORM residues and the amount of NORM waste. This could be achieved, for example, by increasing the efficiency of processes, or through the reuse and recycling of NORM residues.

3.30. Where applicable, the operating organization should maintain up to date plans for residue management and the decommissioning or closure of facilities, as appropriate. These plans should take account of the financial provision throughout the lifetime of the facility, including how it will meet the end state criteria, which also will provide the basis for any financial resources mechanism that is necessary.
4. PROTECTION OF PEOPLE AND THE ENVIRONMENT

GENERAL

4.1. The fundamental safety objective is to protect people and the environment from harmful effects of ionizing radiation [2]; this is to be achieved through compliance with the requirements established in GSR Part 3 [4]. Given the broad spectrum of NORM residues arising from a wide range of NORM activities, it is important that a graded approach to protection and safety in the management of NORM residues be adopted. That is, the protection measures adopted should be commensurate with the magnitude and likelihood of exposures and level of risk.

4.2. The regulatory framework for NORM residue management is based on requirements laid out in GSR Part 3 [4]. Practices or sources may be exempted from some or all of the requirements of GSR Part 3 [4]. With regard to NORM, para. I.4 of GSR Part 3 [4] states (footnote omitted):

“For radionuclides of natural origin, exemption of bulk amounts of material is necessarily considered on a case by case basis by using a dose criterion of the order of 1 mSv in a year, commensurate with typical doses due to natural background levels of radiation.”

4.3. The management of NORM residues is an example of the management of facilities and activities, as defined in GSR Part 3 [4]. Radiation protection considerations are therefore governed by the principles of justification, optimization and (for planned exposure situations) dose limitation. The justification principle should be applied to proposed new NORM activities before making modifications to processes that would affect the generation of NORM residues and during licence renewal.

4.4. In the management of NORM residues, a safety culture that encourages continuous improvement and a questioning and learning attitude to protection and safety is required to be fostered and sustained (see Requirement 12 of GSR Part 2 [23]).

4.5. Radioactive discharges to the environment from NORM facilities and activities that are subject to authorization should be controlled in accordance with a licence issued by the regulatory body. Recommendations on the regulatory control of discharges are provided in GSG-9 [13].
4.6. Radon exposures associated with the management of NORM residues should be controlled in accordance with the requirements established in GSR Part 3 [4] (e.g. see para. 3.4(c) and (d) of GSR Part 3 [4]). Recommendations on the assessment of radon (and thoron) exposures in workplaces, and on the protection of workers from such exposures, are provided in GSG-7 [12].

PLANNED EXPOSURE SITUATIONS

Occupational exposure

4.7. In general, occupational radiation protection in the management of NORM residues involves the consideration of three main exposure pathways:

(a) External exposure to radiation (primarily gamma radiation);
(b) Intakes of radionuclides directly through dust inhalation and ingestion or indirectly through ingestion of contaminated water or food;
(c) Exposure due to radon (and sometimes thoron) released from residues into the air.\(^5\)

Workers might be exposed during the generation of NORM residues; during operations to process, reuse or recycle the residues; or during the long term management of such residues. Exposure might also occur during the handling of contaminated items (e.g. pipes, equipment) and during maintenance and cleaning of facilities.

4.8. Where NORM residues are subject to regulatory control, the operating organization is required to prepare and implement a radiation protection programme (see Requirement 24 of GSR Part 3 [4]). The radiation protection programme should describe the measures taken to ensure that the protection of workers is optimized. Recommendations on the scope and content of the radiation protection programme are provided in GSG-7 [12], which also includes special considerations for mineral processing involving NORM (see paras 9.66–9.72 of GSG-7 [12]).

4.9. Occupational radiation protection in the generation of NORM residues is usually managed as part of the radiation protection programme for the overall process that is generating the residues. For example, radiation protection in the

\(^5\) The terms ‘radon’ and ‘thoron’ include not only the parent radionuclides — \(^{222}\)Rn and \(^{220}\)Rn, respectively — but also their short lived progeny.
generation and handling of uranium mill tailings will be a part of the overall radiation protection programme for the mill. In other cases, a single NORM residue might be the only material in the whole process where the concentration of radionuclides is sufficient to lead to exposures that warrant control; in such cases, the radiation protection programme will be specific to the NORM residue. For example, for a rare earth facility, the main residue of interest might be thorium hydroxide, which presents a significant radiological risk that needs to be managed through the radiation protection programme. Another example is NORM scale in oil and gas production facilities.

4.10. The radiation protection programme is required to include arrangements for the designation of controlled and supervised areas (see paras 3.88–3.92 of GSR Part 3 [4]). Controlled areas are likely to be unnecessary where only materials with low activity concentrations are handled, as is the case in many industrial activities involving NORM (see para. 3.79 of GSG-7 [12]).

**Public exposure**

4.11. As naturally occurring radionuclides are present in the environment and contribute to natural background radiation, it is important to distinguish between exposures arising as a result of NORM activities and NORM residues, and those arising from natural background sources. Establishing baseline information on natural radiation levels is therefore important (see para. 8.50(d)).

4.12. For public exposure, the dose limit is an effective dose of 1 mSv in a year (schedule III of GSR Part 3 [4]). NORM activities are also subject to the radiation protection principle of optimization, for which (for planned exposure situations) the government or the regulatory body is required to establish or approve dose constraints (see para. 3.120 of GSR Part 3 [4]).

4.13. Dose limits and dose constraints for public exposure apply both during operations involving NORM residues — such as generation, reuse or recycling, storage, or disposal of NORM residues — and after the cessation of such operations. During operation, public exposure can be assessed by monitoring

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6 For facilities and sites where NORM industries have been operating for a long time, the monitoring programme for establishing natural radiation levels may need to focus on representative locations away from the immediate vicinity of the site.

7 In special circumstances, a higher value of effective dose could apply in a single year, providing that the average effective dose over five consecutive years does not exceed 1 mSv in a year.
the radionuclides in ambient air or foodstuffs, or by monitoring the discharges and then modelling the transfer of radionuclides through the environment to estimate the subsequent intakes and doses to the public. After the cessation of operations, the end state criteria — in conjunction with institutional control, where appropriate — should ensure that public exposures are below the established dose constraint.

4.14. If several facilities and activities are located on the same site, the dose constraints for public exposure should apply to all sources of planned exposure to which a representative member of the public could be exposed, leaving an appropriate margin for foreseeable future activities at the site that could give rise to additional exposure. As described in para. 4.12, the regulatory body is required to either establish dose constraints or approve dose constraints, for example those that have been proposed by the operating organizations of the facilities and activities on the site.

4.15. There should be reasonable assurance by the operating organization that any control measures implemented will remain effective for a specified period agreed with the regulatory body and that during this period the dose constraint established or approved by the regulatory body will continue to be met.

4.16. The potential for public exposures in excess of the dose constraint due to possible future redevelopment of, or unplanned intrusion into, closed facilities for NORM residue management should be considered in the planning and design as well as in the safety assessment, and appropriate institutional controls should be planned and implemented.

EMERGENCY EXPOSURE SITUATIONS

4.17. For the management of NORM residues, there are very few, if any, credible accident scenarios that could lead to an emergency exposure situation. Consequently, arrangements for emergency preparedness and response, as described in GSR Part 7 [10], are unlikely to be required.

4.18. Engineering controls (e.g. the surface cover of a tailings dam) could fail because of natural processes (e.g. erosion), or other incidents might occur that result in the release of increased amounts of radionuclides to the environment. Such scenarios might have some radiation exposure implications; however, other

8 ‘Scenario’ is defined as a postulated or assumed set of conditions or events [1].
(non-radiological) risks will generally dominate. Due consideration should be given to the probability of failure of such controls and to the likely impact in terms of the overall integrity of the facility and any public exposure or environmental consequences. Such events, however, generally do not fall within the definition of a radiological emergency. The management of non-radiological emergencies is outside the scope of this Safety Guide (see para. 1.18).

4.19. If the results of the safety assessment demonstrate that an emergency exposure situation could occur in a NORM facility or activity, an adequate level of emergency preparedness and response is required, in accordance with the requirements established in GSR Part 7 [10]. Recommendations and guidance supporting the implementation of GSR Part 7 [10] are provided in IAEA Safety Standards Series No. GS-G-2.1, Arrangements for Preparedness for a Nuclear or Radiological Emergency [24], and IAEA Safety Standards Series No. GSG-2, Criteria for Use in Preparedness and Response for a Nuclear or Radiological Emergency [25].

4.20. In most cases, any deviations from normal operations and small scale incidents should be managed within the framework established for planned exposure situations. In the event of such circumstances, some arrangements might be needed for dealing with public concerns (e.g. the provision of information) and for the management of non-radiological hazards (e.g. chemicals) present at the site; however, the establishment of either on-site or off-site emergency plans, in accordance with GSR Part 7 [10], is not warranted.

EXISTING EXPOSURE SITUATIONS

4.21. Three categories of existing exposure situations involving NORM residues can potentially give rise to public exposures or occupational exposures (see para. 5.1 of GSR Part 3 [4]), as follows:

(a) Exposure due to contamination of areas by residual radioactive materials: This exposure can occur in contaminated areas containing NORM residues from past activities that were never subject to regulatory control or that were subject to regulatory control but not in accordance with the requirements of the safety standards. Where persons have access to sites containing residual NORM contamination, exposure can arise directly from those residues. More commonly, exposures occur in the area surrounding the site owing to radionuclides being dispersed by airborne or water-borne pathways and by the emanation of radon.
(b) Exposure due to commodities deriving from residual radioactive materials: This exposure can occur where NORM (irrespective of the activity concentration) is present in commodities, including fertilizers, soil amendments and construction materials, or as residual radioactive material in the environment.

(c) Exposure due to other natural sources: This exposure can occur where NORM is present in other materials and the activity concentration of radionuclides in either the uranium decay chain or the thorium decay chain does not exceed 1 Bq/g and the activity concentration of $^{40}$K does not exceed 10 Bq/g.

4.22. For existing exposure situations involving NORM residues in which doses are less than 1 mSv in a year, further action with respect to radiological controls would not normally be warranted. Where annual effective doses exceed 1 mSv, a protection strategy should be developed and implemented to ensure that any remedial action is justified and that protection and safety is optimized, in accordance with Requirement 48 of GSR Part 3 [4]. Recommendations on the remediation of contaminated sites from past practices are provided in GSG-15 [14].

4.23. The management of NORM residues in existing exposure situations is generally outside the scope of this Safety Guide (see para. 1.16). However, there might be circumstances where the regulatory body determines that the most appropriate protection strategy (see paras 5.4 and 5.5 of GSR Part 3 [4]) in a particular existing exposure situation is to utilize the system of regulatory control applied to planned exposure situations. This guidance is not intended to preclude such actions on the part of the regulatory body.

PROTECTION OF THE ENVIRONMENT


“Protection of the environment includes the protection and conservation of: non-human species, both animal and plant, and their biodiversity; environmental goods and services, such as the production of food and feed; resources used in agriculture, forestry, fisheries and tourism; amenities used in spiritual, cultural and recreational activities; media, such as soil, water and air; and natural processes, such as carbon, nitrogen and water cycles.”

4.25. In many cases, the standard of radiation protection to protect people from harmful effects of ionizing radiation means that specific consideration of effects in
the environment might not be necessary (see para. 1.21 of IAEA Safety Standards Series No. GSG-10, Prospective Radiological Environmental Impact Assessment for Facilities and Activities [26]). Furthermore, in many cases, protection of the environment from the non-radiological (i.e. chemical and physical) impacts (see paras 4.26, 4.27) of NORM activities is likely to dominate the decision making process. Nevertheless, there is a need to demonstrate that the environment is protected from harmful effects of ionizing radiation in situations in which NORM residues are released to the environment. The radiological environmental impact assessment should assess such impacts and, where necessary, identify additional control measures. Recommendations on assessing the radiological environmental impact for facilities and activities are provided in GSG-10 [26].

NON-RADIOLOGICAL CONSIDERATIONS

4.26. Non-radiological hazards might arise directly from toxic contaminants, such as heavy metals, or from toxic contaminants that can indirectly cause harmful effects. An example of the latter is acid-forming materials (such as sulphides), which might lead to the dispersion of otherwise relatively benign forms of toxic contaminants in the general environment. Other concerns might arise not from the NORM residues themselves but from materials associated with their generation or management. Examples of this are excessive amounts of sediment entering water bodies, having been eroded from the cover of a management facility for NORM residues, or discharge of process water or mine water with high salinity to a receiving water course. It is important that the overall planning of the management of NORM residues include a broad assessment of all the potentially harmful agents and effects likely to be involved, and that appropriate control measures be adopted.

4.27. In many cases, the non-radiological risks are of greater concern than the radiation risks. Arrangements are necessary between the involved regulatory bodies to maintain a consistent approach to the management of all hazards and to clearly assign the tasks and responsibilities of each regulatory body.
5. SYSTEM FOR REGULATORY CONTROL

GENERAL

5.1. The number of facilities involved in the processing of minerals and raw materials is very large, but experience suggests that only certain processes in some facilities can result in doses above 1 mSv in a year due to NORM [17]. The selection and application of regulatory controls should be commensurate with the associated hazards and risk. While the criteria for applying regulatory controls should be based on reasonable and prudent precautions to ensure safety, it should be recognized that an inappropriate application of regulatory controls could result in many facilities and activities being regulated without net benefit. For this reason, the concept of a graded approach is especially important in defining the scope of regulatory control. Before introducing regulatory controls for the purpose of radiation protection, the regulatory body should consider the regulations and controls that are already in place (i.e. for non-radiological purposes) and aim to integrate with these existing controls.

5.2. NORM residues arising from uranium production should always be under regulatory control. To determine the optimum regulatory approach for other NORM residues, the regulatory body should understand how, when and where natural radionuclides could occur in the NORM activities listed in Section 2. The regulatory body should therefore consider the processes, the materials and the residues in more detail — including an initial estimate of occupational exposures and public exposures — and consider the added cost associated with the regulation of residue management in comparison with the benefits achievable.

INVENTORY OF NORM FACILITIES AND ACTIVITIES FOR REGULATORY CONSIDERATION

5.3. Creating a list of the NORM activities that are potentially of regulatory concern is the first step in the regulatory control process. These activities can be identified by operating organizations and by the regulatory body. The list can be developed using the information in Section 2 and Annex I, adjusted to take account of national circumstances. The regulatory body may decide that activities other than those listed in para. 2.3 should be included in the scope of the regulations if there are indications that exposure of workers or the public cannot be disregarded in terms of radiation safety. The regulatory body should then update the list accordingly.
5.4. A detailed understanding of NORM activities is essential for the proper implementation of the graded approach. Therefore, the regulatory body should compile an inventory of the NORM facilities and activities that generate or manage NORM residues, including a description of the processes and materials, and the associated occupational exposures and public exposures. Information on sampling NORM residues and determining the radioactive content is provided in Annex II.

5.5. The list of NORM activities for consideration for regulatory control should not be limited to those activities that generate NORM residues. Attention should also be paid to activities involving reuse and recycling, disposal in landfill sites, and other long term management options.

5.6. With regard to the residues from uranium mining, especially bulk waste rock materials, the activity concentrations of radionuclides in the uranium and thorium decay chains are, in most cases, less than 1 Bq/g. As indicated in para. 4.21, the requirements for existing exposure situations normally apply to such materials; however, a safety assessment is generally considered to be mandatory for these residues from uranium mining, which are managed in accordance with the requirements for planned exposure situations.

