

Hanford Site Composite Analysis: Radionuclide Selection for Groundwater Pathway Evaluation

Prepared for the U.S. Department of Energy
Assistant Secretary for Environmental Management

Contractor for the U.S. Department of Energy
under Contract DE-AC06-08RL14788

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Terms

| | |
|----------------|--|
| BBI | Best Basis Inventory |
| CA | composite analysis |
| CERCLA | <i>Comprehensive Environmental Response, Compensation, and Liability Act of 1980</i> |
| COPC | contaminant of potential concern |
| DOE | U.S. Department of Energy |
| ERDF | Environmental Restoration Disposal Facility |
| FFTF | Fast Flux Test Facility |
| IDF | Integrated Disposal Facility |
| K _d | distribution coefficient |
| LLBG | low-level burial ground |
| PA | performance assessment |
| TC&WM EIS | Tank Closure and Waste Management Environmental Impact Statement |
| WMA | waste management area |

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1 Introduction

The updated Hanford Site Composite Analysis (CA) will provide an all-pathways dose projection to a hypothetical future member of the public from all planned low-level radioactive waste disposal facilities and potential contributions from all other projected end-state sources of radioactive material left at the Hanford Site following site closure. Its primary purpose is to support the decision-making process of the U.S. Department of Energy (DOE) under DOE O 435.1-1, *Radioactive Waste Management*, related to managing low-level waste disposal facilities at the Hanford Site.

A key aspect of conducting a CA is selecting the radionuclides to be analyzed. This document describes the selection process for radionuclides to be included in the quantitative analysis.

2 Background

The disposed inventory estimate is perhaps the most important component of the CA, as it directly affects the future radiological impacts following site closure. The primary purpose is to estimate radionuclide inventory from site inception to closure. Because of the inclusive nature of a CA, all relevant contaminants are identified and initially considered. Then, subsets of contaminants appropriate for quantitative analysis are selected. Reducing the number of radionuclides for inclusion in the quantitative analysis helps focus budget and resources on simulating only those radionuclides that are likely to contribute to the total dose to the receptor above a threshold value.

3 Screening Methodology

The approaches adopted in the three prior site-wide studies were evaluated to help develop a radionuclide screening process for the CA.

For the Hanford Site CA update, the methodology for selecting radionuclides to be included is based, in part, on the following aspects that influence the scope and approach:

- Use information from past Hanford Site-wide studies to guide the methodology for screening radionuclides. Three past studies related to waste site evaluation and radionuclides inventories provide valuable insights into the radionuclide selection process:
 1. CA of radionuclides conducted in 1998, documented in PNNL-11800, *Composite Analysis for Low-Level Waste Disposal in the 200-Area Plateau of the Hanford Site*, and PNNL-11800 Addendum 1, *Addendum to Composite Analysis for Low-Level Waste Disposal in the 200 Area Plateau of the Hanford Site*.
 2. A site-wide inventory of radionuclides conducted in 2006, documented in PNNL-15829, *Inventory Data Package for Hanford Assessments*, hereinafter referred to as the 2006 Data Package.
 3. A site-wide analysis of cumulative impacts from radionuclides and chemicals documented in DOE/EIS-0391, *Final Tank Closure and Waste Management Environmental Impact Statement for the Hanford Site, Richland, Washington*, hereinafter referred to as the TC&WM EIS.
- Include any new information since the past site-wide studies were conducted, including the Environmental Restoration Disposal Facility (ERDF), Waste Management Area (WMA) C, and the Integrated Disposal Facility (IDF) performance assessments, and the updated Soil Inventory Model (ECF-HANFORD-17-0079, *Hanford Soil Inventory Model (SIM-v2) Calculated Radionuclide Inventory of Direct Liquid Discharges to Soil in the Hanford Site's 200 Areas*).

3.1 Approaches Considered in Past Site-wide Studies to Select Radionuclides

The approaches used in past site-wide studies (the 1998 CA, the 2006 Data Package, and the TC&WM EIS) are considered in the following subsections.

3.1.1 Initial 1998/2001 Composite Analysis

In the 1998 CA, radionuclides were selected primarily based on those identified as potentially significant contributors to dose in the 200 West and 200 East performance assessments and ERDF remedial investigation/feasibility study. In addition, other studies were reviewed to identify radionuclides unique to specific types of wastes or closed facilities, and to identify key radionuclides in immobilized low-activity radioactive waste from single- and double-shell tanks and residing in burial grounds. Different lists of radionuclides were developed for groundwater and air pathways.

The selection process assumed that sources outside of the Central Plateau would be remediated and not represent significant sources of radionuclides following site closure. It also assumed eight of the nine production reactors would be disposed on the Central Plateau; the ninth reactor had been declared a national historic monument and was expected to remain along the Columbia River.

3.1.2 2006 Data Package

In the 2006 Data Package, radionuclides were selected using the data quality objective process. The intent was to identify those radionuclides that had been observed in the environment or had sufficient inventory in waste sites to potentially impact human or ecological health. The screening process reviewed all groundwater monitoring data from 1990 to December 2002 using the following steps:

- Retain all sample results above detection levels.
- Retain all samples not rejected by data quality assurance checks.
- Retain all radionuclides with a half-life greater than 10 years.
- Identify all samples above drinking water standards.
- Identify all radionuclides that have regional or Hanford Site scale distribution (specifically, radionuclides present at more than one or two points in the aquifer).
- Identify all radionuclides with a temporal distribution of more than a single moment in time.
- Add radionuclides that could have a future impact as indicated by performance assessments and environmental impact statement studies.

This process resulted in 16 radionuclides being retained for quantitative analysis.

3.1.3 Tank Closure and Waste Management Environmental Impact Statement

The intent of the TC&WM EIS screening processes was to focus attention on the constituents that control the impacts to groundwater. Separate screening processes were conducted for sites evaluated for cumulative impacts, for the alternatives analysis, and for human health impacts. Contaminants of potential concern (COPCs) assessed for ecological impacts are also summarized below.

3.1.3.1 Cumulative Impacts Analysis

For the sites evaluated under the cumulative impacts analysis, the initial list included radionuclides with half-lives greater than 10 years. Constituents were considered to pose a potential health risk from ingestion if they had a maximum contaminant level or were listed in the Integrated Risk Information

System as having a health-based ingestion standard. As described in Appendix S of the TC&WM EIS, the screening process was intended to select those constituents appropriate for a groundwater release scenario; thus, for radionuclides, “...only groundwater consumption was considered, release was assumed to be partition limited, and decay during transport was considered” (DOE/EIS-0391, p. S-16). Relative impacts were based on the distribution of radionuclides in the cumulative impacts inventory. The initial list was screened, removing radionuclides contributing less than one percent of the impacts under drinking water consumption scenarios and chemicals present at levels below health-based limits. The screening resulted in a final set of 14 radioactive constituents (DOE/EIS-0391, p. S-16).

3.1.3.2 Alternatives Impacts Analysis

For sites evaluated under the alternative analysis, different processes were used to select constituents for tank closure, Fast Flux Test Facility (FFTF) decommissioning, and waste management alternatives.

The Best Basis Inventory (BBI), which included 46 radionuclides and 24 chemicals, was used as the initial list of constituents to consider for evaluating the tank closure and waste management alternatives. Constituents were screened out if they contributed less than one percent of impacts on drinking water ingestion for the chemicals, and on intruder or drinking water consumption scenarios for the radionuclides. As described in Appendix D of the TC&WM EIS:

“Not all constituents are important in the exposure scenarios used to assess TC & WM EIS alternative implementation impacts. Thus, to focus attention on the constituents that control the impacts, DOE performed an initial screening analysis. For radionuclides, groundwater release and direct intrusion scenarios were considered. For the groundwater release screening scenario, only drinking water consumption was considered. Release was assumed partition limited, and decay during transport was considered. For the direct intrusion screening scenario, inadvertent soil ingestion and inhalation pathways were considered.”

“The analysis estimated relative impacts based on distribution of radionuclides in the BBI for all tanks. Radionuclides contributing less than 1 percent of impacts under intruder or well scenarios were eliminated from the detailed analysis. To screen for hazardous chemicals, drinking water ingestion impacts were estimated for the 24 BBI chemical constituents, and those contributing more than 99 percent of impacts were selected for detailed analysis. In addition, reported tank concentrations were reviewed and compared with health-based limits (DOE 2003a)¹; chemical COPCs, when compared with health-based limits (DOE 2003a), were added to the initial list of screened chemicals.” (DOE/EIS-0391, p. D-3).”