GRADED APPROACH TO REGULATION

5.7. For NORM activities subject to the requirements for planned exposure situations, a graded approach to regulatory control is required, in accordance with Requirement 6 of GSR Part 3 [4]. As such, the application of the requirements for planned exposure situations to NORM activities needs to be commensurate with the characteristics of the NORM activity and with the magnitude and likelihood of the exposures. Where an existing formal regulatory process (e.g. licensing) is already in place for managing residues, that process should be followed. An example of the application of the graded approach to NORM residues is given in Annex III.

5.8. Important features of the graded approach in planned exposure situations are provisions for exemption and clearance based on established criteria (see schedule I of GSR Part 3 [4]), as well as the application of different levels of regulatory control, as follows:

(a) Exemption (from some or all regulatory requirements);
(b) Notification;
Authorization in the form of registration;
Authorization in the form of licensing.

5.9. The decision on whether a practice is subject to licensing or registration depends on the following:

(a) The estimated public exposures;
(b) The estimated occupational exposures;
(c) The measures that are considered necessary to prevent, limit and control releases of radioactive material and other hazardous substances (solid, liquid or gas) to the environment;
(d) The likelihood of deviations from normal operating conditions.

5.10. A stepwise and graded approach to the regulatory control of NORM residues in accordance with GSR Part 3 [4] is shown in Fig. 1. Details of relevant steps in Fig. 1 are stated sequentially in paras 5.11–5.40.

Notification

5.11. Requirement 7 of GSR Part 3 [4] states: “Any person or organization intending to operate a facility or to conduct an activity shall submit to the regulatory body a notification....” This notification should be made when it is intended to carry out the practice or when it is intended to make any modifications with implications for radiation protection. In this way, the regulatory body remains informed of operations and important changes.

5.12. An operating organization that intends to start an activity that is on the list of identified NORM activities within the State should formally inform the regulatory body of its plans and should include the following information:

(a) The type of intended activity and contact information for the operating organization;
(b) Where applicable, the name, contact details and proof of professional qualification of the radiation protection expert and any other qualified experts;
(c) The location of the facility or activity and details of the surrounding environment;
(d) The process and the processing capacity, including raw materials, discharges and the generation of solid residues;
(e) The radiological characteristics of raw materials, by-products and residues;
(f) The plan for managing NORM residues.
FIG. 1. Stepwise and graded approach to the regulatory control of naturally occurring radioactive material (NORM) residues in accordance with GSR Part 3 [4].
5.13. As stated in para. 3.7 of GSR Part 3 [4]:

“Notification alone is sufficient provided that the exposures expected to be associated with the practice or action are unlikely to exceed a small fraction, as specified by the regulatory body, of the relevant limits, and that the likelihood and magnitude of potential exposures and any other potential detrimental consequences are negligible.”

In such cases, there may be no need for any further action by the operating organization or the regulatory body. Nevertheless, the recommendations provided in this Safety Guide can still be used as guidance to encourage best practice in the management of NORM residues. Notwithstanding this, the operating organization should notify the regulatory body when there are changes that might lead to an increase of occupational exposures or public exposures.

5.14. For the recycling of NORM residues into construction materials, reference levels are required to be established by the regulatory body or other relevant authority, in accordance with Requirement 51 of GSR Part 3 [4]. In addition, manufacturers and suppliers should provide the relevant authority with information on the activity concentration of radionuclides in construction materials (see para. 4.14 of SSG-32 [15]).

5.15. If the NORM activity involves disposal of bulk amounts of residues, a screening safety assessment (see paras 5.17–5.20) is very likely to be necessary, and a further detailed safety assessment might also be necessary (see Section 7).

5.16. For other activities involving NORM residues (i.e. those not recycled into construction materials or disposed of in bulk amounts), if the activity concentration of any radionuclide in the uranium decay chain or the thorium decay chain exceeds 1 Bq/g (or 10 Bq/g for $^{40}$K), a screening assessment should be carried out for decision making on exemption from any further regulatory requirements.

**Screening safety assessment**

5.17. Upon receiving notification from the operating organization (or if the regulatory body has identified an activity belonging to the list compiled as described in paras 5.3–5.6, or if the regulatory body considers that a facility or activity
should be added to this list), the regulatory body may request that the operating organization undertake a screening assessment that includes the following:

(a) The baseline radiological conditions on the site and in the surrounding environment (see para. 4.11);
(b) Further details on the radiological characteristics of the raw materials, processed materials and residues, and where these occur in the facility;
(c) The estimated magnitude of the doses to workers and to the public arising from the NORM residues, including the impact on these estimated doses of implementing alternative options for the long term management, reuse and recycling of NORM residues;
(d) Any protection measures that have been implemented for workers and for the public.

5.18. The screening assessment should be specific to a particular facility or activity, where specific information is available, and the assessment method and the period over which it is conducted should be agreed with the regulatory body. The assessment could be based on existing information relating to the facility or activity and its processes and residue management methods. Alternatively, the assessment might be based on an agreed monitoring programme that is designed to provide more data. In some cases, the screening assessment could be based on assessments undertaken for other similar facilities, activities or processes that involve similar materials.

5.19. Possible outcomes of the screening assessment include exemption, authorization by registration\(^9\) (including a periodic review) (see paras 5.28–5.31) or authorization by licensing (see paras 5.32–5.34). If the estimated effective dose, excluding the contribution from the emanation of radon\(^10\), to workers or to the public exceeds 1 mSv in a year, a more detailed safety assessment (see paras 5.24–5.27) should be undertaken, as described in Section 7, and the facility or activity may need to be authorized.

5.20. In the event of a significant change in the process, or where external events (e.g. flooding, fire, land slippage, subsidence) have affected the facility or activity,

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\(^9\) A form of authorization for facilities and activities of low or moderate risk whereby the person or organization responsible for the facility or activity has, as appropriate, prepared and submitted a safety assessment of the facilities and equipment to the regulatory body.
\(^10\) This does not imply that control of radon is excluded. The exposure resulting from radon needs to be assessed to support regulatory decision making and supporting measures for protection and mitigation.
a new screening assessment might be necessary. The operating organization and the regulatory body should review the situation after a mutually agreed period to check whether the conclusions of the screening assessment are still valid.

**Exemption**

5.21. The regulatory body may decide that the optimum regulatory option is not to apply regulatory requirements to the operating organization. The mechanism for implementing such a decision is the granting of an exemption for some or all aspects of the facility or activity and from some or all of the regulatory requirements. As stated in para. I.1 of GSR Part 3 [4]:

“The general criteria for exemption of a practice or a source within a practice from some or all of the requirements of [GSR Part 3] are that:

(a) Radiation risks...are sufficiently low as not to warrant regulatory control...; or
(b) Regulatory control...would yield no net benefit, in that no reasonable measures for regulatory control would achieve a worthwhile return in terms of reduction of individual doses or of health risks.”

5.22. For NORM activities, the general criteria for exemption are deemed to have been met if the doses to workers and the public (as determined in the screening assessment) from the activity are 1 mSv in a year or less and the NORM activities do not pose an environmental risk. As stated in para. I.4 of GSR Part 3 [4] (footnote omitted):

“exemption of bulk amounts of material is necessarily considered on a case by case basis by using a dose criterion of the order of 1 mSv in a year, commensurate with typical doses due to natural background levels of radiation.”

5.23. Specific exemption criteria may be used for the disposal of residues in small quantities or where several radiation facilities and activities are located at the same site. In granting an exemption, the regulatory body may choose to exempt the operating organization from some or all of the regulatory requirements, including liability. The regulatory body should choose to grant a partial exemption in cases where certain specific control measures are considered to achieve a net benefit.
Safety assessment

5.24. If the screening assessment indicates that doses might exceed 1 mSv in a year, a more detailed safety assessment should be conducted for a period of time agreed with the regulatory body. As described in Section 7, this safety assessment might include the following:

(a) A detailed baseline survey of the site and its surrounding environment;
(b) The use of assumptions and exposure scenarios that are more realistic than those used in the screening assessment;
(c) The collection of more specific data to improve the estimation of the source of exposure, the exposure pathways and the resulting doses;
(d) More complex models to estimate exposures;
(e) Results of workplace measurements.

5.25. If the safety assessment demonstrates that the expected doses are of the order of 1 mSv in a year, the regulatory body can still grant a partial exemption, subject to certain conditions, such as enhanced monitoring by the operating organization or the regulatory body as well as regulatory inspections.

5.26. If the expected doses are of the order of 1 mSv in a year and have the potential to slightly exceed 1 mSv in a year, the regulatory body may authorize the practice by registration.

5.27. Where the safety assessment demonstrates that doses will exceed 1 mSv in a year, regulatory authorization incorporating further regulatory controls is needed and appropriate, and these controls should be placed on the operating organization through the granting of a licence by the regulatory body.

Registration

5.28. Registration is the appropriate form of authorization in cases where the operating organization has to meet only limited obligations to ensure that workers, the public and the environment are adequately protected. These obligations would typically involve measures to keep exposures under review and to ensure that the management of NORM residues and the impacts of discharges to the environment and on working conditions are such that protection and safety is optimized, with doses not approaching or exceeding the established dose constraints or the authorized limits for discharges.
5.29. As stated in footnote 19 to para. 3.8 of GSR Part 3 [4]:

“Typical practices that are suitable for registration are those for which: (i) safety can largely be ensured by the design of the facilities and equipment; (ii) the operating procedures are simple to follow; (iii) the training requirements for safety are minimal; and (iv) there is a history of few problems relating to safety in operations. Registration is best suited to those practices for which operations do not vary significantly.”

5.30. For NORM activities authorized by registration, a graded approach to other regulatory processes, including review, assessment and inspection of facilities and activities, should also be applied. The facility or activity will not need a complex programme for radiation protection and for managing NORM residues; instead, this programme might be integrated into the overall programme for health and safety. Such facilities and activities will require a safety assessment and a radiological environmental impact assessment; however, generic assumptions and simple calculations are likely to be more appropriate than the more complex safety assessments set out in Section 7.

5.31. For facilities and activities subject to registration, the strategies for NORM residue management set out in Section 6 and the safety considerations for long term management set out in Section 8 can be regarded as providing useful guidance on achieving best practice, but they should be implemented only to the degree appropriate to the level of risk.

**Licensing**

5.32. Licensing is the appropriate form of authorization for NORM activities in which an acceptable level of protection can only be ensured through the enforcement of more stringent measures to control radiation exposures. This is the highest level of the graded approach to regulation and is normally used for practices involving exposure to the following residues:

(a) Residues that are generated in very substantial quantities (e.g. by uranium production facilities);
(b) Low volume residues containing radionuclides with a high activity concentration;
(c) Residues that are discharged to the environment in significant quantities.

5.33. Licensed facilities and activities should undertake a radiological environmental impact assessment and ensure that the safety assessment addresses
the recommendations provided in Section 7. A specific programme for the management of NORM should be developed and fully documented and should be made available for regulatory review. Licensed facilities and activities should be subject to regular regulatory supervision.

5.34. For facilities and activities subject to licensing, the strategies for NORM residue management set out in Section 6 and the safety considerations for long term management set out in Section 8 represent the general expectations in terms of the control measures that should be implemented. The regulatory body should specify in the licence conditions the measures necessary to effectively manage the risks.

Clearance

5.35. Clearance is defined as the removal of regulatory control by the regulatory body from radioactive material or radioactive objects within notified or authorized facilities and activities [1], thus allowing the material or objects to be removed from the site without any further restrictions.

5.36. Clearance of NORM residues within notified or authorized NORM facilities and activities is dependent on the characteristics of the residues, including the activity concentration of nuclides, physical and chemical form, quantity, and potential risk.

5.37. As stated in para. I.10 of GSR Part 3 [4]:

“The general criteria for clearance are that:

(a) Radiation risks arising from the cleared material are sufficiently low as not to warrant regulatory control, and there is no appreciable likelihood of occurrence for scenarios that could lead to a failure to meet the general criterion for clearance; or
(b) Continued regulatory control of the material would yield no net benefit, in that no reasonable control measures would achieve a worthwhile return in terms of reduction of individual doses or reduction of health risks.”

5.38. In accordance with para. I.12(b) of GSR Part 3 [4], NORM residues may be cleared without further consideration, provided that the activity concentration of each radionuclide in the uranium decay chain or the thorium decay chain is below 1 Bq/g and the activity concentration of $^{40}$K is below 10 Bq/g. The clearance of
NORM residues containing activity concentrations above these values may be appropriate in certain situations, provided that the regulatory body is satisfied that future exposures from such residues will not require the reinstatement of controls.

5.39. As stated in para. I.12(c) of GSR Part 3 [4] (footnote omitted):

"Radioactive material…may be cleared without further consideration provided that: …For radionuclides of natural origin in residues that might be recycled into construction materials, or the disposal of which is liable to cause the contamination of drinking water supplies, the activity concentration in the residues does not exceed specific values derived so as to meet a dose criterion of the order of 1 mSv in a year, which is commensurate with typical doses due to natural background levels of radiation."

5.40. As stated in para. I.13 of GSR Part 3 [4]:

"Clearance may be granted by the regulatory body for specific situations, on the basis of the criteria of paras I.10 and I.11 [of GSR Part 3], with account taken of the physical or chemical form of the radioactive material, and its use or the means of its disposal65. Such clearance levels may be specified in terms of activity concentration per unit mass or activity concentration per unit surface area.

65For example, specific clearance levels may be developed for metals, for rubble from buildings and waste for disposal in landfill sites."

Therefore, specific clearance levels may be developed for scenarios and pathways specific to NORM residues. In terms of the processing of NORM and the management of NORM residues, it may be appropriate to establish a single set of levels both for exemption and clearance.

FINANCIAL PROVISIONS

5.41. The objective of the provision of financial resources is to protect the government and society from liabilities arising from the failure of the operating organization to adequately construct, operate, decommission or ensure the effective closure of a site containing NORM residues.

5.42. As described in para. 3.9, the government should establish a regulatory framework that allows the regulatory body to require financial resources from the
operating organization to cover all costs (including any extra costs incurred owing to the existence of NORM residues) associated with decommissioning or long term institutional control of a site containing NORM residues. These finances should be accessible only for the purpose of decommissioning or closure and any long term institutional control.

5.43. To determine the amount of financial resources needed, the regulatory framework should include provisions that require the operating organization to submit, prior to construction and operation of a facility, a plan that provides details of decommissioning and closure, including any long term management of NORM residues, and how the end state criteria will be achieved. The plan should include cost estimates for completing the work and should be subject to regulatory approval as a condition of commencing operations.

5.44. With regard to disposal facilities, the regulatory body should require the operating organization to establish a mechanism to ensure that adequate funds are available for closure and for any ongoing institutional control. The amount of funding needed will vary with time, as liabilities increase owing to the impact of operations and decrease with any progressive decommissioning, where applicable. Funding estimates should become more accurate as the scheduled final decommissioning and closure approaches. For many NORM residues, the liability and the financial resources should address both the radiological and the non-radiological aspects.

5.45. The regulatory framework should include the condition that the requirement for financial resources cannot be terminated without regulatory approval. Operating organizations could become insolvent at any time; therefore, the funds need to be in place prior to the creation of liabilities. The availability and assurance of financial resources should be reviewed at a frequency commensurate with the liabilities incurred by the NORM residue.

INTERESTED PARTIES

5.46. The regulatory body and the operating organization are required to consult with interested parties (see, e.g., Requirement 36 of GSR Part 1 (Rev. 1) [3], Requirement 5 of GSR Part 2 [23] and Requirement 3 of GSR Part 3 [4]). For facilities and activities involving the management of NORM residues, the regulatory body should ensure that the operating organization undertakes a consultation process with interested parties, when deemed necessary by the regulatory body. This consultation should also be consistent with the graded
approach to regulation; for activities subject to authorization, this should be a condition of licensing. Radioactive material attracts a large amount of public scrutiny, even when the associated radiation risk is low. Consultation with affected interested parties is required to be an open and inclusive process (see para. 4.67 of GSR Part 1 (Rev. 1) [3]).