The screening resulted in 10 radionuclides and 10 chemicals being selected for detailed analysis, listed in Table D-2 in the TC&WM EIS (p. D-4). One of the radionuclides, Americium-241, is applied to the intruder scenarios only via the inhalation pathway. Although Appendix D mentions that other COPCs were added to the list from the screening conducted for the cumulative impact analysis (last paragraph, p. D-3), the tables comparing tank alternatives only list 9 radionuclides and 10 chemicals (for examples, see Tables D-35 through D-60).

¹ DOE 2003a from the quoted material references: DOE (U.S. Department of Energy), 2003a, *Environmental Impact Statement for Retrieval, Treatment, and Disposal of Tank Waste and Closure of Single-Shell Tanks at the Hanford Site, Richland, WA: Inventory and Source Term Data Package*, DOE/ORP-2003-02, Rev. 0, Office of River Protection, Richland, Washington, April 17.

For the waste management alternatives, three categories of waste were considered: secondary low-level waste and mixed low-level wastes managed at three Hanford facilities; onsite non-*Comprehensive Environmental Response, Compensation, and Liability Act of 1980* (non-CERCLA) waste, non-tank-activity waste; and offsite waste.

Secondary low-level waste and mixed low-level wastes from the operation of three sites were evaluated: low-level burial ground (LLBG) 218-W-5 (trenches 31 and 34), the Waste Receiving and Processing Facility, and the T Plant complex. To evaluate the waste management alternatives, the same 9 radionuclides and 10 chemicals as the tank alternatives were considered, but only three of the chemicals were evaluated because inventories for the other 7 chemicals were not included in the cited report (p. D-129, footnote 'a' to Table D-82).

No screening process was described in Section D.3.5, "Radionuclide and Chemical Inventory Estimates for Onsite Non-CERCLA, Non-Tank-Activity Waste." In the table summarizing the inventory of non-CERCLA, non-tank-activity waste, the same nine radionuclides as in the tank alternatives are reported, but an expanded list of 19 chemicals is reported, based on a Solid Waste Information Tracking System forecast from fiscal year 2006 to fiscal year 2035 as reported in SAIC, 2011, *Waste Inventories Reference Mapping*.

Similarly, no screening process was described in Section D.3.6 "Projected Volumes, Radionuclide and Chemical Inventories for Offsite Waste." Inventories from projected waste volumes that could be shipped to Hanford list the same nine radionuclides as in the tank alternatives but list 15 chemical constituents.

To evaluate the FFTF alternatives, inventories of various radionuclides and chemicals were obtained from existing reports, such as *FFTF Radioactive and Hazardous Materials Inventory* (in DOE/EIS-0391), *FFTF-18346, Technical Information Document for the Fast Flux Test Facility Closure Project Environmental Impact Statement*, and Kidd, 2005, *Activation of the FFTF Biological Shield Wall*. The process used to screen the many reported constituents to the selected four radionuclides and three chemical constituents was described in the TC&WM EIS as follows: "Matching the list of radionuclides and chemicals identified in the above tables with the COPCs identified in Appendix D, Section D.1.1, resulted in a report of the following radionuclides (in curies)..." (p. D-119). The "above tables" refers to tables of inventories reported from the various sources, with differing numbers of COPCs, from seven to 31. Section D.1.1 is "Current Tank Inventory of Radioactive and Chemical Constituents" and includes Table D-2, "Constituents Selected for Detailed Analysis," which lists the 10 radionuclides and 10 chemicals mentioned above. In the FFTF alternatives analysis, four radionuclides and three chemicals were reported in the tables and figures showing the inventories of COPCs.

3.1.3.3 Human Health Impacts Analysis

In Appendix Q of the TC&WM EIS, "Long-term Human Health Dose and Risk Analysis," the screening process is described as follows: Using the inventories in Appendix D for the alternatives analysis and Appendix S for the cumulative impacts analysis, relative impacts were estimated based on the distribution of radionuclides in wastes associated with tanks, FFTF decommissioning, the IDF, the proposed River Protection Project Disposal Facility, and cumulative analysis sites. Radionuclides contributing less than one percent of impacts for intruder (inadvertent soil ingestion and inhalation) or drinking water scenarios and chemicals contributing less than one percent of drinking water impacts were screened out. The result was a list of 14 radionuclides and 26 chemical constituents (Table Q-1, p. Q-2).

3.1.3.4 Ecological Impacts Analysis

A screening process to select COPCs to assess ecological impacts was not described in Appendix P, "Ecological Resources and Risk Analysis." Appendix P describes potential ecological impacts of airborne

releases during operations and groundwater discharges under various alternatives. Appendix P states “Concentrations of radionuclides and chemicals resulting from deposition of airborne contaminants during construction and operations associated with the alternatives were predicted, as described in Appendix G.” (p. P-6). However, Appendix G, “Air Quality Analysis,” states “This appendix presents information on the nonradiological air quality impacts that could result from emissions associated with construction, operations, deactivation, and closure activities under the various alternatives...” (p. G-1). The only constituents described in Appendix G are nonradiological ambient air pollutants such as carbon monoxide; PM₁₀; and sulfur dioxide and other pollutants such as benzene, mercury, formaldehyde, and 1,3-butadiene.

Appendix P also describes that predicted seep, sediment pore water, sediment, and surface water “...concentrations were calculated from the modeled groundwater concentrations at the Columbia River resulting from the varying radioactive and chemical COPC inventories in place under the different alternatives (see Appendix O).” (p. P-46). Appendix O, “Groundwater Transport Analysis,” describes the particle-tracking method used to implement the contaminant transport model. Radionuclides included in the particle-tracking analysis were the same as the screened COPCs to assess human health in Appendix Q, except that plutonium-239 and uranium-238 were listed instead of plutonium and uranium isotopes. For chemicals, Appendix P again points to Appendix G.

3.1.3.5 Comparison of Different TC&WM EIS Screening Results

In the TC&WM EIS, the tables of constituents selected for detailed analysis were the same for the human health impacts analysis and the cumulative impacts analysis (DOE/EIS-0391) as shown in Table 1, despite key differences in the screening methodologies. The human health impacts screening considered groundwater release and intruder scenarios while the cumulative impacts screening only considered groundwater consumption. In the human health impacts screening, relative impacts were estimated based on the distribution of radionuclides in multiple types of sources (tanks, FFTF decommissioning, waste proposed for disposal at IDF and the River Protection Project Disposal Facility, and cumulative analysis sites), while in the cumulative analysis screening, the distribution of radionuclides was based only from cumulative analysis sites. This suggests the inventories in the cumulative impact waste sites were the major driver of impacts.

Screening processes conducted to evaluate the tank closure alternatives and human health impacts both considered groundwater release and intruder scenarios. However, the tank closure screening was based on the distribution of radionuclides in the BBI, which includes fewer radionuclides and chemicals than in the initial lists used for the human health screening, and would help explain the shorter screened list for tank closure (Table 1).

Table 1. Selected COPCs Based on Screening Evaluations Conducted in the TC&WM EIS

| Analyte | Analysis of Human Health Impacts (Appendix Q, Table Q-1) ^a and Cumulative Impacts (Appendix S, Table S-8) ^a | Analysis of Tank Closure Alternatives (Appendix D, Table D-2) ^a |
|----------------------|--|--|
| Radionuclides | | |
| Americium-241 | X | X ^b |
| Carbon-14 | X | X |
| Cesium-137 | X | X |

Table 1. Selected COPCs Based on Screening Evaluations Conducted in the TC&WM EIS

| Analyte | Analysis of Human Health Impacts (Appendix Q, Table Q-1) ^a and Cumulative Impacts (Appendix S, Table S-8) ^a | Analysis of Tank Closure Alternatives (Appendix D, Table D-2) ^a |
|----------------------|--|--|
| Gadolinium-152 | X | |
| Hydrogen-3 (tritium) | X | X |
| Iodine-129 | X | X |
| Neptunium-237 | X | X |
| Plutonium isotopes | X | X |
| Potassium-40 | X | |
| Strontium-90 | X | X |
| Technetium-99 | X | X |
| Thorium-232 | X | |
| Uranium isotopes | X | X |
| Zirconium-93 | X | |

a. Source: DOE/EIS-0391, *Final Tank Closure and Waste Management Environmental Impact Statement for the Hanford Site, Richland, Washington*

b. Applies to the inhalation pathway for the intruder scenario analyzed in Appendix Q but not to the EIS alternatives analysis (p. D-3).