5.47. The interested parties that should be involved in the consultation process are as follows:

(a) Residents and landowners;
(b) Indigenous people;
(c) Local communities economically dependent on the operation or the land impacted;
(d) Government agencies, including the regulatory body;
(e) Other interested parties.

5.48. Consultation is a valuable tool in gaining support for a project. Interested parties also need to be part of the decision making process regarding future land uses. This is an important element of setting end state criteria for sites containing radioactive residues.

5.49. A government that is setting up a new NORM regulatory framework should consider, where appropriate, undertaking a public engagement and education programme. This programme should promote awareness among the operating organizations of NORM activities and promote education and training activities for operating organizations and workers involved in NORM activities.

MANAGEMENT SYSTEM

5.50. Requirement 5 of GSR Part 3 [4] requires that protection and safety be effectively integrated into the overall management system, and para. 2.48(b) of GSR Part 3 [4] states (footnote omitted):

“the management system is designed and applied to enhance protection and safety by: ...Describing the planned and systematic actions necessary to provide adequate confidence that the requirements for protection and safety are fulfilled”.

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5.51. Requirements for the management system are established in GSR Part 2 [23]. Recommendations relevant to establishing a management system for NORM residues are provided in the following publications:

(a) IAEA Safety Standards Series, No. GS-G-3.1, Application of the Management System for Facilities and Activities [27];
(b) IAEA Safety Standards Series No. GS-G-3.3, The Management System for the Processing, Handling and Storage of Radioactive Waste [28];

5.52. With respect to facilities and activities relating to the management of NORM residues, the management system will need to address the life cycle of the residues, from their generation until their reuse, long term management or disposal, and the life cycle of the facilities, including siting, design, construction, commissioning, operation, decommissioning or closure and, as appropriate, long term institutional control.

5.53. The management system will need to address the impacts and controls identified in the safety assessment and in the radiological environmental impact assessment. Residue management plans should be established. For uranium production, these plans should cover residue management, radiation protection, environmental management, emergency preparedness and response, decommissioning and closure (as appropriate), monitoring and evaluation, engagement of interested parties, and transport of radioactive material. The recommended contents of a residue management plan and a closure plan applicable to uranium production are provided in Appendix II and Appendix III, respectively. The information in Appendix II and Appendix III might also be applicable, to some extent, to NORM residues of other origin with similar characteristics. Plans for the management of residues from other NORM facilities and activities should be developed, commensurate with the scale of the operation and nature of the risks.

5.54. As stated in Requirement 6 of GSR Part 2 [23], “The management system shall integrate its elements, including safety, health, environmental, security, quality, human-and-organizational-factor, societal and economic elements, so that safety is not compromised.”

Radiation protection should be integrated with and incorporated into management systems for quality assurance, environmental protection and workplace safety. With regard to the management of NORM residues, it is important that radiation
safety not be allowed to compromise protection from more significant workplace hazards or environmental impacts.

5.55. The management system should include measurable performance indicators for radiation protection, including for occupational exposure and public exposure, and in terms of workplace monitoring results.

5.56. As part of the management system, operational limits and conditions 11 should be developed on the basis of the following:

(a) The safety assessment and radiological environmental impact assessment;
(b) Design specifications and operating parameters and the results of commissioning tests;
(c) The key factors and components important to safety;
(d) The consequences of events following the failure of equipment;
(e) The minimum staffing level needed to operate the facility or conduct the activity safely.

5.57. The plans for the management of residues should be reviewed by the operating organization, as follows:

(a) At a frequency agreed with the regulatory body;
(b) Following modifications to the facility, the activity or the types of residue;
(c) As part of the process of periodically reviewing the safety case (see Section 7) for the facility;
(d) Following incidents or near misses 12;
(e) If there are changes in relevant regulatory requirements.

Any changes to the plans for managing NORM residues as a result of these reviews should be subject to regulatory approval.

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11 'Operational limits and conditions' are a set of rules setting forth parameter limits, the functional capability and the performance levels of equipment and personnel approved by the regulatory body for safe operation of an authorized facility [1].

12 A 'near miss' is defined as a potential significant event that could have occurred as a consequence of a sequence of actual occurrences but did not occur owing to the conditions prevailing at the time [1].
6. STRATEGIES FOR NORM RESIDUE MANAGEMENT

GENERAL

6.1. This section provides recommendations on the general approach to NORM residue management in facilities and activities for which authorization by licensing is appropriate, including the application of the graded approach to implement the requirements established in GSR Part 5 [6] for predisposal management of radioactive waste. It covers options for residue management through processing\(^{13}\), reuse and recycling, storage and retrieval, and long term management of NORM residues. Approaches to controlling the generation of NORM residues are also described. More information on the long term management of NORM residues is given in Section 8. For other facilities and activities (i.e. for which licensing is not appropriate), the recommendations in this section can also be useful in terms of continuous improvement and the application of good practices.

6.2. The steps involved in the management of NORM residues are as follows:

(a) Assessment of the potential for generating different types of residue, based on the design and operation of similar facilities;
(b) Measures to control the generation of residues;
(c) Processing (sorting, characterization, segregation and treatment);
(d) Clearance, if applicable;
(e) Reuse and recycling;
(f) Discharge to the environment;
(g) Long term management, including disposal where appropriate.

6.3. Facilities that generate NORM residues should be designed such that protection from exposures arising from the management of such residues is optimized. The design should address the principle of preventing an undue burden on future generations, for example by minimizing waste to be disposed of, minimizing the use of land, avoiding soil degradation, minimizing the use of fresh water, minimizing the project footprint and its potential impacts, and maximizing the reuse and recycling of materials, with due consideration of radiation safety issues and regulatory requirements.

\(^{13}\) ‘Processing’ is considered to be any operation that changes the characteristics of residues, including pretreatment, treatment and conditioning.
6.4. To avoid the need for long term management of residues, the options of clearance, discharge to the environment, reuse and recycling, and authorized disposal (including disposal in existing landfill sites and other waste disposal facilities) should be used to the maximum extent possible, subject to meeting relevant regulatory requirements. The segregation of NORM residues can reduce the volume of material for which long term management is necessary and, as a result, can reduce the amount of land or surface area needed for this purpose. Segregation facilitates the clearance, reuse and recycling of residues, as well as the conditioning and packaging of other NORM residues for transport and long term management off the site.

6.5. The design, construction, operation, decommissioning and/or closure of facilities for the processing, storage and disposal of residues from NORM activities should be undertaken in accordance with the management system outlined in paras 5.50–5.57. In particular, licensed facilities for the management of NORM residues should be constructed, operated, and decommissioned or closed in accordance with plans and procedures approved by the regulatory body. Appendix I describes special considerations for residues arising from uranium production.

6.6. The siting and design of the long term management facility should aim to avoid the need to relocate large quantities of residues when the NORM activities on the site cease. Siting and design are an essential part of the overall project development and should be addressed from the earliest stages of project development, as described in Section 8.

6.7. The decommissioning and/or closure of a residue management facility should be considered in all stages of the NORM activity; that is, during siting, design, construction and operation. Planning for the management of NORM residues should already have been addressed in the siting and design stage and not delayed until the decommissioning or closure stage. For example, taking measures at an early stage to reduce the migration of water-borne and airborne contamination to the surrounding environment will facilitate the subsequent management of the closure stage. During design and operation, attention should be given to the prevention and management of contamination of the plant and pieces of equipment. Consideration should also be given to potential events that might result in the unexpected spread of contamination.

6.8. Section 8 of this Safety Guide outlines the important characteristics and desirable features of the options that should be considered for the long term management of residues from NORM activities that require authorization
through licensing, including considerations in the design, construction, operation, decommissioning and closure of facilities, the release of materials from regulatory control, and the factors to be considered for institutional control of disposal facilities.

DEVELOPMENT AND IMPLEMENTATION OF A RESIDUE MANAGEMENT PLAN

6.9. A residue management plan should be developed, implemented and updated, as necessary, by the operating organization, in compliance with relevant regulatory requirements and in accordance with the operating organization’s policy and strategy for protection and safety, environmental protection, and waste management. The residue management plan should address the various streams of residues, with account taken of their respective characteristics, and address the full life cycle, from the generation of the residue until clearance, discharge, reuse and recycling, or long term management, including final disposal, as appropriate. Further information on a residue management plan for uranium production is given in Appendix II.

6.10. At the design stage of any project, the operating organization should be aware of the quantity and characteristics of all materials, radioactive and non-radioactive, and be able to identify potentially harmful characteristics. This allows for the systematic and iterative consideration of all materials and potential risk at the design stage, when it is easier to provide for proper controls and management. This design work will ultimately support the safety assessment, which in turn will support licensing and other regulatory activities.

6.11. The characterization of residues is an important factor in determining appropriate controls. Characterization helps in developing a complete understanding of the physical, chemical and radiological characteristics of the residues for classification and segregation, transport, processing, reuse and recycling, and long term management, including final disposal.

6.12. The following information should be considered in the characterization of NORM residues:

(a) Sources and quantities of NORM residues;
(b) Physical, chemical and radiological characteristics;
(c) Significant exposure pathways and exposure scenarios;
(d) Predicted radiation exposures and radiological environmental impact from the residues considered;
(e) Predicted impacts and risks from non-radiological components that might affect the radiological characteristics (e.g. the acidic nature of residue might lead to the mobilization of radionuclides);
(f) The measures that could be taken to control exposures, environmental impacts and other risks, including any measures to mitigate the consequences of accidents.

6.13. The development of a cost effective residue management plan can be complex. The process involves evaluating options for siting, design and construction, operation, management of residue streams (e.g. processing, storage, recycling), decommissioning or closure, and long term institutional control. Factors to be taken into account include benefits, costs, detriments, the national policy and strategy, and any regulatory limits and constraints. The process is also iterative, as different options are evaluated. For many NORM residues, non-radiological environmental considerations will predominate the process.

6.14. The evaluation criteria and procedures used to select the preferred options and to develop a residue management plan that will achieve the optimal balance between the considerations of regulatory requirements, national policy and strategy, costs, and site and process characteristics should be clearly defined and presented to the different interested parties, including the public.

CONTROL OF RESIDUE GENERATION

6.15. NORM facilities and activities should be designed to reduce, as far as practicable, the volume and radioactivity content of the residues and waste to be managed. This can be accomplished through the choice of appropriate processes that generate less NORM residue, and the reuse and recycling of equipment, materials and residues.

6.16. With regard to design features and operational procedures for controlling the generation of residues, the operating organization should consider the following:

(a) The selection of design options, processes and materials, construction methods, commissioning, and operating procedures that facilitate control of the generation of residues throughout the entire life cycle of the facility, including decommissioning;
The implementation of measures to avoid spills and the classification and designation of areas to prevent the spread of contamination;

Appropriate segregation of the various streams of residues to facilitate subsequent processing and reuse and recycling, where appropriate;

Methods to monitor and control the transfer of natural radionuclides to residues or products.

6.17. The quantities of residues that need long term management should be kept to the minimum practicable. Viable options for the safe reuse or recycling of NORM residues should be sought by the operating organization before designating such residues as NORM waste. Information on the reuse and recycling of NORM residues is given in Annex IV.

PROCESSING

Pretreatment

6.18. Pretreatment generally consists of the collection, characterization, segregation, chemical adjustment and decontamination of equipment contaminated with residues, including interim storage, as necessary.

6.19. The characterization step is important because it provides an opportunity to segregate residues in terms of their physical, chemical and radiological features and so facilitate the subsequent management of the residues, including treatment, storage, clearance, and reuse and recycling.

6.20. Residues should be segregated on the basis of their physical, chemical and radiological characteristics, with account taken of subsequent options for treatment and the potential for generating further (secondary) residues. Segregation should be designed and implemented to reduce the volume of residues and waste that will need long term management. Segregation should facilitate the reuse and recycling of residues. In mining and mineral processing, the segregation of non-mineralized or clean waste rock from mineralized waste rock is a pretreatment activity.

6.21. Scrap items such as pipes, valves, process vessels, pumps and machinery that have been contaminated with NORM residues should be decontaminated where practicable, in the interest of reuse and recycling.
Treatment

6.22. Treatment of NORM residues includes operations intended to improve safety by changing the characteristics of the residues. The basic treatment concepts are volume reduction, radionuclide removal and change of composition. Examples of such operations are incineration of combustible waste or compaction of dry solid waste (volume reduction); evaporation, filtration or ion exchange of liquid streams (radionuclide removal); and precipitation or flocculation of chemical species (change of composition). Often, several of these processes are used in combination to provide effective decontamination of a liquid residue stream. This might lead to further types of secondary residue to be managed (e.g. contaminated filters, spent resins, sludge).

6.23. Other options for liquid residue management include the following:

(a) Diversion of clean water away from sources of contamination;
(b) Reuse of residue water in the process or for dust suppression;
(c) Treatment to separate any solid NORM residues that are suspended in liquids;
(d) Treatment of residual liquid to make it suitable for discharge to the environment;
(e) Optimized processes to reduce the volume.

6.24. Unless the practice or source is exempt, or the residue meets the criteria established for release from regulatory control (see paras 5.35–5.40), authorization for discharges is required (see paras 3.4, 3.123, 3.124 and 3.132–3.134 of GSR Part 3 [4]; further recommendations are provided in GSG-9 [13]).

Conditioning

6.25. Conditioning of NORM residues involves operations that transform the residues into a form suitable for handling, transportation, storage and long term management, including disposal. Conditioning operations include immobilization, stabilization and packaging. Common immobilization methods include solidification of liquid residues, for example in cement. Stabilization methods can include dewatering and chemical adjustment.

6.26. Residues containing hazardous constituents that can become mobile in the environment, or constituents that can enhance the mobility of radionuclides in the environment, should be immobilized, stabilized or otherwise properly controlled. This is particularly important for large volumes of mining and processing
tailings; for stockpiles of NORM residues from processed raw materials, such as phosphogypsum and red mud; and for acid mine drainage.

6.27. Removal of excess water from tailings is important for reducing the potential for seepage of tailings liquor, for allowing the tailings to consolidate to prevent differential settlement, and for producing a firm mass for improved containment. This can be achieved by deposition in thin layers, with each section being allowed to drain and dry by evaporation before the next layer is deposited. Alternatively, the installation of a drainage system prior to or during the emplacement of tailings can produce successful results. The use of wicks driven into the tailings after emplacement has been used with limited success.

REUSE AND RECYCLING

6.28. The implementation of reuse and recycling options should be subject to suitable criteria, especially clearance criteria (including, as appropriate, clearance for specific situations; see paras 5.39 and 5.40). More information on reuse and recycling of NORM residues is given in Annex IV.

6.29. As described in para. 5.39, for radionuclides of natural origin in residues that might be recycled into construction materials, the activity concentration in the construction materials should not exceed specific values derived to meet a dose criterion of the order of 1 mSv in a year, and the aim should be to achieve an optimized activity concentration. Further recommendations on use of residues as building materials are provided in SSG-32 [15]. The reference level of about 1 mSv in a year applies to the dose received from exposure to gamma radiation from the building materials only (i.e. excluding any additional dose from $^{222}$Rn or $^{220}$Rn released from building materials into indoor air (see paras 4.17–4.27 of SSG-32 [15]).

STORAGE AND RETRIEVAL OF RESIDUES

6.30. Storage refers to the placement of the NORM residues in a facility where appropriate containment is provided and with the intention of retrieval of these residues [1]. Storage may take place between or within different residue management steps. In some cases, storage may be used to facilitate the next step in residue management — for example, to act as a buffer within and between residue management steps — or to provide time for the decay of radionuclides until authorized discharge, authorized reuse or recycling, or clearance can be
allowed. For example, some residues might be suitable for storage to allow for
decay of short lived radionuclides such as $^{210}$Po. However, for some residues,
storage might result in the ingrowth of decay products.