3.2 Radionuclide Selection

The approach used to select COPCs for the current study began with development of an initial list of potentially important radionuclides based on the evaluations conducted by the 1998 CA, the 2006 Data Package, the TC&WM EIS, and available performance assessments for the 200 West LLBGs, 200 East LLBGs, ERDF, WMA C, and IDF. The list is presented in Table 2.

Table 2. Contaminants of Potential Concern Identified from Prior Analyses

| COPC | Site-Wide Analyses | | | Performance Assessments | | | | | |
|----------------------|-----------------------|--------------------------------------|-------------------------------|---------------------------------|--------------------------------------|-----------------------------|-----------------------------|-----------------------------|--|
| | CA (PNNL-11800, 1998) | 2006 Data Package (PNNL-15829, 2006) | TC&WM EIS (DOE/EIS-039, 2012) | 200-E LLBGs (WHC-EP-0645, 1995) | 200-W LLBGs (WHC-SD-WM-TI-730, 1996) | ERDF (WCH-520 Rev. 1, 2013) | WMA_C (RPP-ENV-58782, 2016) | WMA_C (RPP-ENV-58782, 2016) | IDF* (RPP-RPT-59958, Rev. B Draft in Progress, 2017) |
| Americium-241 | | | X ^b | | | | | | |
| Carbon-14 | X | X | X | X | X | X | X | X | |
| Cesium-137 | X | X | X | | | | | | |
| Chlorine-36 | X | X | | X | X | X | | | |
| Cobalt-60 | | | | | | X | | | |
| Europium-152 | X | | | | | | | | |
| Gadolinium-152 | | X | | | | | | | |
| Hydrogen-3 (tritium) | X ^c | X | X | X | X | X ^d | X | X | |
| Iodine-129 | X | X | X | X | X | X | X | X | X |
| Molybdenum-93 | | | | | | | X | | |
| Neptunium-237 | | X | X | X | X | | | | |
| Niobium-93m | | | | | | X ^d | X | | |
| Niobium-94 | | | | | | X | | | |
| Plutonium Isotopes | | X | | | | | | | |
| Polonium-209 | | | | X | | | | | |
| Potassium-40 | | | X | | | X | | | |
| Protactinium-231 | | X ^e | | X | | | | | |
| Radium-226 | | X ^f | | | | | | | |
| Radon-222 | | | | | | X | | | |
| Rhenium-187 | | | | X | X | | | | |
| Selenium-79 | X | X | X | X | X | | X | | |
| Strontium-90 | X ^e | X | X | X | X | | | | |
| Technetium-99 | X | X | X | X | X | X | X | X | X |
| Thorium-232 | | | X | | | | | | |
| Tin-116 | | | | | | | X | | |
| Uranium Isotopes | X | X | X | X | X | | X | X ^g | |

Table 2. Contaminants of Potential Concern Identified from Prior Analyses

| COPC | Site-Wide Analyses | | | Performance Assessments | | |
|--------------|-----------------------------|--|-----------------------------------|------------------------------------|---|-----------------------------------|
| | CA (PNNL-11800, 1998) | 2006 Data Package (PNNL-15829, 2006) | TC&WM EIS (DOE/EIS-0391, 2012) | 200-W LLBGs (WHC-EP-0645, 1995) | 200-E LLBGs (WHC-SD-WM-TI-730, 1996) | ERDF (WCH-520 Rev. 1, 2013) |
| Uranium-238 | X ^g | X | X | | | X |
| Zirconium-93 | | | X | | | |

References:

DOE/EIS-0391, *Final Tank Closure and Waste Management Environmental Impact Statement*
PNNL-11800, *Composite Analysis for Low-Level Waste Disposal in the 200 Area Plateau of the Hanford Site*
PNNL-15829, *Inventory Data Package for Hanford Assessments*
RPP-ENV-58782, *Performance Assessment of Waste Management Area C, Hanford Site, Washington*
RPP-RPT-59958, *Performance Assessment for the Integrated Disposal Facility, Hanford Site, Washington*
WCH-520, *Performance Assessment for the Environmental Restoration Disposal Facility, Hanford Site, Washington*
WHC-EP-0645, *Performance Assessment for the Disposal of Low-Level Waste in the 200 West Area Burial Grounds*
WHC-SD-WM-TI-730, *Performance Assessment for the Disposal of Low-Level Waste in the 200 East Area Burial Grounds*

a. 43 radionuclides were included in the IDF Performance Assessment groundwater pathway analysis. However, process model calculations focused almost entirely on I-129 and Tc-99. Impacts due to the remaining radionuclides were evaluated using the integrated system model. Only iodine-129 and technetium-99 were included here.

b. Table D-1 in the TC&WM EIS Appendix D indicates that Americium-241 applies to intruder analysis scenarios only. Appendix Q and Appendix S do not include this comment.

c. Tritium and strontium-90 were included in dose evaluations based on existing plumes, but were not included in the release and vadose zone modeling.

d. Section 4.2.2 of the ERDF Performance Assessment states that "Hydrogen-3 and niobium-93 m do not exist anywhere in the model domain in significant quantities after 1,000 years and decay to insignificant quantities (less than 1E-14 Ci per Ci source) before reaching the water table."

e. Protactinium-231 as a progeny was included in the calculation of uranium-235 dose.

f. Radium-226 as a progeny was included in the calculation of uranium-234 and uranium-238 dose.

g. The contribution of uranium and its progeny to dose was estimated by simulating uranium-238, approximating the abundance of other uranium isotopes using a single set of isotopic ratios, and assuming uranium daughter products move with the parent.

h. Table D-8 of the WMA C Performance Assessment indicates that the base case analysis included uranium-232, uranium-233, uranium-234, uranium-235, uranium-236, and the Uranium-238 daughter products, but these isotopes were not evaluated directly using the STOMP model.

The initial list in Table 2 was screened to identify key radionuclides that could potentially affect a receptor via the groundwater within 10,000 years after site closure. The initial list was also evaluated against current information on site-wide inventories and contaminant mobility. Short-lived radionuclides with a half-life of less than 10 years were screened out.

The following subsections provide information used to support the decision to include or exclude the radionuclides listed in Table 2 during the screening process. Tables 3 and 4 contain the radionuclide half-lives (DOE-STD-1196-2011, *Derived Concentration Technical Standard*) and K_d values used for the previous studies represented in Table 2. K_d values for the 200 West LLBGs Performance Assessment (PA) and the 200 East LLBGs PA were not included since radionuclides were grouped as nonsorbing, slightly sorbing, moderately sorbing, and strongly sorbing and assigned values of 0, 1, 10 and 100 mL/g, respectively.

Table 3. Half-life Values for Potentially Important Radionuclides

| COPC | Half-Life (Years) |
|----------------------|-------------------|
| Americium-241 | 432.2 |
| Carbon-14 | 5700 |
| Cesium-137 | 30.1671 |
| Chlorine-36 | 3.01e+5 |
| Cobalt-60 | 5.2713 |
| Europium-152 | 13.537 |
| Gadolinium-152 | 1.08e+14 |
| Hydrogen-3 (tritium) | 12.32 |
| Iodine-129 | 1.57e+7 |
| Molybdenum-93 | 4000 |
| Neptunium-237 | 2.14e+6 |
| Niobium-93m | 16.13 |
| Niobium-94 | 20300 |
| Plutonium-238 | 87.7 |
| Plutonium-239 | 24100 |
| Plutonium-240 | 6564 |
| Plutonium-241 | 14.35 |
| Plutonium-242 | 3.75e+5 |
| Polonium-209 | 102 |
| Potassium-40 | 1.25e+9 |
| Protactinium-231 | 32800 |
| Radium-226 | 1600 |
| Radon-222 | 0.0105 |

Table 3. Half-life Values for Potentially Important Radionuclides

| COPC | Half-Life (Years) |
|---------------|-------------------|
| Rhenium-187 | 4.12e+10 |
| Selenium-79 | 2.95e+5 |
| Strontium-90 | 28.79 |
| Technetium-99 | 2.11e+5 |
| Thorium-230 | 75400 |
| Thorium-232 | 1.41e+10 |
| Tin-126 | 2.30e+5 |
| Uranium-232 | 68.9 |
| Uranium-233 | 1.59e+5 |
| Uranium-234 | 2.46e+5 |
| Uranium-235 | 7.04e+8 |
| Uranium-236 | 2.34e+7 |
| Uranium-238 | 4.47e+9 |
| Zirconium-93 | 1.53e+6 |