6.31. Storage might be appropriate for materials that are currently uneconomic
to process but that might be subsequently retrieved. In such cases, it is important
that the management plan adequately manage the risks and liabilities associated
with stockpiled residues.

OPTIONS FOR LONG TERM MANAGEMENT OF NORM RESIDUES

6.32. The preferred option for long term management will depend on the
conditions at the facility or the site where the activity is undertaken, and on the
characteristics of the ore body or the process materials, the mining or processing
operation, and the residues generated. When no future use of the NORM residues
is foreseen, the residues should be processed or otherwise prepared so as to meet
acceptance criteria for long term management established with the approval
of the regulatory body. These criteria are required to specify the radiological,
mechanical, physical, chemical and biological properties of the residues (see
para. 4.24 of GSR Part 5 [6]).

**Bulk amounts of residues**

6.33. Bulk amounts of residues represent the greatest challenge, despite their
relatively low activity concentration, because of the large volumes generated
and the presence of very long lived radionuclides and (often) other hazardous
substances, such as heavy metals, acids and alkali. Such residues include mineral
process tailings, raffinates, waste rock, phosphogypsum, red mud from alumina
processing, and fly ash from combustion of coal.

6.34. The best location for long term management facilities depends very much
on the physical quantities of the residues. Bulk amounts of residues such as
mine process tailings and phosphogypsum are often managed in a dedicated
facility at the site where they are generated. In such cases, the siting and design
of the facility is critical to effective and safe long term management. This is
described in Section 8.

6.35. The relocation of large quantities of material is an expensive option and
can affect the viability of a project. Relocating bulk amounts of NORM residues
when a site is shut down would not normally be the optimal strategy for residue
management because of the very large volumes and costs involved. In considering the relocation of bulk amounts of residues, the radiological, non-radiological and environmental impacts introduced by the relocation itself should be taken into account.

6.36. Subject to authorization by the regulatory body, some residues may be suitable for reincorporation into the environment from which they were originally removed, possibly including disposal with other residues or wastes, where allowed by the regulatory body. An example would be monazite sands being reincorporated uniformly into the remediated workings of a mineral sands extraction operation.

**Medium amounts of residues**

6.37. A possible option for medium amounts of residues that can be transported is taking such residues to existing management facilities, or co-locating the residues with other wastes, for example in landfill sites. If on-site management is still considered to be the best option, siting and design are important considerations, as described in Section 8.

**Small amounts of residues with higher activity concentrations**

6.38. Residues that arise in small amounts can be managed at off-site facilities, using a graded approach based on risk evaluation and regulatory approval. For instance, scales with a significant activity concentration are often removed by water jetting techniques; the secondary waste from this decontamination process is further treated according to regulatory requirements. Dispersion of small amounts of high activity concentration residues throughout a large volume of low activity concentration residues might not be appropriate unless this is addressed in the safety case and meets regulatory requirements.

6.39. Small amounts of unmodified residues might be sealed in suitable containers and deposited together with radioactive waste or other hazardous waste in a designated waste facility or special landfill sites, or possibly placed deep within tailings management facilities that are designed for long term management. Possible options for some liquid residues, such as those from in situ leaching of uranium, are injection into suitable geological formations and pretreatment followed by land application.
7. THE SAFETY CASE AND SAFETY ASSESSMENT FOR NORM RESIDUE MANAGEMENT

GENERAL


“Safety has to be assessed for all facilities and activities, consistent with a graded approach. Safety assessment involves the systematic analysis of normal operation and its effects, of the ways in which failures might occur and of the consequences of such failures.”


7.3. A safety case is defined as a collection of arguments and evidence in support of the safety of a facility or activity and will normally include the findings of a safety assessment and a statement of confidence in these findings [1]. Recommendations on the safety case and safety assessment specific to the predisposal management and disposal of radioactive waste are provided in IAEA Safety Standards Series No. GSG-3, The Safety Case and Safety Assessment for the Predisposal Management of Radioactive Waste [30], and in IAEA Safety Standards Series No. SSG-23, The Safety Case and Safety Assessment for the Disposal of Radioactive Waste [31].

7.4. The recommendations in this section apply to NORM residues associated with facilities and activities for which licensing is the appropriate form of authorization (i.e. where an acceptable level of protection and safety can be ensured only through the enforcement of more stringent measures to control radiation exposures). This is the highest level of regulation described in Section 5 and should be applied to those practices described in para. 5.32. For uranium production and other significant NORM facilities and activities, a safety case and a supporting safety assessment will be required, in accordance with para. 4.1 of GSR Part 4 (Rev. 1) [5].

7.5. For facilities and activities relating to long term management (including disposal) of NORM residues (see Section 8), a safety case and safety assessment
should be prepared before the facility is constructed or the activity is commenced. SSG-23 [31] provides further recommendations on the safety case and safety assessment. As stated in para. 1.9 of GSR Part 4 (Rev. 1) [5]:

“For many facilities and activities, environmental impact assessments and non-radiological risk assessments will be required before construction or implementation can commence. The assessment of these aspects will, in general, have many commonalities with the safety assessment that is carried out to address associated radiation risks. These different assessments may be combined to save resources and to increase the credibility and acceptability of their results.”

7.6. A safety assessment is required to be undertaken in conjunction with the planning and design of a proposed facility or activity (see para. 1.8 of GSR Part 4 (Rev. 1) [5]). When planning a NORM residue facility or activity, the operating organization should prepare a safety assessment that demonstrates the safety of the proposed facilities or activities and the compliance of these facilities or activities with regulatory requirements.

7.7. The safety assessment should primarily address the radiological impact on people and the environment in terms of radiation doses and radiation risks. In cases in which non-radiological risks dominate the risks, arrangements between the regulatory bodies involved are necessary to ensure a consistent approach to all hazards and to clearly assign the tasks and responsibilities of each regulatory body.

7.8. A radiological environmental impact assessment should form part of the safety assessment. GSG-10 [26] provides recommendations on a general framework for performing prospective assessments for facilities and activities to estimate the radiological impact on the public and the environment.

7.9. The key points to consider in conducting a safety assessment are as follows:

(a) A graded approach is required in terms of the scope and level of detail of the safety assessment that is carried out for different facilities and activities relating to NORM residue management (see Requirement 1 of GSR Part 4 (Rev. 1) [5]).

(b) The safety assessment is required to be carried out at the design stage or as early as possible for an existing facility or activity (see para. 4.6 of GSR Part 4 (Rev. 1) [5]). The safety assessment is required to cover the full
lifetime of the facility or activity, including decommissioning or closure and post-closure, as appropriate (see para. 1.8 of GSR Part 4 (Rev. 1) [5]).

(c) The safety assessment should identify and assess the impacts of the various streams of NORM residues through all potential exposure pathways. The effects of temporal variations (e.g. groundwater levels, diurnal radon fluctuations) should also be considered, including possible long term effects.

(d) The safety assessment is required to be documented and to show how the assessment has led to improvements in design or operation (see Requirement 20 and para. 4.15 of GSR Part 4 (Rev. 1) [5]).

(e) The safety assessment is required to be updated as necessary to reflect material changes in operation or regulatory requirements (see para. 4.6 of GSR Part 4 (Rev. 1) [5]).

7.10. The operating organization is required to use the safety assessment as an input to establish operational limits and conditions, as well as a monitoring programme and administrative controls (see Requirement 24 of GSR Part 4 (Rev. 1) [5]). The safety assessment should also inform the plan and design criteria for NORM residue management.

7.11. The safety assessment is the primary documentation for the operating organization to submit to the regulatory body when applying for an authorization for a facility or activity. Therefore, it should demonstrate compliance with regulatory requirements, with consideration of the whole life cycle of NORM residue management. An important outcome of the safety assessment is the facilitation of communication between interested parties on issues relating to the facility or activity (see also para. 5.9 of GSR Part 4 (Rev. 1) [5]).

7.12. The various stages in the lifetime of NORM residue facilities (i.e. siting, design, construction, operation, decommissioning or closure, post-closure) and NORM residue activities (i.e. residue generation, processing, reuse and recycling, storage, disposal) should be taken into account in the safety assessment.

7.13. The government should ensure that the regulatory framework (see Section 3) includes provisions for the regulatory review and approval of safety cases, in accordance with the graded approach.

SCOPE OF THE SAFETY ASSESSMENT

7.14. As noted in paras 7.26–7.30, the scope and extent of the safety assessment should be commensurate with the site specific issues relating to NORM residue
management to be addressed. The results of the initial safety assessment should be factored into the selection of the site and the design of the facility for NORM residue management. The assessment should consider all significant scenarios and exposure pathways by which workers, the public and the environment might be subject to a radiological impact. The scope and depth of the safety assessment should be sufficient to identify and evaluate relevant risk components over the lifetime of the facility or activity. The models or methods used should allow the effects of the various hazards associated with different options for NORM residue management to be compared in a consistent manner.

7.15. Both radiological and non-radiological components should be assessed to determine how to optimize protection and safety. The assessment of non-radiological impacts will also be subject to environmental protection legislation and health and safety legislation, as appropriate. While the assessment of non-radiological hazards lies outside the scope of this Safety Guide, the approaches to assessment described here might also be of use in the assessment of hazards and risks posed by non-radioactive components of NORM residues. Equally, existing systems to assess and manage environmental impacts and general health and safety (e.g. for workers) might be valuable in terms of managing radiological risks. This is especially true for NORM facilities and activities for which licensing is not considered appropriate and will allow an effective optimization of protection and safety.

7.16. The safety assessment should include aspects such as the following:

(a) A description of the site and the facility or activity, including relevant structures, systems and components and the characteristics of items important to the safety of the facility or activity.
(b) The maximum expected inventory of radioactivity in raw materials, process equipment, products and NORM residues, together with any associated acceptance criteria.
(c) A description of operations and procedures (inside and outside the facility), including the associated inventories and characteristics of residues.
(d) A description of the management system for protection and safety in relation to NORM residue management.
(e) The systematic identification of hazards for scenarios associated with operational states and accident conditions.
(f) An evaluation of different scenarios, including combinations, which might result in a failure of confinement that leads to a release of radioactive material, to eliminate from further consideration those scenarios of low likelihood or with low potential consequences.
An assessment of doses to workers and the public, including exposure due to radon and/or thoron, where applicable.

An assessment of the likelihood and potential consequences of the release of radioactive material and a comparison of the results of the assessment with regulatory limits and constraints.

The establishment of operational limits and conditions and administrative controls. If necessary, the designs for the management of NORM residues should be modified and the safety assessment should be updated.

Procedures and operational manuals for activities with significant safety implications.

A programme for periodic maintenance, inspection and testing of the plant and equipment.

A description of the monitoring and surveillance programmes.

The training programme for staff.

The emergency plan, if appropriate.

Provisions for occupational radiation protection and for protection of the public and the environment.

Provisions for decommissioning and/or closure, including financial resource requirements, if applicable.

Provisions for the involvement of interested parties.

Record keeping and quality management.

CONDUCTING A SAFETY ASSESSMENT

7.17. To address those aspects listed in para. 7.16, the safety assessment procedure should normally include the following:

Identification and definition of the context of the assessment, including the assessment criteria;

Development and justification of the operational scenarios to be assessed;

Formulation and implementation of models used to calculate radiological impacts;

An analysis of results and a comparison with the assessment criteria;

A description of any revisions of the project or processes;

A description of any reiterations of the assessment undertaken to achieve compliance with the assessment criteria and an optimized level of protection and safety.

7.18. The context for the assessment includes the purpose and scope of the assessment, the philosophy underlying the assessment, the regulatory framework,
the assessment criteria and end points, and the time frame for the assessment. As noted in para. 7.9(b), the assessment is required to cover the full lifetime of the facility, including decommissioning and/or closure and post-closure, as appropriate.

7.19. The description of the site, the facility and the operational activities should be sufficiently detailed to support the development of operational scenarios and the subsequent safety assessment of these scenarios. The scenarios should be specific, where practicable: specific site and facility features, facility specific operational arrangements, and characteristics of NORM residues should be considered and selected. The scenario should cover features, events and processes during operation, closure and post-closure. For example, any factors that affect the stability of a tailings management facility, including natural and human activities, should be sufficiently addressed. It is required that the features, events and processes considered in the safety assessment be addressed systematically (see para. 7.16) and that the identification of scenarios relevant to safety be justified (see para. 4.51 of GSR Part 4 (Rev. 1) [5]).

7.20. Once the scenarios have been developed, the corresponding assessments should be carried out, with account taken of the application of the graded approach. This is commonly undertaken using assessment models. A useful approach is a site model that considers the potential pathways by which radioactivity might move through the environment. This site model should consider the inventories of NORM residues, as well as their physical and chemical characteristics, and the location of any NORM (including raw materials, residues and waste), together with a description of any non-radiological hazards. The assessment model may be developed from one or more of the following components: specialist knowledge, conceptual site models, mathematical modelling and computer simulations. Often, specific models may need to be developed, for example to consider particular processes. For the purposes of safety assessment, any individual components need to be linked in such a way that it is possible to assess the potential radiological impacts of the facility or activity as a whole.

7.21. The safety assessment should also consider the following:

(a) The baseline concentration (see para. 8.50) and end points for the assessment, together with a justification for their selection;
(b) The timescale for the assessment;
(c) If several facilities or activities exist or are planned for the same site, the cumulative impact of all such facilities and activities;
(d) Initiating events, including internal events, external events and human induced events (see paras 4.5 and 4.22 of GSR Part 4 (Rev. 1) [5]);
(e) The use of both conservative and realistic calculations in completing the assessment;
(f) For disposal facilities, the need for any ongoing institutional control after closure and the duration of any such control;
(g) In relation to disposal facilities, the loss of institutional control after closure, including the possibility of inadvertent human intrusion;
(h) The use of sensitivity analyses and the approach to uncertainties in the safety assessment (see Requirement 17 of GSR Part 4 (Rev. 1) [5]).

7.22. Upon completion of the safety assessment, the radiation risks associated with each scenario should be quantified, screened and ranked in such a manner as to direct resources towards the most significant hazards associated with the facility or activity. Any scenarios lacking the potential to cause any significant harm to people or the environment can be removed from further consideration in the safety assessment. In the re-evaluation of a safety assessment, any such decisions should be reviewed to check that they remain valid.

7.23. The safety assessment is required to be submitted by the operating organization to the regulatory body as part of the authorization process (see para. 1.2 of GSR Part 4 (Rev. 1) [5]). The output from the safety assessment will also form part of the safety case that is required for certain facilities and activities (see Requirements 13–16 of GSR Part 5 [6]).

7.24. If the results of the safety assessment do not demonstrate compliance with regulatory requirements, the project components should be revisited and revised, as necessary, to achieve the necessary level of compliance. It is not sufficient that the calculated doses are below dose constraints; the project should be reassessed to demonstrate that protection and safety is optimized. This step should be repeated, as necessary, to “provide the highest level of safety that can reasonably be achieved throughout the lifetime of the facility or activity” (para. 3.21 of SF-1 [2]). The safety case should not be finalized until this iterative process is completed.

7.25. The operating organization should ensure that any calculations undertaken as part of the safety assessment are sufficient to enable comparisons with the assessment end points and with any additional safety or performance criteria specified by the regulatory body. Guidance on the application of the safety assessment results should be provided by the operating organization when applying for a licence. For example, it should be explained how the safety
assessment results (end points) demonstrate compliance with regulatory criteria (e.g. safety targets).