Source: DOE-STD-1196-2011, *Derived Concentration Technical Standard*

Table 4. Radionuclide K_d Values from Past Studies

| COPC | 1998 CA ^a | 2006 Data Package ^b | TC&WM EIS ^c | ERDF PA ^d | WMA C PA ^e | IDF PA ^f |
|----------------------|----------------------|--------------------------------|------------------------|----------------------|-----------------------|---------------------|
| Americium-241 | 300 | | 1900 | 300 | 600 | 300 |
| Carbon-14 | 5 | 0 | 4 | 0.5 | 1 | 5 |
| Cesium-137 | 1500 | 2000 | 80 | 2000 | 100 | 2000 |
| Chlorine-36 | 0 | 0 | | 0 | | 0 |
| Cobalt-60 | 1200 | | | 10 | 0 | 2000 |
| Europium-152 | 300 | 200 | | 300 | 10 | 300 |
| Gadolinium-152 | | | 5 | | | |
| Hydrogen-3 (tritium) | 0 | 0 | 0 | 0 | 0 | 0 |
| Iodine-129 | 0.5 | 0.2 | 0 | 0.2 | 0.2 | 0.25 |
| Molybdenum-93 | | | | 0 | | |
| Neptunium-237 | 15 | 10 | 2.5 | 10 | 10 | 15 |
| Niobium-93m | 300 | | | 0 | 0 | 0 |

Table 4. Radionuclide K_d Values from Past Studies

| COPC | 1998 CA ^a | 2006 Data Package ^b | TC&WM EIS ^c | ERDF PA ^d | WMA C PA ^e | IDF PA ^f |
|--------------------|----------------------|--------------------------------|------------------------|----------------------|-----------------------|---------------------|
| Niobium-94 | 300 | | | 0 | 0 | |
| Plutonium isotopes | 200 | 600 | 150 | 600 | 600 | 150 |
| Polonium-209 | | | | | | |
| Potassium-40 | | | 15 | 0 | | |
| Protactinium-231 | 15 | | | | 300 | 15 |
| Radium-226 | 20 | | | 20 | 10 | 14 |
| Radon-222 | | | | | 0 | |
| Rhenium-187 | | | | | | |
| Selenium-79 | 0 | 5 | | 5 | 0.1 | 7 |
| Strontium-90 | 20 | 22 | 10 | 20 | 10 | 14 |
| Technetium-99 | 0 | 0 | 0 | 0 | 0 | 0 |
| Thorium-232 | 1000 | | 3200 | 1000 | 300 | 1000 |
| Tin-126 | 300 | | | 50 | 0.5 | 300 |
| Uranium isotopes | 3 | 0.8 | 0.6 | 0.8 | 0.6 | 1 |
| Zirconium-93 | 1000 | | 600 | 1000 | 300 | 1000 |

Sources:

- a. PNNL-11800, *Composite Analysis for Low-Level Waste Disposal in the 200 Area Plateau of the Hanford Site*, Appendix E, Table E.10 (K_d Best Estimates for Low Organic/Low Salts/Near Neutral).
- b. PNNL-14702, *Vadose Zone Hydrogeology Data Package for Hanford Assessments*, Table 4.11 (K_d Best estimates for low organic/low salt/near neutral, intermediate impact - sand or groundwater).
- c. DOE/EIS-0391, *Final Tank Closure and Waste Management Environmental Impact Statement*, Appendix N, Table N-2.
- d. WCH-515 Rev. 0, *Parameter Uncertainty for the ERDF Performance Assessment Uncertainty and Sensitivity Analysis*, Table 25. Best estimates for low organic/low salt/near neutral waste chemistry, not impacted sand.
- e. RPP-ENV-58782 Rev. 0, *Performance Assessment of Waste Management Area C, Hanford Site, Washington*, Table 6-11.
- f. RPP-RPT-59958, *Performance Assessment for the Integrated Disposal Facility, Hanford Site, Washington*, Table 4-33 (Best estimates for far field sand sequence with natural recharge (no impact from wastes)).

3.2.1 Radionuclides Screened from the Initial List of COPCs

Americium-241

Only the TC & WM EIS included americium-241 in the list of selected COPCs. Table D-2 in TC & WM EIS Appendix D, *Waste Inventories*, indicates that americium-241 was considered for intruder analysis scenarios only. Appendix Q and Appendix S do not include this limitation. A review of the TC & WM EIS vadose zone simulations showed that americium-241 was included in 211 of the transport simulations, but reached the water table in only one simulation with a cumulative release of only 4.6E-13 Ci.

Reported K_d values for americium-241 in the TC & WM EIS, 1998 CA, and the ERDF, WMA C, and IDF PAs ranged from 300 to 1,900 mL/g. Based on these high K_d values and minimal groundwater

impact predicted by the TC & WM EIS vadose zone simulations, americium-241 will be removed from the list of proposed COPCs.

Cesium-137

Cesium-137 was included in the list of selected COPCs for two of the past site-wide studies: the 2006 Data Package and the TC&WM EIS. Reported K_d values for cesium-137 in the TC&WM EIS, the 2006 Data Package, the 1998 CA, and the ERDF, WMA C, and IDF PAs ranged from 80 to 2,000 mL/g. PNNL-13895, *Hanford Contaminant Distribution Coefficient Database and Users Guide*, states that "Under normal Hanford conditions, Cs(I) adsorption is high with K_d values in excess of 1,000 mL/g" and "it appears that Cs(I) transport through the Hanford Site vadose zone and groundwater will be negligible except under conditions of extremely high salt concentration".

A review of the TC&WM EIS vadose zone simulations showed that cesium-137 was included in 250 of the transport simulations, but reached the water table in only three locations (Gable Mtn Pond, 216-A-5, and 218-W-2A_Burial Ground). However, the TC&WM EIS K_d of 80 mL/g appears to be based on a value from PNNL-14702, *Vadose Zone Hydrogeology Data Package for Hanford Assessments*, for IDF vitrified waste for intermediate impact sand. For the same intermediate impact sand and low organic/low salt/near neutral waste chemistry, the "best" value in PNNL-14702 is 2,000 mL/g. PNNL-14702 also states "For cesium, the best estimate K_d value selected for most Hanford impact zones and waste categories is 2,000 mL/g with a range of 200 to 10,000."

Based on the high K_d values, cesium-137 will be removed from the list of proposed COPCs.

Cobalt-60

Only the WMA C PA included cobalt-60 in the list of selected COPCs. RPP-ENV-58782, *Performance Assessment of Waste Management Area C, Hanford Site, Washington*, states that "Among radionuclides, the only contaminant producing nonzero concentrations at 100 m from the WMA C fenceline in the compliance period is ^{99}Tc . Other mobile contaminants such as ^3H , ^{60}Co , and $^{93\text{m}}\text{Nb}$ decay to insignificant quantities before reaching the water table".

PNNL-13895 states that "The general conclusions that can be drawn from these results are 1) Co(II) is highly immobile under normal Hanford groundwater conditions...".

Based on these observations and a half-life of less than 10 years, cobalt-60 will be removed from the list of proposed COPCs.

Europium-152

Only the 2006 Data Package included europium-152 in the list of selected COPCs. Reported K_d values for europium-152 in the 1998 CA, the 2006 Data Package, and the ERDF and IDF PAs ranged from 200 to 300 mL/g. For the WMA C PA, a K_d value of 10 mL/g was reported for europium-152, which was then excluded from consideration due to the K_d value being greater than 1.5 mL/g. The WMA C PA references PNNL-17154, *Geochemical Characterization Data Package for the Vadose Zone in the Single-Shell Tank Waste Management Areas at the Hanford Site*, as the source for the europium-152 10 mL/g K_d value. In PNNL-17154 the 10 mL/g K_d is assigned as the "best" value for all europium isotopes in sand size sediments under intermediate impact conditions. For the same sand size sediments under no impact conditions, the "best" K_d value is 300 mL/g.

Based on the high K_d values, europium-152 will be removed from the list of proposed COPCs.

Gadolinium-152

Only the TC & WM EIS included gadolinium-152 in the list of selected COPCs. In the inventory tables in the TC&WM EIS, only one site was reported with an inventory of gadolinium-152, 3.39×10^{-3} curies at the 218-W-3A Burial Ground. A review of the TC&WM EIS vadose zone simulations showed that gadolinium-152 did not emerge to groundwater in 10,000-year evaluation period. Because there was no impact to groundwater for the single gadolinium-152 source, gadolinium-152 will be removed from the list of proposed COPCs.

Molybdenum-93

Only the ERDF PA included molybdenum-93 in the list of selected COPCs. Since molybdenum-93 was identified as a selected COPC in only a single PA, composite impacts do not need to be evaluated. Molybdenum-93 will be removed from the list of proposed COPCs.

Niobium-93m

Niobium-93m was included in the list of selected COPCs for two PAs: the ERDF PA and the WMA C PA. WCH-520, *Performance Assessment for the Environmental Restoration Disposal Facility, Hanford Site, Washington*, states that, for the ERDF PA, “Hydrogen-3 and niobium-93m do not exist anywhere in the model domain in significant quantities after 1,000 years and decay to insignificant quantities (less than 1 E-14 Ci per Ci source) before reaching the water table.” RPP-ENV-58782 states that “Among radionuclides, the only contaminant producing nonzero concentrations at 100 m from the WMA C fenceline in the compliance period is ⁹⁹Tc. Other mobile contaminants such as ³H, ⁶⁰Co, and ^{93m}Nb decay to insignificant quantities before reaching the water table.” Based on the lack of impact at the water table for these two PAs, niobium-93m will be removed from the list of proposed COPCs.

Niobium -94

Only the ERDF PA included niobium-94 in the list of selected COPCs. Since niobium-94 was identified as a selected COPC in only a single PA, composite impacts do not need to be evaluated. Niobium-94 will be removed from the list of proposed COPCs.

Plutonium isotopes

Only the TC&WM EIS included plutonium in the list of selected COPCs. A review of the TC&WM EIS vadose zone simulations showed that plutonium-239 was included in 264 of the transport simulations, but reached the water table in only three locations (Gable Mtn Pond, 216-A-5, and 200-E-78). Total cumulative releases to the water table were 2.65E-3 Ci for Gable Mtn Pond, 1.92E-6 Ci for 216-A-5, and 1.58E-6 Ci for 200-E-78.

Reported K_d values for plutonium isotopes in the TC&WM EIS, the 1998 CA, the 2006 Data Package, and the ERDF, WMA C, and IDF PAs ranged from 150 to 600 mL/g. PNNL-13895 states that “Based on the limited data available for Pu, it appears that Pu will be fairly immobile except at very low pH values or high ethylenediaminetetraacetic acid concentrations.” Based on the high K_d values and minimal groundwater impact predicted by the TC&WM EIS vadose zone simulations, plutonium will be removed from the list of proposed COPCs.

Polonium-209

Only the 200-W LLBGs PA included polonium-209 in the list of selected COPCs. Since polonium-209 was identified as a selected COPC in only a single PA, composite impacts do not need to be evaluated. Polonium-209 will be removed from the list of proposed COPCs.

Potassium-40

Potassium-40 was included in the list of selected COPCs for the TC&WM EIS and the ERDF PA. A review of the TC&WM EIS vadose zone simulations showed that potassium-40 did not reach groundwater in any of the 10 transport simulations where it was included. WCH-520 states that, for the ERDF PA, “For K-40 and Rn-222, there is no limit calculated because K-40 occurs naturally in the soils (it was not generated during the Hanford reactor operations)...”. Based on the lack of impact to groundwater, potassium-40 will be removed from the list of proposed COPCs.

Protactinium-231

Protactinium-231 was included in the list of selected COPCs for the 2006 Data Package and the 200-W LLBGs PA. Protactinium-231 as a progeny will be included in the calculation of uranium-235 dose. Protactinium-231 will be removed from the list of proposed COPCs.

Radon-222

Only the WMA C PA included radon-222 in the list of selected COPCs. RPP-ENV-58782 indicates that radon-222 was included to complete the uranium decay chain to calculate radon flux. Since radon flux calculations will not be included as part of the CA, and radon-222 has a half-life of less than 10 years, radon-222 will be removed from the list of proposed COPCs.

Selenium-79

Selenium-79 was included in the list of selected COPCs for two of the past site-wide studies (1998 CA and 2006 Data Package), and three Performance Assessments (200 West LLBGs PA, 200 East LLBGs PA and WMA C PA). Although selenium-79 was predicted to be a groundwater dose contributor in some of the earlier studies, this would not be the case for K_d values currently considered to be appropriate for selenium-79. As discussed in Appendix A, the understanding of selenium-79 K_d has progressed over time. During the early studies, the K_d for selenium-79 was assumed to be 0 mL/g (i.e., no retardation). Current estimates, based on site-specific data, are higher, ranging from 3 to 10 mL/g (PNNL-13895), assuming low selenium concentrations and near neutral conditions. In the immediate vicinity of waste sites, the selenium K_d may be lower due to higher selenium concentrations or basic conditions, but throughout most of the vadose zone, conditions favoring the higher K_d range should apply.

Also, the SIM inventory (ECF-HANFORD-17-0079, Appendix J) for selenium-79 was less than 2.3 Ci for all historical liquid discharges included in the SIM. Based on the relatively high K_d , limited inventory, exclusion from the TC&WM EIS list of COPCs, and lack of impact for studies with higher K_d values (DOE/ORP-2000-24, *Hanford Immobilized Low-Activity Waste Performance Assessment: 2001 Version*, WCH-520, *Performance Assessment for the Environmental Restoration Disposal Facility, Hanford Site, Washington*, and RPP-RPT-59958, *Performance Assessment for the Integrated Disposal Facility, Hanford Site, Washington*), selenium-79 will be removed from the list of proposed COPCs.

Thorium-232

Only the TC&WM EIS included thorium-232 in the list of selected COPCs. A review of the TC&WM EIS vadose zone simulations showed that thorium-232 did not reach groundwater in any of the 195 transport simulations where it was included.

Reported K_d values for thorium-232 in the TC & WM EIS, the 1998 CA, and the ERDF, WMA C, and IDF PAs ranged from 300 to 3,200 mL/g. Also, the SIM inventory (ECF-HANFORD-17-0079,2017) for thorium-232 was less than one hundredth of a Ci for all historical liquid discharges included in the SIM.

Based on the high K_d values and low inventory, thorium-232 will be removed from the list of proposed COPCs.

tin-126

Only the WMA C PA included tin-126 in the list of selected COPCs. Since tin-126 was identified as a selected COPC in only a single PA, composite impacts do not need to be evaluated. Tin-126 will be removed from the list of proposed COPCs.

Zirconium-93

Only the TC&WM EIS included zirconium-93 in the list of selected COPCs. A review of the TC&WM EIS vadose zone simulations showed that zirconium-93 did not reach groundwater in any of the 170 transport simulations where it was included.

Reported K_d values for zirconium-93 in the TC & WM EIS, the 1998 CA, and the ERDF, WMA C, and IDF PAs ranged from 300 to 1,000 mL/g. Based on these high K_d values, zirconium-93 will be removed from the list of proposed COPCs.

3.2.2 Radionuclides Retained from the Initial List of COPCs

Tritium, iodine-129, neptunium-237, technetium-99, and the uranium isotopes are known leading dose contributors and, as such, will be retained in the final list of COPCs. Carbon-14, chlorine-36, and rhenium-187 were included in multiple studies where they were predicted to be groundwater dose contributors. These radionuclides will be retained in the final list of COPCs. Since strontium-90 is found in groundwater in the 200 Area in concentrations that exceed the drinking water standard (DOE/RL-2016-67), strontium-90 will be retained in the final list of COPCs. Additionally, radium-226 will be retained and thorium-230 will be added to evaluate the decay chain: uranium-234 > thorium-230 > radium-226. Table 5 shows the initial list of potential COPCs and the reason for retaining or removing each radionuclide from the final COPC list.