**GRADED APPROACH TO SAFETY ASSESSMENT**

7.26. It is important that a graded approach to conducting safety assessment be applied and that existing occupational health, safety and environmental control measures be taken into account. For NORM residues, in many cases very simple assumptions and calculations may be more appropriate than undertaking a detailed and complicated safety assessment. Furthermore, additional controls should be applied to the management of NORM residues only where these controls are necessary to reach an optimum level of radiation protection.

7.27. Safety assessment is a systematic process (see para. 3.15 of SF-1 [2]), and the resources devoted to safety assessment are required to be proportionate to the risks that need to be managed (see para. 3.2 of GSR Part 4 (Rev. 1) [5]). For more complex projects, the safety assessment should be iterative, with each iteration contributing to the optimization of protection and safety.

7.28. Due account needs to be taken of social and economic factors when determining the optimum level of protection and when determining the optimum level of regulatory intervention (see paras 3.23 and 3.24 of SF-1 [2]). As such, while the safety principles are the same for managing any radioactive residues, regardless of origin, there are likely to be significant differences in the practical focus of individual programmes for NORM residue management in order to optimize protection.

7.29. Paragraph 3.3 of GSR Part 4 (Rev. 1) [5] states:

“The main factor to be taken into consideration in the application of a graded approach is that the safety assessment shall be consistent with the magnitude of the possible radiation risks arising from the facility or activity. The approach also takes into account any releases of radioactive material in normal operation, the potential consequences of anticipated operational occurrences and possible accident conditions, and the possibility of the occurrence of very low probability events with potentially high consequences.”
7.30. Three aspects are to be considered in the application of a graded approach (see paras 3.3 and 3.4 of GSR Part 4 (Rev. 1) [5]):

(a) The magnitude of the possible radiation risks;
(b) The use of proven practices, procedures and designs to manage risk;
(c) The complexity of the facility or activity.

7.31. The application of the graded approach should be reassessed as the safety assessment progresses and a better understanding is obtained of the radiation risks arising from the facility or activity. The regulatory body should consider granting exemption from specific regulatory requirements if the safety assessment demonstrates that such requirements will not be effective in terms of the optimization of protection and safety.

DOCUMENTATION OF THE SAFETY CASE AND THE SAFETY ASSESSMENT

7.32. As stated in Requirement 15 of GSR Part 5 [6] in respect of predisposal radioactive waste management facilities and activities:

“The safety case and its supporting safety assessment shall be documented at a level of detail and to a quality sufficient to demonstrate safety, to support the decision at each stage and to allow for the independent review and approval of the safety case and safety assessment. The documentation shall be clearly written and shall include arguments justifying the approaches taken in the safety case on the basis of information that is traceable.”

7.33. Any assumptions made, or generic information used, in the safety case are required to be justified in the documentation (see para. 5.9 of GSR Part 5 [6]). For facilities or activities that involve long time frames, a plan for adequate record keeping over the expected project life should be provided as part of the safety case.

7.34. Some regulatory bodies might not have in-depth experience and expertise in the regulation of facilities and activities involving NORM residues. In such cases, the regulatory body may need to seek cooperation and advice from relevant expert agencies and staff when reviewing and assessing the safety case and the safety assessment.
PERIODIC SAFETY REVIEWS

7.35. The safety assessment is required to be periodically reviewed (see Requirement 24 of GSR Part 4 (Rev. 1) [5]) at predefined intervals in accordance with regulatory requirements (see para. 5.12 of GSR Part 5 [6]). In accordance with Requirement 16 of GSR Part 5 [6], the safety case and supporting safety assessment are expected to be reviewed and updated in the following circumstances:

(a) When there is any material change to the facility or activity, or a change in the radionuclide inventory that might affect safety;
(b) When changes occur to the site that might impact the facility or activity, such as encroaching industrial or municipal development;
(c) When significant changes in knowledge and understanding occur, for example from new research data or from monitoring and operating experience;
(d) When there is an emerging safety issue due to a regulatory concern or an incident;
(e) Periodically, at predefined periods, as specified by the regulatory body;
(f) When regulatory requirements change.

8. SAFETY CONSIDERATIONS FOR LONG TERM MANAGEMENT OF NORM RESIDUES

GENERAL

8.1. This section applies to facilities for the long term management of NORM residues for which authorization by licensing is appropriate, as described in Section 5. This should be applied to those practices described in para. 5.32, which include uranium production and other significant NORM facilities and activities for which a safety case and a supporting safety assessment are required, as described in Section 7.

8.2. The siting, design, construction, operation, and decommissioning and/or closure of residue management facilities should meet the requirements established by the regulatory body, including any licensing conditions, through all these phases. When residues have no foreseen further use and are neither exempted nor cleared from regulatory control, the requirements for disposal of radioactive waste
established in SSR-5 [8] should be applied in accordance with a graded approach. The requirements for decommissioning are established in GSR Part 6 [7].

8.3. The optimum location for long term management of residues depends very much on the physical quantities of the residues. Bulk amounts of residues such as waste rock and tailings of uranium mining and milling and phosphogypsum are generally managed at the site where they are generated. In selecting the site for management of bulk amounts of NORM residues, consideration should be given to the benefits of relocating and consolidating residues to limit the number of residue management sites.

8.4. The construction of a facility for managing large volumes of NORM residues, such as uranium mine or process tailings, is generally a long term project involving significant costs; therefore, any issues in terms of siting, design or construction should be identified before work begins — or as early in the process as possible — to avoid unexpected costs. Repairs or other remedial measures on completed constructions will most likely be economically prohibitive, time consuming and, in some cases, impracticable.

8.5. It is important that effective verification and quality control measures be in place during site characterization, design and construction to ensure that any engineered structures such as dams, berms, engineered liners and compacted layers meet the design specifications. The quality control programme should also involve testing of construction materials (e.g. tills, clay) to ensure that they meet the design standards and specifications.

SITING

8.6. In selecting a site for large volumes of residues, an important consideration is to minimize the dependence on active institutional controls. The final optimized choice of site, obtained using the conceptual design for residue management, should be assessed, and the resulting safety assessment, which includes the environmental impact assessment, should be submitted to the regulatory body for review and approval. The choice of the location of a facility for the management of residues should take into consideration long term stability and the need to optimize protection and safety for people and the environment for the expected lifetime of the facility during normal operation and possible accident conditions. In selecting a site, consideration should be given to features that might help control the further generation of residues, for example features that minimize the secondary contamination of environmental matrices, such as soil or seepage.
water. For uranium production and other NORM facilities, non-radiological environmental protection issues will usually dominate the decision making.

8.7. The long term management facility for bulk amounts of residues is usually near the site where the residues are generated. It is, however, essential to identify the optimum site through a stepwise site selection programme and site characterization programme (see also Requirement 15 of SSR-5 [8] in relation to site characterization for a disposal facility). A preliminary evaluation of site characteristics should be made to identify any restrictions, in terms of radiological and environmental factors, at each proposed location and to allow the selection of a small number of locations and possible preliminary design concepts for which the impacts can then be evaluated in detail.

8.8. Characterization of the site is especially important when selecting a location for the long term management of bulk amounts of residues. Understanding the site, including temporal fluctuations, before design decisions for long term management are made is very important. The site characterization information that is needed to support design decisions includes the following:

(a) Local climate and meteorology;
(b) Geography and geomorphology;
(c) Structural geology and seismology;
(d) Geochemistry (of natural and process materials);
(e) Mineralogy;
(f) Surface water and groundwater hydrology;
(g) Flora and fauna, including any protected and endangered species;
(h) Local land management;
(i) Population distribution and local land use;
(j) Archaeological and heritage issues;
(k) Socioeconomic issues.

DESIGN AND CONSTRUCTION

8.9. A long term management facility for NORM residues or NORM waste should be designed and constructed with the following objectives:

(a) To minimize water infiltration;
(b) To maintain long term stability and integrity of containment;
(c) To maximize the use of inert and stable materials as confinement barriers;
To place residues and waste below ground level to minimize the effects of potential surface erosion that could lead to the failure of the facility or accidental release of contaminated material;

To minimize the surface area impacted by the facility;

To minimize the impact on the surrounding environment during operations and after decommissioning or closure;

To minimize the potential for groundwater contamination;

To minimize the need to retrieve or relocate residues before the closure of a disposal facility;

To minimize the possibility of inadvertent intrusion;

To facilitate the implementation of surveillance, maintenance and controls during operations and, where appropriate, post-closure;

To minimize the number of residue management sites through the consolidation of residues.

8.10. The design of a long term residue management facility should follow good practice (and best practice, to the extent practicable) and meet the applicable regulatory requirements for protection and safety. Factors that should be considered in the design process include the following, as appropriate:

(a) Site characteristics (see para. 8.8);

(b) Residue characteristics including volume and chemical, physical and radiological properties;

(c) The capacity of the facility, to ensure that sufficient space will be available during operation and during decommissioning or closure (including consideration of foreseeable accident scenarios);

(d) Residue conditioning, including neutralization, precipitation, thickening and evaporation;

(e) The potential for retrieval of residues for relocation, reuse or recycling (including processing for further resource extraction);

(f) Drainage and liquids management, including seepage collection and treatment;

(g) The acid generating potential of the residues;

(h) Radiation protection measures, which might include shielding, containment, and measures to control radon and dust;

(i) Site access control and control of access to controlled areas;

(j) Results of inspections of the residues and their containment and any non-compliance issues;

(k) Ventilation of facilities, including the filtration of exhaust air discharged to the atmosphere;
The permeability of any cover and base, and the permeability criteria that are acceptable considering the site and residue characteristics, including those relating to intrusion, the leaking of liquids and the emanation of radon;

(m) Provisions for environmental monitoring, including groundwater well installations, and water and air sampling stations for effluent discharges or airborne releases;

(n) Provisions to facilitate maintenance work and eventual decommissioning and/or closure;

(o) Long term stability and erosion control (e.g. dams, berms, slopes, covers) in relation to natural weathering processes and extreme natural events (e.g. flooding, droughts, tornadoes, earthquakes);

(p) Control of inadvertent intrusion by people, plants or animals.

8.11. A detailed engineering design can be carried out after the site selection and the conceptual design have been approved by the regulatory body. At this stage, a further safety assessment, including optimization of protection, should be performed. If significant changes are made to the design of the management facilities at any stage, a further safety assessment, including optimization of protection, should be undertaken.

8.12. The detailed design should be supported by the safety assessment (see Section 7) and, where appropriate, by fieldwork and laboratory and/or pilot plant studies. The design should take account of plans for the management of residues and waste. Such plans will include, for example, the management of tailings and waste rock; proposals for effluent treatment, seepage controls and operational monitoring; and a consideration of closure and post-closure management.

8.13. A quality control programme for construction should be established at an early stage in the design process; this programme should be clearly defined and documented, and reassessed periodically. The effective implementation of a robust quality control programme involves well trained and dedicated staff. The quality control programme should specify the tests to be carried out, including the test objectives and the design criteria to be met, and any other measures that are necessary to ensure completion of the construction in accordance with the detailed design.

8.14. During the conceptual design stage for a disposal facility, a preliminary closure plan should be prepared that identifies and ranks the available options for closure according to the results of the safety assessment and the optimization of protection. The preliminary closure plan should also specify the financial resources necessary for the preferred option and take into account the post-closure land
use options. The preliminary closure plan should be submitted to the regulatory body for approval.

OPERATION

8.15. Facilities for the long term management of NORM residues or waste should be operated in accordance with the residue and/or waste management plan that was developed and modified in a manner consistent with the safety assessment and in accordance with the authorization issued by the regulatory body. This plan should describe in detail all aspects of the management of the residues or waste. The plan should be consistent with the quality assurance programme and should include provisions for the following:

(a) Detailed and documented procedures for operation, maintenance, monitoring, quality assurance, safety and, as appropriate, security;
(b) Training of personnel in the implementation of the procedures;
(c) Adequate surveillance and maintenance of all the structures, systems and components important to safety;
(d) The designation of controlled and supervised areas, as appropriate (see Requirement 24 of GSR Part 3 [4]);
(e) Procedures for the clearance of materials removed from the site;
(f) Timely submission to the regulatory body of inspection reports, monitoring results and reports on unusual occurrences;
(g) The development of emergency plans, where appropriate (see paras 4.17–4.20);
(h) The review and updating of the management plan;
(i) The regular updating of the inventory register of waste deposited.

8.16. The operating organization should ensure that the residue management plan and operating procedures are followed. The management plans should be modified and updated to take into account feedback and lessons identified from the operation of the facility. This is important for maintaining the desired level of protection and safety during operation and, where appropriate, after closure.

8.17. The regulatory body should review and approve the residue management plan and verify that operating procedures are followed by the operating organization during operation and decommissioning or closure. The regulatory body should implement a suitable system to audit and inspect the operating organization’s compliance with the approved residue management plan. If the operating
organization fails to satisfactorily follow the approved residue management plan, the regulatory body should take appropriate action to address the non-compliance.

8.18. As with other aspects of NORM residue management, the regulatory body should take a graded approach to regulatory oversight, commensurate with the scale of the risks under normal operation and from foreseeable incident scenarios.

8.19. The operating organization should take measures, on the basis of the safety assessment, to limit the release of radionuclides to the environment in liquid and airborne effluents. Measures should be taken to ensure that solid residues and waste remain under proper control so that the misuse of tailings and other NORM residues is avoided. Releases of radon or radioactive dusts into the atmosphere, and of radium and other radionuclides into surface water and groundwater by surface runoff or leaching from solid residues or waste, should be minimized.

8.20. In specific cases, a confined water covering over tailings placed in a pit may be used as a radon barrier, thereby obviating the need to perform dewatering to any significant degree. Plans for the closure of facilities that rely on water coverings should consider the placement of the tailings (above ground or below ground), the local climate and the likelihood of the water cover being passively maintained over the long term. Water covers are generally used only as temporary or interim radon barriers for residues placed above ground or, in the case of residues placed below ground, where conditions do not support a permanent water cover.

DECOMMISSIONING OF FACILITIES AND CLOSURE OF FACILITIES

8.21. Requirements for the closure of disposal facilities are established in SSR-5 [8], and requirements for decommissioning of facilities are established in GSR Part 6 [7]. Recommendations on decommissioning are provided in IAEA Safety Standards Series No. WS-G-5.1, Release of Sites from Regulatory Control on Termination of Practices [32], and IAEA Safety Standards Series No. WS-G-5.2, Safety Assessment for the Decommissioning of Facilities Using Radioactive Material [33].

8.22. When a facility for the long term management of NORM residues and waste is shut down, both decommissioning (i.e. of buildings and services used for the management of residues) and closure (i.e. of the part of the site in which waste
has been disposed) might be necessary. In such cases, the process would comprise the following steps:

(a) Design considerations and early planning;
(b) Preparation and approval of the final plans for decommissioning and for closure;
(c) Decommissioning of buildings and other structures;
(d) Management of residues and waste resulting from decommissioning activities;
(e) Closure of the disposal facility;
(f) Completion of final radiation survey;
(g) Implementation of institutional controls, if necessary;
(h) Consideration of final land use and infrastructure use.

8.23. A preliminary plan for decommissioning and/or closure should be prepared during the design phase prior to construction of the facility. The preliminary decommissioning and/or closure plan should identify and rank the available options for safely managing residues and waste according to the safety assessment and the end state criteria, with the goal of selecting a preferred option in which protection and safety is optimized. The preliminary plan should also specify the provision of the financial resources necessary for the preferred option. The preliminary plan for decommissioning and/or closure should be subject to regulatory review and approval and to periodic revision.

8.24. Long term protection and safety in the management of residues and waste relies primarily on passive means to minimize the need for significant and ongoing maintenance. The passive safety features that are used will depend on the amount and types of residue or waste. For example, uranium process tailings should be stabilized and covered by soil or water to limit radon emissions, and liners are often used and necessary to reduce the chance of groundwater contamination.

8.25. Prior to decommissioning or closure, regulatory criteria should be established for the clearance, reuse and recycling of materials (see paras 6.28 and 6.29). Criteria should also be established, as appropriate, for equipment, structures and the site in terms of, for example, the following:

(a) Removal of equipment and structures from regulatory control;
(b) Reuse and recycling of equipment, structures and material;
(c) Release of the site for unrestricted or restricted use.
8.26. Progressive closure and decommissioning in stages should be undertaken to the extent reasonably practicable during operation.

8.27. The plan for decommissioning and/or closure should be subject to review on the following basis:

(a) Periodically, to take into account ongoing operations, the results of monitoring and any measures implemented for contamination control;
(b) Following modifications made to the facility or the types or quantities of NORM residue being managed;
(c) If there are changes in regulatory requirements or anticipated future uses of the land.

8.28. Recommendations on financial provisions are provided in paras 5.41–5.45. The operating organization should periodically review the financial resources and the plan for decommissioning and/or closure during operation of the facility to ensure that adequate funds are available to cover the full costs of meeting the end state criteria.

8.29. As stated in Requirement 11 of GSR Part 6 [7] (footnote omitted), “Prior to the conduct of decommissioning actions, a final decommissioning plan shall be prepared and shall be submitted to the regulatory body for approval.”

8.30. As stated in Requirement 19 of SSR-5 [8]:

“A disposal facility shall be closed in a way that provides for those safety functions that have been shown by the safety case to be important after closure. Plans for closure, including the transition from active management of the facility, shall be well defined and practicable, so that closure can be carried out safely at an appropriate time.”

8.31. The final decommissioning and/or closure plan is required to be approved by the regulatory body (see Requirement 11 of GSR Part 6 [7] and Requirement 19 of SSR-5 [8]) prior to the initiation of decommissioning and/or closure activities. The final decommissioning and/or closure plan should address at least the following elements:

(a) An assessment of the post-decommissioning and/or post-closure risks to people and the environment.
(b) Land ownership and future land use.
(c) End state criteria — radiological, environmental and landform — and how they are to be met.

(d) Alternatives considered as a means of achieving the required end state.

(e) Concurrent rehabilitation measures.

(f) Provisions for premature closure.

(g) Decommissioning and decontamination procedures and techniques, including the following:
   (i) The reuse and recycling of residues and plant structures, equipment and items containing or contaminated by NORM;
   (ii) The management of NORM residues arising from the decontamination and decommissioning of the facility.

(h) A planned timescale for decommissioning and/or closure.

(i) The need for any remediation of land areas.

(j) The final radiation survey of the site.

(k) The need for any long term institutional control, including monitoring and surveillance.

(l) Involvement of interested parties.

(m) A summary of costs.

(n) A summary of assumptions and uncertainties.

(o) Updates and revisions of the preliminary closure plan.

8.32. NORM residues that arise from operation and from decommissioning can potentially use the same long term facilities for the management of NORM residues. The decommissioning plan should consider the effects of mixing materials from various waste streams and the implications for consolidation and differential settlement.

8.33. Both decommissioning and closure will involve consideration of the non-radiological constituents of NORM residues and waste, and in many cases these non-radiological considerations will be the dominant factors.

8.34. A decommissioning and/or closure report needs to be prepared by the operating organization to confirm that the end state of the facility or site has been achieved, as specified in the approved final decommissioning and/or closure plan. The report should be subject to review and approval by the regulatory body.

8.35. After review of the final decommissioning and/or closure report, and any other verification measures deemed necessary, the regulatory body will decide on the termination of the authorization for a facility following decommissioning and/or closure and on the release of the facility with or without restrictions (see Requirement 15 of GSR Part 6 [7] and paras 5.10 and 5.14 of SSR-5 [8]).
8.36. A system is required to be established to ensure that all safety related records relevant to the decommissioning and/or closure of a facility are maintained (see para. 9.7 of GSR Part 6 [7] and paras 3.15 and 5.13 of SSR-5 [8]). This system should involve the operating organization, the regulatory body, the government and any other entity responsible for implementing long term management and institutional control. The system should be designed to ensure that any persons wishing to access the site are informed about the previous presence of a facility on the site and about the nature of the activities that were conducted on the site.

LONG TERM MANAGEMENT AND INSTITUTIONAL CONTROLS

8.37. If a site cannot be released for unrestricted use, the use of the site should be restricted, and appropriate institutional controls will be necessary to ensure protection of people and the environment over the long term. As stated in para. 1.22(iii) of SSR-5 [8], “institutional controls are put in place to prevent intrusion into facilities and to confirm that the disposal system is performing as expected by means of monitoring and surveillance.” Control may be active (e.g. by means of monitoring, surveillance, remedial work, water diversion and treatment, and fences) or passive (e.g. by means of land use controls, markers and records).

8.38. The long term management period begins when operational buildings and supporting services have been decommissioned, all engineered confinement and isolation features have been put in place, and any remaining facilities are in the final configuration. In accordance with Requirement 22 of SSR-5 [8], after decommissioning actions and closure are complete, the safety of the long term management facility is required to be provided for primarily by means of passive features, including the characteristics of the site and the final covering that has been put in place, together with institutional control measures such as markers (see paras 3.48 and 5.9 of SSR-5 [8]).

8.39. Where institutional controls are considered necessary, a custodian organization for these controls will be necessary; this custodian can be the government (usually an agency other than the regulatory body) or a qualified private entity. The custodian should provide periodic reports to the regulatory body or the government on the situation at the site.

8.40. If active controls are warranted, the operating organization should provide sufficient funds to implement and maintain monitoring, surveillance and control of the facility throughout the necessary time period. The site and any residues therein should not become a financial burden on the government or the public.
8.41. The operating organization is responsible for preparing a proposed programme for long term management of the site for review and approval by the regulatory body (see Requirement 22 of SSR-5 [8]). The design of the programme should be based on safety assessments as described in Section 7, in which impacts on people and the environment over an appropriate period into the future have been considered.

8.42. The safety case prepared by the operating organization should state the period over which institutional controls are planned to remain in force, and this should be subject to approval by the regulatory body. Scenarios postulating human intrusion, failure of engineered structures and changes in environmental conditions should be considered in the safety assessment (see Section 7).

8.43. As part of a long term management programme, all relevant records of the characteristics of closed residue management facilities and of restrictions on land use and ongoing monitoring and/or surveillance measures should be maintained in accordance with applicable legal requirements. Such records should be made available to interested parties upon request (see also paras 3.16 and 5.13 of SSR-5 [8]).

8.44. For some sites currently in operation, or some sites resulting from past practices, the goal of using primarily a passive approach might not be fully achievable (see para. 6.3 of SSR-5 [8]). In such cases, efforts have to be made to minimize the number of active controls.

MONITORING AND SURVEILLANCE

8.45. Requirements 10 and 21 of SSR-5 [8] address monitoring and surveillance programmes for disposal facilities; more detailed recommendations are provided in SSG-31 [16]. Further information on monitoring and surveillance programmes at uranium production facilities is given in Ref. [34].

8.46. The operating organization is required to develop and implement a monitoring and surveillance programme (see Requirements 10 and 21 of SSR-5 [8]); this programme should be subject to regulatory approval. The programme should be conducted and reviewed periodically by the operating organization prior to, during and after operation, decommissioning and closure. The regulatory body should inspect and verify monitoring results throughout the lifetime of the facility and the period of long term management. The institutional controls (see para. 8.37) should
ensure that the monitoring and surveillance programme is robust and continues, as necessary, following closure.

8.47. The monitoring and surveillance programme consists of continuous or periodic observations and measurements to evaluate and verify the behaviour of the residue management facility. The programme includes the measurement of radiological, environmental and engineering parameters. The results of this programme should be used to evaluate the impact of the facility on people and the environment and to support decision making at various stages in the lifetime of the facility.

8.48. The types, duration and frequency of monitoring should be adapted to each period in the lifetime of a facility: the pre-operational period, the operational period (including decommissioning operations) and the post-closure period (see para. 1.22 of SSR-5 [8]).

8.49. A graded approach should be taken to adapt the level of detail (e.g. duration, frequency, locations for sampling, parameters to be monitored) in the monitoring programme so that it is commensurate with the level of risk associated with the facility.

**Pre-operational period**

8.50. The pre-operational period includes site evaluation (selection, verification and confirmation) and safety assessment and design studies. The objectives of the monitoring and surveillance programme during the pre-operational period are the following:

(a) To contribute to the characterization of the site and the evaluation of the suitability of the site.
(b) To provide input for the design and construction of the facility.
(c) To provide input necessary for the operational and post-closure safety cases.
(d) To establish baseline conditions, including a determination of the existing level of natural radioactivity at the site, for comparison with later monitoring results. This is especially important in respect of NORM residues, because the same radionuclides are already present in nature.
(e) To aid in designing the monitoring programme for the operational period.
Operational period

8.51. The objectives of the monitoring and surveillance programme during the operational period are the following:

(a) To demonstrate the protection of workers;
(b) To provide data to confirm the performance of the long term management facility;
(c) To check the performance of effluent treatment and control systems and of abatement systems for airborne releases, as appropriate;
(d) To provide early warning of any deviations from normal operation;
(e) To provide data on the discharge of radionuclides (e.g. rates, concentrations, composition) to the environment for use in predictive modelling and determination of exposures to the public;
(f) To evaluate compliance with regulatory requirements;
(g) To provide information to and support communication with interested parties.

Post-decommissioning and post-closure period

8.52. The monitoring and surveillance programme for the period after decommissioning and closure should be conducted to demonstrate that the facility is performing as predicted and should be used for the following:

(a) To detect abnormal concentrations of radionuclides in the environment that could be attributable to the long term management facility;
(b) To verify the performance and integrity of barriers;
(c) To validate the achievement of post-closure radiological objectives;
(d) To inform decisions on controls, such as moving from active institutional control to passive institutional control to unrestricted release;
(e) To determine the need for, and type of, monitoring and surveillance activities to be conducted during any institutional control period;
(f) To satisfy the principle of openness and transparency of information for interested parties;
(g) To evaluate compliance with regulatory requirements.

8.53. The monitoring and surveillance programme should specify the parameters to be monitored, the locations and frequencies for measurements and sampling, and the procedures for analysis and reporting, including the setting of appropriate action levels. Such a programme should include measurements of the following:
(a) Indicators of environmental impacts, such as levels of radionuclides and non-radiological contaminants in air, water and soil;
(b) The physical integrity of structures and systems for the containment of NORM residues;
(c) Parameters that assist in the interpretation of data, such as meteorological data, operational process data and waste stream data.

8.54. Annex I to Ref. [34] provides an example of the typical content of a long term surveillance plan for a uranium mill tailings site in the post-closure period. This example plan can also be adapted to the surveillance of facilities for other NORM residues with similar characteristics, with account taken of the graded approach.
Appendix I

SPECIAL CONSIDERATIONS FOR RESIDUES FROM URANIUM PRODUCTION

I.1. Uranium production generates various residue streams, including mill tailings, waste rock, mineralized waste rock and process water, including leaching solutions. Rainfall, snowmelt runoff and seepage from stockpiles and areas of uranium process plants should also be managed. In addition, the residue management programme should take into account used pipes, process vessels, filters and mine waters.

URANIUM MINING WASTE ROCK

I.2. Bulk waste rock from uranium mining warrants long term management because of the large volumes generated, the presence of long lived radionuclides and heavy metals, and the potential to generate acidic drainage.

I.3. Bulk waste rock from uranium mining contains all the radionuclides in the original ore in secular equilibrium but at lower activity concentrations. The concentrations of the radionuclides in the uranium decay chain are mostly below 1 Bq/g; however, this can still result in public exposure above 1 mSv in a year. Some options for reusing waste rock materials exist, for example for road construction or as backfilling material.

I.4. In most cases, waste rock is heaped up close to the mine on ground where there is little possibility of negatively impacting water bodies. Seepage water is collected by a drainage system to prevent or reduce the migration of radionuclides into groundwater.

I.5. After closure of the mine (or progressively during operation), waste rock heaps are covered to reduce the infiltration of rainwater. In general, two types of cover are utilized, depending on the potential for acidification of the waste rock material. In particular, different designs should be considered when dealing with alkaline waste rock and pyritic waste rock. Thicker and multilayer cover might be necessary to avoid acidification due to pyrite oxidation and consequently the leaching of radionuclides.

I.6. Options for managing waste rock and mineralized waste rock include use as backfill materials in open pits and underground mines and use in construction
at the mine site. Covering mineralized waste rock with inert waste rock should be considered. As with bulk amounts of mineral processing residues, the stability of piles of waste rock, and their resistance to erosion and rainwater infiltration, should be considered to ensure that these piles do not result in unacceptable environmental impacts on the water catchment area (e.g. acid mine drainage).

I.7. Co-placement of waste rock with tailings is a concept that can be considered for both underground and above ground management options in mining situations. Appropriate cover should be put in place to both inhibit the release of radon to the air and prevent potential human intrusion. However, the chemical and mechanical compatibility of the combined material should be considered.

URANIUM MILL TAILINGS

I.8. Uranium mill tailings represent a challenge in terms of long term management because of the large volumes generated; the presence of long lived radionuclides, heavy metals and chemical hazards; and the potential to generate acidic drainage.

I.9. Tailings contain all the radionuclides in the original ore, at concentrations near their concentration in ore, with the exception of the uranium isotopes and their immediate short lived decay products. Approximately 75% of the original radioactivity present in the uranium ore is retained in the tailings. Tailings are usually discharged as slurry containing about 20–50% solids into a purpose built water-retaining structure or impoundment, either above or below ground level.

I.10. There are few options for reusing tailings. Tailings, particularly the coarser size fractions, might be of use as a component of mine fill; however, engineering considerations can make this problematic, as tailings slimes do not consolidate well on their own. Tailings can also be processed to recover uranium; however, the radiological implications of any such reuse would need to be considered.

I.11. The key issues that should be considered in the design of a tailings management facility include the following:

(a) The stability of the pit, underground mine void or surface impoundment in relation to natural processes such as earthquakes, floods and erosion;
(b) The hydrological, hydrogeological and geochemical characteristics of the site;
(c) The chemical and physical characteristics of the tailings in relation to the potential for generation and transport of contaminants;
The volume of material that will be retained on the site as waste;

The use of neutralization agents, radium precipitating additives, artificial or natural liners, radon barriers and evaporation circuits, depending on the reliability, longevity and durability of such measures.

A thorough investigation of these issues should be undertaken at an early stage when considering options for the management of tailings.

I.12. Relocating tailings to a more favourable site for closure would not normally be the optimum residue management strategy because of the large volumes of waste involved. However, if relocation of the waste is being considered, the optimization of all significant radiological and non-radiological impacts that might be introduced by the relocation itself, including issues relating to the transport of large volumes of waste, should be taken into account.

I.13. The design of a facility for the management of tailings should incorporate drainage systems to consolidate tailings before closure and to reduce excess pore water pressure. In the case of a surface impoundment or a pit, this could be achieved by the installation of a drainage system prior to or during the emplacement of tailings, or by the use of wicks driven into the tailings after emplacement. The base and cap of the impoundment should be built to minimize the release of contaminants, if possible using material of natural origin. The addition of a stabilizing agent (e.g. cement) to the tailings immediately prior to their deposition has the potential to significantly reduce the permeability of the tailings mass, thus retarding the transport of contaminants and binding any pore water. However, in certain cases, a confined water covering in a pit can possess excellent characteristics as a radon barrier, thereby obviating the need to perform dewatering to any significant degree.