Table 5. COPC Screening Results

| COPC | Rationale |
|----------------------|--|
| | Retain |
| Carbon-14 | Key Contributor to Dose. |
| Chlorine-36 | Key Contributor to Dose. |
| Hydrogen-3 (tritium) | Key Contributor to Dose. |
| Iodine-129 | Key Contributor to Dose. |
| Neptunium-237 | Key Contributor to Dose. |
| Radium-226 | Added decay chain U234 > Th230 > Ra226. |
| Rhenium-187 | Dose contributor in the 200-E LLBGs PA and the 200-W LLBGs PA. |
| Strontium-90 | Current groundwater concentrations. |
| Technetium-99 | Key Contributor to Dose. |
| Uranium isotopes | Key Contributor to Dose. |

Table 5. COPC Screening Results

| COPC | Rationale |
|--------------------|---|
| Add | |
| Thorium-230 | Added decay chain U234 > Th230 > Ra226. |
| Eliminate | |
| Americium-241 | High K_d values. |
| Cesium-137 | High K_d values. |
| Cobalt-60 | Half-life less than 10 yr. |
| Europium-152 | High K_d values. |
| Gadolinium-152 | No impact to groundwater for the TC & WM EIS. |
| Molybdenum-93 | Only identified in the ERDF PA, so composite impacts do not need to be evaluated. |
| Niobium-93m | No impact to groundwater for the two PAs where niobium-93m was evaluated. |
| Niobium-94 | Only identified in the ERDF PA, so composite impacts do not need to be evaluated. |
| Plutonium isotopes | High K_d values. |
| Polonium-209 | Only identified in the 200-W LLBGs PA, so composite impacts do not need to be evaluated. |
| Potassium-40 | No impact to groundwater for the TC&WM EIS or ERDF PA. |
| Protactinium-231 | Protactinium-231 will be included as a progeny in the calculation of uranium-235 dose. |
| Radon-222 | Half-life less than 10 yr. |
| Selenium-79 | Relatively high K_d , limited inventory, and lack of impact for studies with higher K_d values. |
| Thorium-232 | High K_d values. |
| Tin-126 | Only identified in the WMA C PA, so composite impacts do not need to be evaluated. |
| Zirconium-93 | High K_d values. |

4 Summary

This document describes the screening and selection process for radionuclides to include in the Hanford Site CA. This screening approach was based on methods adopted in three prior site-wide studies: the 1998 CA, the 2006 Data Package, and the TC&WM EIS. Sixteen radionuclides (Table 6) were selected for the Hanford Site CA groundwater pathway evaluation.

Table 6. Selected Contaminants for Groundwater Pathway Detailed Evaluation

| Contaminant |
|-------------|
| Carbon-14 |
| Chlorine-36 |

Table 6. Selected Contaminants for Groundwater Pathway Detailed Evaluation

| Contaminant |
|----------------------|
| Hydrogen-3 (tritium) |
| Iodine-129 |
| Neptunium-237 |
| Radium-226 |
| Rhenium-187 |
| Strontium-90 |
| Technetium-99 |
| Thorium-230 |
| Uranium-232 |
| Uranium-233 |
| Uranium-234 |
| Uranium-235 |
| Uranium-236 |
| Uranium-238 |

5 References

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Appendix A

Selenium-79 Review

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A1 Documented Se-79 K_d Values

Appendix B (Selenium-79 K_d Values) lists the distribution coefficients (K_{ds}) in the DOE/EIS-0391, *Final Tank Closure and Waste Management Environmental Impact Statement for the Hanford Site, Richland, Washington* (hereinafter referred to as the TC&WM EIS), composite analyses (CAs), performance assessments (PAs), and related or referenced documents. Appendix B includes only those K_d values that were identified as sand size or with no size description; silt-size, gravel corrected, and carbonate-dominated values were not included. There is a progression from an assumed value of 0 mL/g in the early documents (WHC-EP-0645, *Performance Assessment for the Disposal of Low-Level Waste in the 200 West Area Burial Grounds*, hereinafter referred to as the 200-W LLBGs PA; WHC-SD-WM-TI-730, *Performance Assessment for the Disposal of Low-Level Waste in the 200 East Area Burial Grounds*, hereinafter referred to as the 200-E LLBGs PA; and PNNL-11800, *Composite Analysis for Low-Level Waste Disposal in the 200 Area Plateau of the Hanford Site*, hereinafter referred to as the 1998 CA) to a value of 4 mL/g (based on site specific data [DOE/ORP-2000-24, *Hanford Immobilized Low-Activity Waste Performance Assessment: 2001 Version*, hereinafter referred to as the 2001 ILAW PA]), to a “best” non-impacted value of 5 or 7 (depending on the document) for the later documents. Table A-1 lists the least impacted (by waste chemistry) “best” Se-79 K_d values found for each of the studies being reviewed. At first glance, the Waste Management Area (WMA) C PA looks to be an exception with a value of 0.1 mL/g. However, following the references (see Section A1.5) shows that this value is for the intermediate impact zone.

Table A-1. Selenium K_d Values for the EIS, CAs, and PAs

| COPC | 1998 CA | 2006 Data Package | 200-W LLBGs PA | 200-E LLBGs PA | TC&W M EIS | ERDF PA | WMA C PA | IDF PA |
|-------------|---------|-------------------|----------------|----------------|------------|---------|----------|--------|
| Selenium-79 | 0 | 5 | 0 | 0 | Not Listed | 5 | 0.1 | 7 |

References:

- DOE/EIS-0391, *Final Tank Closure and Waste Management Environmental Impact Statement*.
- PNNL-11800, *Composite Analysis for Low-Level Waste Disposal in the 200 Area Plateau of the Hanford Site*.
- PNNL-14702, *Vadose Zone Hydrogeology Data Package for Hanford Assessments*.
- RPP-ENV-58782, *Performance Assessment of Waste Management Area C, Hanford Site, Washington*.
- RPP-RPT-59958, *Performance Assessment for the Integrated Disposal Facility, Hanford Site, Washington*.
- WCH-515, *Parameter Uncertainty for the ERDF Performance Assessment Uncertainty and Sensitivity Analysis*.
- WHC-EP-0645, *Performance Assessment for the Disposal of Low Level Waste in the 200 West Area Burial Grounds*.
- WHC-SD-WM-TI-730, *Performance Assessment for the Disposal of Low Level Waste in the 200 East Area Burial Grounds*.

PNNL-13895, *Hanford Contaminant Distribution Coefficient Database and Users Guide*, provides the following summary regarding selenium:

Selenium. A fair number of Se(VI) K_d values have been determined using natural Hanford sediment and are listed in Table 14. These results indicate that at trace concentrations, adsorption of Se(VI) to Hanford sediment is low to moderate with K_d values ranging from 3 to 10 mL/g. At higher Se(VI) concentrations, the K_d values are lower (0 to 3 mL/g). Acidic conditions increase Se(VI) adsorption, and basic conditions reduce adsorption. This is consistent with the anionic character of Se(VI).

Most of the selenium concentrations in the vadose zone would likely be “trace concentrations”, with a K_d of 3 to 10 mL/g based on PNNL-13895.

A1.1 200-W LLBGs PA, 200-E LLBGs PA, and 1998 CA

The 200-W LLBGs PA, 200-E LLBGs PA, and 1998 CA predicted that Se-79 will be a dose contributor. Figures A-1 and A-2 show that groundwater dose for Se-79 exceeded the Tc-99 dose for the 200-W LLBGs PA and the 200-E LLBGs PA evaluations. For the 1998 CA, the Se-79 cumulative release is only slightly less than U-238 and more than C-14 and I-129 (Table A-2). All three of these early studies had a Se-79 K_d of 0 mL/g.

| 200-W LLBGs C PA (WHC-EP-0645) 1995 | | | | | |
|--|-----------------------------|----------------------------|---------------------------------|-------------|---------|
| Radionuclide | All-pathways dose (mrem/yr) | Groundwater dose (mrem/yr) | Population dose (person-rem/yr) | Dose ratios | |
| | | | | All path/GW | Pop./GW |
| ³ H | 4.1 E-03 | 3.6 E-03 | NA | 1.1 | NA |
| ¹⁴ C | 2.7 E+02 | 1.2 E+02 | 20 | 2.2 | 0.17 |
| ³⁶ Cl | 3.0 E+03 | 1.7 E+02 | 360 | 17 | 2.1 |
| ⁷⁹ Se | 8.5 E+02 | 4.8 E+02 | 78 | 1.8 | 0.16 |
| ⁹⁹ Tc | 2.5 E+02 | 7.6 E+01 | 20 | 3.3 | 0.26 |
| ¹²⁹ I | 2.9 E+04 | 1.6 E+04 | 2100 | 1.8 | 0.13 |
| ¹⁸⁷ Re | 7.0 E-01 | 4.8 E-01 | 0.051 | 1.5 | 0.11 |
| ²³⁷ Np | 1.6 E+03 | 1.4 E+03 | 110 | 1.1 | 0.076 |
| ²⁰⁹ Po | 3.3 E+01 | 2.9 E+01 | 2.3 | 1.1 | 0.078 |
| ²³¹ Pa | 4.2 E+04 | 3.9 E+04 | 2900 | 1.1 | 0.076 |
| U | 1.6 E+04 | 1.4 E+04 | 1100 | 1.1 | 0.080 |