I.14. To avoid undue burden on future generations, a passive approach to design for closure is preferable to a design that needs significant and ongoing maintenance. Such a passive approach is generally best achieved by disposal in pits excavated specifically for this purpose, in mined-out pits, in underground mine voids or in natural water bodies, where appropriate. This option might eliminate or significantly reduce the need for surface disposal of tailings.

I.15. The decision on which approach to take should be optimized so as to match barrier characteristics with available site conditions. Mine or process residues disposed of below ground level are less susceptible to surface erosion and to intrusion. Subsurface placement generally necessitates less maintenance than surface tailings impoundments and eliminates the risk of a dam or dyke structural
failure. Closure entails sealing the openings to the underground disposal facility, thereby isolating it from the surface. While buried tailings are less vulnerable to erosion, they might be more vulnerable to groundwater fluctuation if the water level is not deep, or they might be closer to the water table in the event of liner failure.

I.16. In the case of long term management of tailings in underground mines, the increase in structural integrity gained by using concrete with the tailings mass might allow mining to be continued nearby. Prior to adopting this strategy, possible chemical interactions between the stabilizing agent, the tailings and the host rock should be carefully investigated to ensure that the transport of contaminants would not be enhanced at some time in the future or impact the active mine workings or the workers.

I.17. For the disposal of tailings underground, provided that the probabilities of geological disturbance to the site and of human intrusion into the site are deemed to be sufficiently low, it might be that no further controls are necessary other than archiving details of the location and characteristics of the waste and monitoring the site for a limited period.

I.18. Practical engineering solutions can be identified for some site specific problems associated with below ground tailings disposal facilities. For example, if the hydraulic conductivity of the tailings mass is greater than that of the surrounding host rock, the use of a highly permeable envelope surrounding the tailings should be considered as a means of diverting water around the tailings. In the case of a small and confined aquifer intersecting a pit or underground mine wall, localized grouting should be considered.

I.19. It is possible that the below ground disposal of mine tailings at a particular site might not be feasible, either owing to site specific problems for which no engineering solutions can be identified (e.g. when placement is likely to result in contamination of groundwater) or owing to prohibitive costs. In such cases, the use of engineered surface impoundments might be the only viable option and should be considered.

I.20. For options involving the management of tailings in above ground impoundments, the tailings should be contained within low permeability engineered structures so as to reduce seepage. An above ground closure option would usually necessitate having greater institutional control than is needed for an underground option. Monitoring and maintenance programmes should
be implemented during the operational, closure and post-closure periods. This approach would entail lower initial costs but higher continuing costs.

I.21. A cover system that is designed to limit infiltration and radon emissions is necessary for bulk amounts of residues placed above ground. Cover materials that have been effective in reducing radon emissions include water, earthen materials, geosynthetics such as geomembranes and geosynthetic clay liners, and evapotranspirative barriers. Simple covers might contain one type of material; however, robust combinations of different materials are often necessary.

I.22. Cover systems designed to limit infiltration and radon emissions might comprise a lateral drainage layer, consisting of either coarse sand or gravel above a low permeability clay layer, and a top layer of durable rock for erosion protection. Depending on the climate and environment, a vegetative cover for erosion control, stabilization and limiting infiltration might be employed.

I.23. For placement of waste in a pit, the necessary degree of passive control can be achieved either by backfilling and capping with natural materials or by establishing a permanent water pond over the tailings. The latter option might also involve the application of a low permeability cover for the waste to reduce contact with the pond water. The subsurface conditions should be fully investigated to gain sufficient understanding to be able to ensure that the hydraulic pressure over the backfilled pit will not result in problems with groundwater contamination in the future.

I.24. The diffusion coefficient for radon in a saturated soil can be several orders of magnitude lower than that for radon in a dry soil. A water covering or saturated cover layer might therefore serve as an effective radon barrier, although in dry environments a different approach is necessary.

I.25. Depending on the risk of contamination, a groundwater monitoring programme should be considered in order to avoid creating areas that will require remediation in the future.

I.26. In addition to the emplacement of tailings in above ground impoundments, open pits and underground mine voids, there are other options for tailings management, such as the deposition of tailings in lakes. Monitoring and/or geochemical modelling should be undertaken, where appropriate, to show that a reducing environment has been established. However, some of these alternative options might not be acceptable to the regulatory body or the public and would need further study and evaluation.
Other disposal strategies for mill tailings may be appropriate, and they should be evaluated on a case by case basis. For example, small quantities of mill tailings might be accepted for disposal in a facility designed for low level radioactive waste, provided that the waste acceptance criteria of the facility are complied with.

**RESIDUES FROM HEAP LEACHING**

Heap leaching is a method used for processing low-grade uranium ore and typically involves the treatment of crushed or pelletized ore grade material with acid or alkali (or bacteria) on large engineered pads on the surface. Stope leaching or block leaching of uranium ore underground is also conducted. Most heap leaching operations generate medium quantities of residues; however, some operations are quite large and generate residues in bulk amounts.

Surface heap leaching facilities need efficient containment and liquid collection systems, base liners and leak detection systems to collect the leachate for further processing and to protect the surface environment and groundwater resources.

Residues from heap leaching consist of process liquids generated during operation, the leached ore and, potentially, a continuing release of solutions due to infiltration of the closed facility. During operation, waste process solutions can be collected, treated and sent to adjacent evaporation ponds and/or injected into deep injection wells. In some cases, a separate residue storage dam might be necessary, with characteristics similar to those of a tailings dam.

An important consideration is locating the heap leach pad to facilitate decommissioning and isolation of the resulting residues without relocation. Heap flushing and neutralization might be conducted at the same time as decommissioning. Following decommissioning, long term management of the NORM residues might still be necessary.

**RESIDUES FROM IN SITU LEACHING OF URANIUM**

In situ leaching is carried out by drilling a pattern of injection and extraction wells into the ore body and then circulating a leach liquor that is either acid or alkali depending on the host sediments and ores. The uranium is extracted from the resulting ‘pregnant solution’ by conventional solvent extraction or ion exchange methods, and the now ‘barren solution’ is reconstituted and re-injected.
into the leaching field. No conventional tailings are generated, but large volumes of liquid and small to medium amounts of solid residues can be generated.

I.33. A small fraction (0.5–2%) of the leach liquor is bled off, and this bleed stream constitutes the largest volume of liquid residues from the process. Large volumes of liquid residues can also be generated where reconstitution of the ore body aquifer is undertaken following completion of the operation, for example from flushing of the aquifer. Smaller volumes of liquid water are generated from normal facility operation, including from the washdown of equipment and from spillages.

I.34. If the bleed stream is evaporated, elevated concentrations of radionuclides can remain, and if the bleed is treated chemically to remove radionuclides, these will usually be recovered in solid or slurry form.

I.35. In some cases, selenium and radium are removed prior to land application or re-injection of the resulting water. In these cases, small amounts of residues will need to be managed and ultimately disposed of.

I.36. The ore body aquifer might need pretreatment prior to mining, commonly to remove calcium, and the resulting precipitates can contain elevated radium concentrations.

I.37. Liquid residues can be reduced or eliminated by evaporation, or discharged into aquifers or surface water bodies in accordance with the relevant discharge conditions approved by the regulatory body. Injection into deep (and preferably well confined) aquifers is a possible solution, as is injection into shallower aquifers, typically the mining aquifer itself.

I.38. In cases of injection of liquid residues into aquifers, an environmental impact assessment involving detailed hydrogeological modelling of the situation should be undertaken. Techniques for the restoration of groundwater can include natural attenuation, groundwater flushing to accelerate natural attenuation, injection of reducing agents, or groundwater sweep and reverse osmosis. The more intensive restoration techniques should progressively be used, as necessary (i.e. if the effect of restoration is not proving adequate), to achieve the agreed end state criteria for closure of the facility on a reasonable timescale. More intensive methods require more energy and surface infrastructure, generate waste streams, and incur additional costs. Best practice is therefore to use the restoration technique that will achieve the end state criteria for closure on an agreed timescale.
with the minimum environmental impact. Long term monitoring or institutional control may be required.

I.39. Solid radioactive residues generated by an in situ leaching facility can include used pipes, pumps, filters and other equipment contaminated with soil and sludge from ponds and from evaporation of waste liquids. These might be managed in a purpose built management facility that is usually on the site, or else such residues are taken to an off-site residue management facility.
Appendix II

RESIDUE MANAGEMENT PLAN FOR URANIUM PRODUCTION

II.1. The content of a residue and waste management plan for a uranium production facility could include the following:

(a) A description of the processes in which the residues and waste are generated by the facility;
(b) A description of each of the residue streams and waste streams and the measures taken to prevent these streams from arising or to minimize these streams;
(c) The limits and conditions necessary for the waste to be managed safely;
(d) A comprehensive list of the current and anticipated residues and waste arising from, and inventories at, the facility;
(e) A definition of the waste management principles and objectives at the facility;
(f) Identification of residue and waste management options and associated steps, as well as interdependencies between these steps;
(g) A justification of the selection of appropriate management options based on the information above and on international good practices;
(h) A demonstration that the residue and waste management plan is compatible with the national policy and strategy;
(i) A demonstration, if necessary, of how the safety case is affected by the residue and waste management plan (e.g. how a modification of the plan to incorporate longer storage than the building was originally designed for would impact the safety case).

II.2. The plan should include provisions for the following:

(a) Keeping the generation of residues and waste to the minimum practicable, in terms of type, activity and volume, by using suitable technologies;
(b) Possible reuse and recycling of materials;
(c) Appropriate classification and segregation of waste, and maintenance of an accurate inventory for each residue stream and waste stream, with account taken of the available options for clearance or disposal;
(d) Collection, characterization and safe storage of residues and waste;
(e) Adequate storage capacity for the residues and waste that are expected to be generated (conditioned and unconditioned) and an additional reserve storage capacity;
(f) Ensuring that stored residues and waste can be retrieved at any time within the anticipated storage period;

(g) Techniques and suitable procedures available for the retrieval of stored residues and waste;

(h) Processing of radioactive waste to comply with waste acceptance requirements and to ensure safe storage and long term management including disposal of residues for which no further use is foreseen;

(i) Safe handling and transport of residues and waste, if necessary;

(j) Adequate control of discharges of effluents to the environment.
Appendix III

CLOSURE PLAN FOR A TAILINGS MANAGEMENT FACILITY AT A URANIUM PRODUCTION SITE

III.1. A closure plan for a tailings management facility at a uranium production site could include the following:

(a) Introduction, including site location and history:
   (i) Amounts and types of material produced;
   (ii) Activities undertaken;
   (iii) Previous site assessments;
   (iv) Applicable regulatory end state criteria to be met;
   (v) Current environmental and radiological conditions.

(b) Geology and seismology:
   (i) Stratigraphic features;
   (ii) Structural and tectonic features;
   (iii) Geomorphic features;
   (iv) Seismicity and ground motion estimates.

(c) Geotechnical stability:
   (i) Site and uranium mill tailings characteristics;
   (ii) Slope stability;
   (iii) Settlement;
   (iv) Liquefaction potential;
   (v) Engineering design of the disposal cell cover;
   (vi) Construction considerations;
   (vii) Hydraulic conductivity of the disposal cell.

(d) Surface water hydrology and erosion protection:
   (i) Hydrological description of site;
   (ii) Flooding determinations;
   (iii) Water surface profiles, channel velocities, and shear stresses;
   (iv) Design of erosion protection;
   (v) Design of erosion protection covers;
   (vi) Protecting water resources.

(e) Groundwater protection:
   (i) Standards for groundwater quality;
   (ii) Monitoring results (baseline, during operation and post-operational);
   (iii) Environmental impact assessment;
   (iv) Corrective action assessment;
   (v) Groundwater corrective action and compliance monitoring plans.
(f) Air quality:
(i) Standards for air quality;
(ii) Monitoring results (baseline, during operation and post-operational).

(g) Radiation protection:
(i) Engineered cover of the tailings management facility (type of material, thickness, ability to prevent radon emissions in the long term);
(ii) Attenuation of radon releases;
(iii) Attenuation of gamma radiation;
(iv) Radioactivity content of the cover.

(h) Closure plan for the site:
(i) Types of restriction to the site for long term access control;
(ii) Site access and the need for institutional controls;
(iii) Discussion of the long term stability and containment of residues and waste;
(iv) The proposed types of engagement with interested parties;
(v) A description of the final form of land features, including demographics and possible receptors;
(vi) Schedule and budget.
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Annex I

EXAMPLES OF NORM RESIDUES TO BE ASSESSED FOR POSSIBLE REGULATORY CONTROL

I–1. Table I–1 provides an indicative list of residues that might need regulatory consideration. It is based on table 1 of Ref. [I–1].

<table>
<thead>
<tr>
<th>TABLE I–1. RESIDUES TO BE ASSESSED FOR POSSIBLE REGULATORY CONTROL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Category</td>
</tr>
<tr>
<td>----------------------------</td>
</tr>
<tr>
<td>By-products</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>Slags</td>
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<tr>
<td></td>
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<tr>
<td></td>
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<tr>
<td></td>
</tr>
<tr>
<td>Scales, sludge sediments</td>
</tr>
<tr>
<td>and other residues</td>
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<tr>
<td></td>
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<tr>
<td>Precipitator dust</td>
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<td></td>
</tr>
</tbody>
</table>

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REFERENCES TO ANNEX I

Annex II

SAMPLING NORM RESIDUES AND DETERMINING RADIONUCLIDE ACTIVITY CONCENTRATIONS

INTRODUCTION

II–1. For activities involving naturally occurring radioactive material (NORM), it is useful to conduct an initial screening assessment designed to eliminate from further regulatory consideration a facility or activity that poses a low level hazard. This will normally involve the sampling and analysis of NORM materials, residues and waste to determine the radionuclide activity concentrations. Further information can be found in Refs [II–1, II–2].

II–2. The most probable radionuclides for which the activity concentrations need to be determined are as follows:

— For the uranium decay chain: $^{238}$U, $^{230}$Th, $^{226}$Ra, $^{210}$Pb and $^{210}$Po.
— For the thorium decay chain: $^{232}$Th, $^{228}$Ra, $^{224}$Ra and $^{228}$Th.

SAMPLING OF MATERIAL

II–3. Collecting representative samples is a prerequisite for obtaining reliable results. Sampling positions and numbers of samples are also important. The quantities of material containing NORM can be very large and exhibit a significant range of activity concentrations owing to the inhomogeneous distribution of radionuclides. The activity concentration might also vary over time. To the extent practicable, both these variations need to be taken into account when developing a suitable material sampling strategy.

II–4. The number of samples collected for analysis is important for obtaining a reasonable estimate of the average activity concentration: the greater the number of samples collected and analysed, the greater the confidence in the results obtained. A point is reached, however, where any further gain in accuracy is minimal compared with the additional time and resources needed to analyse more samples. The accuracy of results is also affected by other factors, such as the degree to which the samples are representative of the material as a whole.
MEASUREMENT ACCURACY AND QUALITY ASSURANCE

II–5. Adequate confidence in the results of analyses is ensured if the samples are analysed at a suitably accredited laboratory and if the level of accuracy of the analytical technique is commensurate with the activity concentration criterion against which the material is being compared. If an accredited laboratory is not available, the analytical techniques can at least be validated against appropriate reference materials. Problems due to cross-contamination between samples and contamination of equipment can be avoided by exercising an appropriate level of care during sampling and at the laboratory.

II–6. The distribution of activity concentrations in a material might span an order of magnitude or more. The lower limit of detection of the analysis needs to be well below the activity concentration level against which the measurements are being compared. For instance, when a material is being compared against a value of 1 Bq/g for radionuclides in the uranium decay chain and thorium decay chain (or 10 Bq/g for $^{40}$K), a lower limit of detection of 0.1 Bq/g (1 Bq/g for $^{40}$K) would be appropriate.