Figure A-1. Table 4-22 of the 200-W LLBGs PA

200-E LLBGs C PA (WHC-SD-WM-TI-730)
1996

Table 4-14. Radionuclide Dose Estimates for Groundwater Pathways.*

| Radionuclide | Drinking Water Dose (mrem/yr) | All-Pathways Dose (mrem/yr) | Dose ratio (All-Pathways/groundwater) |
|-------------------|-------------------------------|-----------------------------|---------------------------------------|
| ³ H | 0.11 | 0.12 | 1.1 |
| ¹⁴ C | 1,100 | 2,400 | 2.2 |
| ³⁶ Cl | 1,510 | 26,700 | 17 |
| ⁷⁹ Se | 4,200 | 7,400 | 1.8 |
| ⁹⁹ Tc | 650 | 2,100 | 3.3 |
| ¹²⁹ I | 141,000 | 292,000 | 1.8 |
| ¹⁸⁷ Re | 4 | 6 | 1.5 |
| ²³⁷ Np | 1,964,000 | 2,216,000 | 1.1 |
| U | 1,209,000 | 1,330,000 | 1.1 |

*These doses are from the base-case analysis assuming a 1-Ci inventory per radionuclide and Category 1 infiltration conditions (5 cm/yr). The values does not represent actual inventory projections and associated doses.

Figure A-2. Table 4-14 of the 200-E LLBGs PA

Table A-2. 1998 CA Cumulative Release from All Sources to the Water Table from 1940 to 3000.

| Radionuclide | Activity (Ci) |
|--------------|---------------|
| Tc-99 | 1460 |
| Cl-36 | 11.67 |
| U-238 | 8.67 |
| Se-79 | 8.03 |
| C-14 | 4.58 |
| I-129 | 2.1 |

Note: Estimated from Figures 4.5 to 4.10 from PNNL-11800, *Composite Analysis for Low-Level Waste Disposal in the 200 Area Plateau of the Hanford Site*

A1.2 2006 CA (Incomplete)

PNNL-15829, *Inventory Data Package for Hanford Assessments*, references the 2001 ILAW PA as the “primary source of the selection data” regarding the inclusion of Se-79 in the Hanford Assessments.

A review of the 2001 ILAW PA shows that Se-79 was not a significant contributor to dose within the 10,000-year evaluation period.

The K_d used for Se-79 changed from 0 mL/g in the 1998 ILAW PA to 4.0 mL/g in the 2001 ILAW PA. Section 4.3.6 of the 2001 ILAW PA states:

“In the 1998 ILAW performance assessment (Mann 1998a), the most restrictive impact was caused by the drinking water dose from beta- and photon-emitting radionuclides. At 10,000 years, this dose was calculated to be 2.0 mrem in a year resulting mainly from ^{99}Tc (75 percent) and ^{79}Se (20 percent). This performance assessment shows much lower numbers at 10,000 years (0.010 mrem/y). The highest value calculated for the beta/photon drinking water dose is 0.013 mrem/year at about 76,500 years. Table 4-4 shows the major contributions at 1,000 years and 10,000 years to the estimated beta and photon drinking water dose at a well 100 m downgradient from the disposal facility. Figure 4-17 shows the time dependence. In this assessment, ^{99}Tc is still the most important radionuclide, contributing approximately 58 percent of the dose at 1,000 and 10,000 years. However, the next most important radionuclide is ^{129}I , which contributes approximately 42 percent at 1,000 and 10,000 years. **The switch of selenium and iodine is a direct result of site specific data increasing selenium's K_d from 0. to 4.0 mL/g and decreasing iodine's K_d from 3.0 to 0 mL/g (see Section 3.4.3.3).**”

Section 7.7 of the 2001 ILAW PA states:

“In the 1998 ILAW PA, ^{79}Se was assumed to be mobile because no Hanford Site-specific data were available that indicated otherwise. Since then, we have learned that the half-life of ^{79}Se is longer than was believed. Also, **disposal-site specific information has shown that selenium transport in the vadose zone is chemically retarded.**”

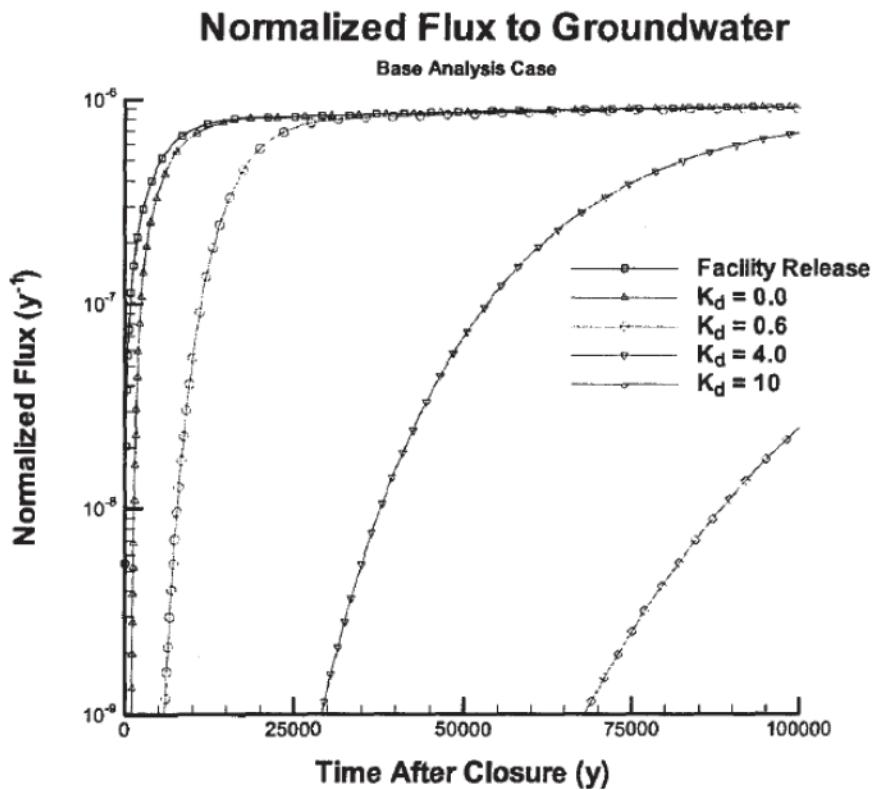
Because of this change in the K_d value, Se-79 does not reach the water table within 10,000 years for the 2001 ILAW PA Base Analysis Case. Section 4.3.4 of the 2001 ILAW PA states:

“Figure 4-10 shows the contaminant flux summed over horizontal distance as a function of time and K_d bin. Only the mobile contaminants reach the groundwater during the time of compliance (the first 1,000 years). At 10,000 years, the slightly retarded contaminants ($K_d = 0.6 \text{ mL/g}$) also are beginning to reach the groundwater, but their inventory-normalized contribution is still approximately one order of magnitude less than the mobile contribution. **Higher K_d contaminants ($K_d \geq 4 \text{ mL/g}$) do not contribute to the estimated doses at 10,000 years and are even less important.**”

Figure A-3 is a copy of Figure 4-10 from the 2001 ILAW PA. It shows that, for the Base Analysis Case, radionuclides with a K_d of 4.0 mL/g (including Se-79) do not reach the water table until after 25,000 years.

Figure 4-10. Inventory-Normalized Contaminant Flux Summed Over Horizontal Distance as a Function of Time and K_d Bin (Logarithmic Scale).

“Facility Release” refers to values just below the disposal facility.