ANALYTICAL TECHNIQUES

II–7. Having defined the main radionuclides of interest (e.g. on the basis of knowledge of the process or from a search of the relevant literature) and the necessary measurement sensitivity, appropriate analytical protocols can be considered. Analysis techniques for determining activity concentrations of individual radionuclides in solid materials can be time consuming and expensive. The techniques employed for a particular sample therefore need to be chosen carefully.

II–8. For a general screening of the total radioactivity, it might be adequate to perform gross alpha–beta counting, applying suitable corrections for self-absorption, where appropriate. This technique is a relatively quick and inexpensive method for determining the total activities (or activity concentrations) of alpha emitting and beta emitting radionuclides, from which the ratio of the two can be obtained. This technique does not give information on individual radionuclides; however, the alpha–beta ratio can provide some indication of the radionuclide composition, which can be useful in deciding on subsequent analysis steps. If the total activity concentration is less than the activity concentration criterion for individual radionuclides, no further analysis is necessary. Counting times are selected to obtain the necessary lower limit of detection for the materials concerned.
II–9. For analysis of individual radionuclides of interest, the following analytical techniques [II–1] can be applied:

(a) X-ray fluorescence spectrometry: This method is widely used to measure the elemental composition of materials and is suitable for the rapid determination of uranium and thorium. There are two types of spectrometer, both of which can be used for this application:
   (i) Wavelength dispersive spectrometers, in which photons are separated by diffraction on an analysing crystal before being detected.
   (ii) Energy dispersive spectrometers, in which the energy of the photon is determined when it is detected. These spectrometers are smaller and less expensive than wavelength dispersive spectrometers, and the measurement is faster; however, the resolution and the limit of detection are not as good.

(b) Inductively coupled plasma atomic emission spectroscopy: This method is used for the chemical analysis of aqueous solutions of rocks and other materials and is suitable for the determination of a wide range of major elements and a limited number of trace elements. Sample preparation involves the digestion of the powdered material with 40% (vol./vol.) hydrofluoric acid mixed with either perchloric or nitric acid. Some minerals, such as chromite, zircon, rutile and tourmaline, will not completely dissolve using this digestion procedure. For samples containing substantial amounts of these minerals, X-ray fluorescence analysis is probably more appropriate.

(c) Inductively coupled plasma mass spectrometry: This method is used to determine trace elements in aqueous solutions. The technique is suitable for the determination of uranium and thorium. The sample preparation procedure is the same as that for inductively coupled plasma atomic emission spectroscopy.

(d) High energy gamma spectrometry (high purity germanium crystal detector): This technique provides a quantification of radionuclides such as $^{226}$Ra, $^{228}$Ra, $^{228}$Th and (if needed) $^{40}$K. The method can also be used to quantify the $^{238}$U concentration, although the lower limit of detection is relatively poor.

(e) Low energy gamma spectrometry (high purity germanium crystal or lithium drifted silicon crystal detector): This technique provides a quantification of $^{238}$U and $^{210}$Pb (as well as $^{235}$U). The technique can also provide a determination of $^{226}$Ra (as well as other radionuclides such as $^{227}$Ac, $^{231}$Pa and $^{230}$Th), but with a poorer lower limit of detection.

(f) Alpha spectrometry: Sample digestion followed by various chemical separation techniques and then alpha spectrometry can be used to determine each of the NORM alpha emitters. This technique is commonly used for the
quantification of $^{210}$Po. However, different digestion methods can produce large variations in the results.

(g) Scintillation detector system: Where the radionuclide composition is well known, the activity might be determined by means of a handheld gamma spectrometry instrument.

II–10. The application of the techniques described in para. II–9 is summarized in Table II–1. The minimum sample size needed is in each case about 10 g, although for techniques involving high energy gamma spectrometry, larger samples (up to 1 kg) are preferred. When undertaking analyses for elemental uranium or thorium, the following conversions from ppm to Bq/g can be used:

- $1 \text{ ppm uranium} = 0.012436 \text{ Bq/g } ^{238}\text{U}$;
- $1 \text{ ppm thorium} = 0.004057 \text{ Bq/g } ^{232}\text{Th}$.

II–11. For material associated with most NORM industrial processes, it is adequate to have a basic analytical infrastructure consisting of X ray fluorescence spectrometry in combination with a background shielded, thin window, high purity germanium crystal gamma spectrometry system. Only in those processes where $^{210}$Po is of concern will radiochemical techniques in combination with alpha spectrometry be necessary.

TABLE II–1. ANALYTICAL TECHNIQUES FOR DETERMINING RADIONUCLIDE ACTIVITY CONCENTRATIONS

<table>
<thead>
<tr>
<th>Radionuclide</th>
<th>Suitable technique</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>U-238, Th-232</td>
<td>X ray fluorescence spectrometry, inductively coupled plasma atomic emission spectroscopy, inductively coupled plasma mass spectroscopy</td>
<td>Sensitivity of 1 ppm uranium or thorium achievable with any of these techniques (equivalent to about 0.01 Bq/g U-238 and 0.004 Bq/g Th-232)</td>
</tr>
</tbody>
</table>
TABLE II–1. ANALYTICAL TECHNIQUES FOR DETERMINING RADIONUCLIDE ACTIVITY CONCENTRATIONS (cont.)

<table>
<thead>
<tr>
<th>Radionuclide</th>
<th>Suitable technique</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ra-226,</td>
<td>High energy gamma spectrometry</td>
<td>The presence of uranium can interfere with the direct determination of Ra-226</td>
</tr>
<tr>
<td>Ra-228,</td>
<td></td>
<td>For indirect determination of Ra-226, gas-tight sealing of the sample for 3 weeks is needed to ensure equilibrium with progeny (Pb-214, Bi-214)</td>
</tr>
<tr>
<td>Th-228 (and K-40)</td>
<td></td>
<td>To achieve a lower limit of detection of 0.1 Bq/g, the detector needs to be shielded from background radiation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High relative efficiency (&gt;25%) and high resolution high purity germanium detectors are needed</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Counting times of a few hours per sample will be adequate</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Self-absorption corrections are necessary for high density materials (&gt; 2.5 g/cm³)</td>
</tr>
<tr>
<td>Pb-210</td>
<td>Low energy gamma spectrometry</td>
<td>Self-absorption corrections are necessary</td>
</tr>
<tr>
<td></td>
<td></td>
<td>To achieve a lower limit of detection of 0.1 Bq/g, the detector needs to be shielded from background radiation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Counting times of a few hours per sample will be adequate</td>
</tr>
<tr>
<td>Po-210</td>
<td>Sample digestion + alpha</td>
<td>Microwave acid digestion might be necessary</td>
</tr>
<tr>
<td></td>
<td>spectrometry</td>
<td>Validated radiochemical separation techniques are needed</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Counting times of a few hours per sample will be adequate to achieve a lower limit of detection of 0.1 Bq/g</td>
</tr>
</tbody>
</table>

REFERENCES TO ANNEX II

Annex III

EXAMPLE OF APPLICATION OF THE GRADED APPROACH IN THE MANAGEMENT OF NORM RESIDUES

GENERAL

III–1. This annex provides further information on various aspects for which the graded approach can be applied and includes an example of the application of the graded approach to naturally occurring radioactive material (NORM) residue management.

III–2. Applying the graded approach is intended to ensure that the level of effort applied to achieving protection and safety is optimized (i.e. the level of effort is commensurate with the magnitude of the radiation risks and their amenability to control).

III–3. For NORM, the ideal regulatory infrastructure has flexibility in terms of the control measures needed for different conditions, based on specific criteria. These different levels could be accommodated through suitable regulations or a system for authorization that explicitly accommodates different levels of control that are based on NORM residue characteristics and the associated levels of risk.

III–4. The first step in establishing a regulatory framework for the management of NORM residues is to determine a list of activities that need to be considered for regulatory control. The list provides the opportunity for the national authorities to proactively investigate particular sectors and, on the basis of an initial assessment, exclude those that do not warrant further investigation. A periodic review of the list is advisable.

III–5. Classification of residues into different categories may provide a framework to readily apply a graded approach to the way these residues are processed or disposed of (i.e. exemption, unconditional clearance, specific clearance, disposal in conventional landfill sites, disposal in a facility for NORM residues or a radioactive waste disposal facility). A graded system of control based on categories of NORM residues should be developed from knowledge of the NORM residues, including their radiological and chemical characteristics, and the control measures (regulatory or otherwise) that are already in place.
III–6. As noted in Section 5 of this Safety Guide, the graded approach will also define the authorization process in terms of registration or licensing. This can be taken further by applying a graded approach in the form of different categories of licence or by adapting the authorization conditions to the specific characteristics of the NORM residues.

III–7. The graded approach will also influence the strategy that is applied to the radiological and chemical characterization of residues, including the level of detail needed, for example in terms of the dose rate from the NORM residues, the activity concentrations of radionuclides, the doses received by workers and by the public, and the impact on the environment.

III–8. The graded approach is also applied to the degree of detail contained in the safety assessment. In many cases, the application of a generic approach, or the use of simple and pragmatic rules or models, will be sufficient to estimate occupational and public exposures. If a specific assessment is performed, the complexity of the assessment and the effort involved are expected to be commensurate with the magnitude of the risks.

EXAMPLE OF THE GRADED APPROACH, BELGIUM

III–9. The Belgian Federal Agency for Nuclear Control (FANC) and regional environmental authorities authorize operating organizations that generate, process and dispose of NORM residues.

III–10. The list of activities that require regulatory control is consistent with Ref. [III–1]. The list is reviewed and amended through the royal decree on radiation protection (Ref. [III–2]) to include any additional activities that are newly identified. Up to now, the majority of NORM residues have arisen from the phosphate industry (gypsum and CaF$_2$ sludge) and the titanium dioxide industry (TiO$_2$ filter cake).

III–11. According to the Belgian radiation protection regulations [III–2], NORM residues are subject to regulatory control when they contain radionuclides of natural origin with an activity concentration exceeding the exemption levels described in Ref. [III–3] (i.e. 0.5 Bq/g for $^{238}$U and $^{232}$Th in secular equilibrium with their progeny). If the activity concentration exceeds these levels, any operating organization that processes or disposes of the residues is required to notify FANC. These levels are thus applied to determine exemption from notification; they are not intended to define what is ‘radioactive’ and ‘non-radioactive’ waste.
III–12. If the notification level is exceeded, the residue may be accepted by treatment facilities for non-radioactive waste, subject to a set of generic conditions. On basis of the information provided in the notification by the operating organization, FANC imposes waste acceptance criteria on the treatment facility, derived from a generic safety assessment. These acceptance criteria consist of limits on the maximum activity concentration per batch of NORM residues as well as a limit on the total quantities of NORM waste that can be annually disposed of to landfill. Other limits are applied to the activity concentration of the end product and/or to the residues from the processing operations. These generic acceptance criteria, shown in Table III–1, are imposed on the waste facility in the form of conditions attached to an authorization by registration.

III–13. If these generic acceptance criteria are not met, the waste facility (or the facility that generates the NORM residue) needs to submit a detailed safety assessment demonstrating that the impact to the public of the waste disposal or processing is less than 0.3 mSv in a year.

III–14. The level of detail needed in the safety assessment is defined by FANC on a case by case basis. In some cases (e.g. a phosphogypsum stack), the operating organization is allowed to refer to an assessment performed for facilities with similar characteristics. The conclusions of the environmental impact assessment performed for the environmental permit will also be taken into consideration.

<table>
<thead>
<tr>
<th>Type of processing</th>
<th>Maximum activity concentration (Bq/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Input (single batch of residues)</td>
</tr>
<tr>
<td>Disposal to a landfill for hazardous waste</td>
<td>50</td>
</tr>
<tr>
<td>Disposal to a landfill for non-hazardous waste</td>
<td>10</td>
</tr>
<tr>
<td>Incineration</td>
<td>10</td>
</tr>
</tbody>
</table>
III–15. If it is demonstrated that public exposures are lower than 0.3 mSv in a year, FANC imposes specific conditions in the form of a registration or a licence, such as a monitoring programme for the relevant radionuclides.

III–16. If the detailed safety assessment indicates that public exposure can exceed 0.3 mSv in a year, the NORM residue is treated as radioactive waste and managed by the Belgian National Waste Agency.

REFERENCES TO ANNEX III


Annex IV

REUSE AND RECYCLING OF NORM RESIDUES

IV–1. Reuse can be defined as the reutilization of materials for their original purpose, either in their original form or in a recovered state. Recycling is the utilization of materials, tools and equipment for other than the original purpose, with or without treatment. The reuse and recycling options are attractive in cases in which there is a strong economic incentive to use large amounts of naturally occurring radioactive material (NORM) residues and to avoid the costs associated with long term management. The decision of whether to reuse and/or recycle residues depends on many factors that are specific to the type of residue, the activities concerned and the situation in the State. Implementation of reuse and recycling options requires the establishment of suitable criteria, together with a suitable measurement methodology and suitable instrumentation.

IV–2. Mixing of NORM residues with other materials might be considered as a means to facilitate reuse and recycling. Although the Euratom Basic Safety Standards [IV–1] prohibit the deliberate dilution of radioactive materials for the purpose of releasing them from regulatory control, the mixing of materials that takes place in normal operations (i.e. where radioactivity is not a consideration) is not subject to this prohibition. The competent authority might authorize, in specific circumstances, the mixing of radioactive and non-radioactive materials for the purpose of reuse or recycling.

IV–3. Some examples of reuse and recycling of NORM residues are described in the following paragraphs.

SCRAP METAL

IV–4. Contaminated scrap metal from NORM activities can, in many cases, be decontaminated by various methods. Details of decontamination methods for equipment in the oil and gas industry, as well as information on measurement principles and instrumentation, are given in Ref. [IV–2]. The decontaminated metals can be recycled.

IV–5. The contaminated scrap might also be melted in dedicated furnaces: natural radionuclides normally transfer to the slag, leaving the metal clean for reuse. Depending on the activity concentration, the slag can also be reused if regulatory requirements can be met. Melting of contaminated scrap is
generally a regulated practice and complies with requirements established by the regulatory body.

IV–6. The transport of contaminated items is subject to the requirements established in IAEA Safety Standards Series No. SSR-6 (Rev. 1), Regulations for the Safe Transport of Radioactive Material, 2018 Edition [IV–3], when the activity concentration limits for exempt material and the activity limits for exempt consignments are exceeded.

SLAG

IV–7. Slag from NORM activities can be used as landfill or in road construction. An example of the latter is the use of slag from the thermal phosphorus production industry in road construction in the Netherlands [IV–4].

FLY ASH

IV–8. In many cases, fly ash from coal fired stations is recycled into building materials, for instance as additives to concrete or in lightweight building materials. In some States, the use of fly ash in concrete blocks for building construction is not of concern where the activity concentration is well below 1 Bq/g. In other States, regulations specify the maximum permissible activity concentrations in concrete and in imported building materials, such as certain types of cement.

PHOSPHOGYPSUM

IV–9. There are several options for the recycling of phosphogypsum, such as use as a fertilizer additive and use in road construction and in building materials; detailed information can be found in Ref. [IV–4]. Treatments to improve soils for agricultural use often employ natural gypsum, but phosphogypsum might also be recycled for use in soils. However, as well as the radiological issues associated with this option, non-radiological contaminants, such as cadmium and fluorine, have an impact on the suitability of recycling this residue in agriculture.

IV–10. Phosphogypsum, when subjected to compaction, can be transformed into a solid of valuable strength. It has been used as a binder to stabilize soil and as a replacement for shell and clay in road and parking lot construction. These uses result in significant savings in cost compared with the traditional method of construction. Radiation monitoring during road construction indicated no significant radiological hazards, either to the construction workers or to members of the public living in the area.
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