Source: DOE/ORP-2000-24, *Hanford Immobilized Low-Activity Waste Performance Assessment: 2001 Version*

Figure A-3. Figure 4-10 of the 2001 ILAW PA

Based on these observations, Se-79 should not have been included in the 2006 Data Package screened radionuclide list using the 2001 ILAW PA as a basis for selecting Se-79 since, with its K_d of 4.0 mL/g, Se-79 would not have reached the water table within 10,000 years under the 2001 ILAW PA Base Analysis Case. Also, the K_d of 5 mL/g selected for Se-79 for the incomplete 2006 CA and documented in PNNL-14702, *Vadose Zone Hydrogeology Data Package for Hanford Assessments*, should result in minimal to no impact to the groundwater from Se-79.

A1.3 TC&WM EIS

Radionuclide screening for the TC&WM EIS is discussed in three of the appendices: Appendix D, Appendix Q, and Appendix S. The next three paragraphs show excerpts from these appendices.

Appendix D:

“The BBI includes quantity estimates of 46 radionuclides and 24 chemical constituents. Not all constituents are important in the exposure scenarios used to assess TC & WM EIS alternative implementation impacts. Thus, to focus attention on the constituents that

control the impacts, DOE performed an initial screening analysis. For radionuclides, groundwater release and direct intrusion scenarios were considered. For the groundwater release screening scenario, only drinking water consumption was considered. Release was assumed to be partition limited, and decay during transport was considered. For the direct intrusion screening scenario, inadvertent soil ingestion and inhalation pathways were considered.

The analysis estimated relative impacts based on distribution of radionuclides in the BBI for all tanks. Radionuclides contributing less than 1 percent of impacts under intruder or well scenarios were eliminated from the detailed analysis.”

“The screening of the BBI for the groundwater scenarios resulted in reduction of the original set of 46 radionuclides and 24 chemical constituents to a final set of ten radionuclides and ten chemical constituents that was used in the analysis of the tank waste. However, a screening of the cumulative impacts analysis data resulted in the addition of other COPCs that are not included in Table D-2. Appendix Q provides details on this screening.”

Appendix Q:

“The process of impacts analysis is iterative in nature, with execution of initial passes through the steps at a high level so as to screen out less important conditions and produce a manageable set of scenarios for analysis. An initial iteration through the procedure was used to establish the number of constituents to be included in the analysis. For radionuclides in this screening analysis, groundwater release and direct intrusion scenarios were considered. For the groundwater release screening scenario, only drinking water consumption was considered, release was assumed to be partition limited, and decay during transport was considered. For the direct intrusion scenario, inadvertent soil ingestion and inhalation pathways were considered. The analysis involved estimation of relative impacts based on the distribution of radionuclides in all tanks; FFTF decommissioning; waste proposed for disposal at IDF-East, IDF-West, and the RPPDF; and contamination in place at cumulative analysis sites. In reviewing constituents at a given source area, radionuclides contributing in combination less than 1 percent of impacts for intruder or well scenarios were not included in the detailed analysis.”

“The screening resulted in reduction of the original set of radioactive and chemical constituents to a final set of 14 radioactive and 26 chemical constituents, which represents both alternatives and cumulative impact sources.”

Appendix S:

“The initial list of radionuclides included those with half-lives greater than 10 years, and the initial list of chemicals included those with a health risk from ingestion—that is, they have maximum contaminant levels or are listed in the Integrated Risk Information System as having health-based ingestion standards. Not all the radioactive and chemical constituents on the initial list are important in exposure scenarios used to assess cumulative impacts in this TC & WM EIS. Therefore, to focus attention on constituents that control impacts, an additional screening analysis was performed. The primary focus of that analysis was to consider groundwater release scenarios for cumulative impacts analysis sources and to ensure consistency with the screening done for the alternatives

analysis, allowing for cumulative impacts to be added to the alternatives impacts. For radionuclides, only groundwater consumption was considered, release was assumed to be partition limited, and decay during transport was considered. For analysis purposes, estimation of relative impacts was based on the distribution of radionuclides in the cumulative impacts inventory. Radionuclides contributing less than 1 percent of impacts under well scenarios were eliminated from the detailed analysis.”

“As indicated in Table S-8, the screening resulted in reduction of the original set of radioactive and chemical constituents to a final set of 14 radioactive and 26 chemical constituents, which includes those constituents also identified for the alternatives impacts analysis. Appendix Q of this TC & WM EIS provides further description of the screening process for the radioactive and chemical constituents identified for the groundwater analysis.”

All three appendices state that radionuclides contributing less than 1 percent of impacts were eliminated from the detailed analysis. This statement indicates that selenium-79 was determined to contribute little to no impact under the assumptions of the TC&WM EIS.

A1.4 ERDF PA

WCH-520, *Performance Assessment for the Environmental Restoration Disposal Facility, Hanford Site, Washington* (hereinafter referred to as the ERDF PA), used a Se-79 K_d of 5 mL/g. Se-79 does not reach the water table within the 10,000-year period for the ERDF PA compliance case evaluation, as noted in Section 4.2.3 of the ERDF PA:

“During the post-compliance period 1,000 to 10,000 years after closure, chlorine-36, technetium-99, niobium-94, molybdenum-93, and iodine-129 breakthrough at the point of compliance (100 m downgradient of the ERDF) as shown in Figure 4-9. Iodine-129 is the only radionuclide with a K_d value greater than zero to do so.”

This can be seen in Figure 4-9 of WCH-520.

A1.5 WMA C PA

Figure A-4 shows that RPP-ENV-58782, *Performance Assessment of Waste Management Area C, Hanford Site, Washington* (hereinafter referred to as the WMA C PA) predicted a Se-79 peak groundwater concentration that was half that of U-238 and over twice the I-129 concentration. The WMA C PA states that “The K_d values are chosen assuming low-salt, near-neutral waste chemistry in the vadose and saturated zone.” The Se-79 K_d of 0.1 mL/g used for the WMA C PA is referenced to PNNL-17154, *Geochemical Characterization Data Package for the Vadose Zone in the Single-Shell Tank Waste Management Areas at the Hanford Site*. This value applies to the “Best” value for sand size sediments in the intermediate impact zone (see Tables 3.3, 3.7, 3.9, 3.13, 3.17, 3.21, and 3.23 of PNNL-17154). PNNL-17154 defines the intermediate zone as “Zones in which the acidic or basic nature of the wastes was expected to have been largely neutralized by reaction with the natural sediment.” Table 3.4 in PNNL-17154 lists a Se-79 “Best” K_d of 5 mL/g for sand size sediments in natural pore waters/groundwater. This value is in the range given by PNNL-13895, agrees with the value used in the ERDF PA, and is similar to the value used in RPP-RPT-59958, *Performance Assessment for the Integrated Disposal Facility, Hanford Site, Washington* (hereinafter referred to as the IDF PA).

Also, Table 4.11 in PNNL-14702 gives a “Best” Se-79 K_d value of 5 mL/g for sand size sediments in the intermediate impact zone assuming a low-salt/near-neutral waste chemistry.

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| Radionuclide or Nonradiological Contaminant | Nominal K_d value (mL/g) | Maximum Concentration during Compliance Time Frame (pCi/L) | Point of Calculation where Maximum Concentration Occurs | Years after Closure of Maximum Concentration | Maximum Concentration during Sensitivity/Uncertainty Time Frame (pCi/L) | Point of Calculation where Maximum Concentration Occurs |
|---|----------------------------|--|---|--|---|---|
| Iodine-129 | 0.2 | 0 | — | 6,540 | 0.004 | PoCal 4 |
| Selenium-79 | 0.1 | 0 | — | 3,770 | 0.01 | PoCal 5 |
| Tin-126 | 0.5 | 0 | — | 10,000 | 0.05 | PoCal 5 |
| Technetium-99 | 0 | 0.1 | PoCal 5 | 1,550 | 30 | PoCal 4 |
| Uranium-238 | 0.6 | 0 | — | 10,000 | 0.02 | PoCal 3 |

1

Figure A-4. Table 7-3 of the WMA C PA

A1.6 IDF PA

Table 4-33 in IDF PA lists Se-79 “reasonably conservative” and “best” K_d values of 1 and 2 mL/g for chemically impacted far field in sand, and values of 3 and 7 mL/g for far field in sand with natural recharge (i.e., no impact from wastes). These K_d values are referenced to PNNL-13037, *Geochemical Data Package for the 2005 Hanford Integrated Disposal Facility Performance Assessment*. Based on Figure 1-4 in the IDF PA (shown here as Figure A-5), Se-79 does not reach the water table within the 10,000-year period.

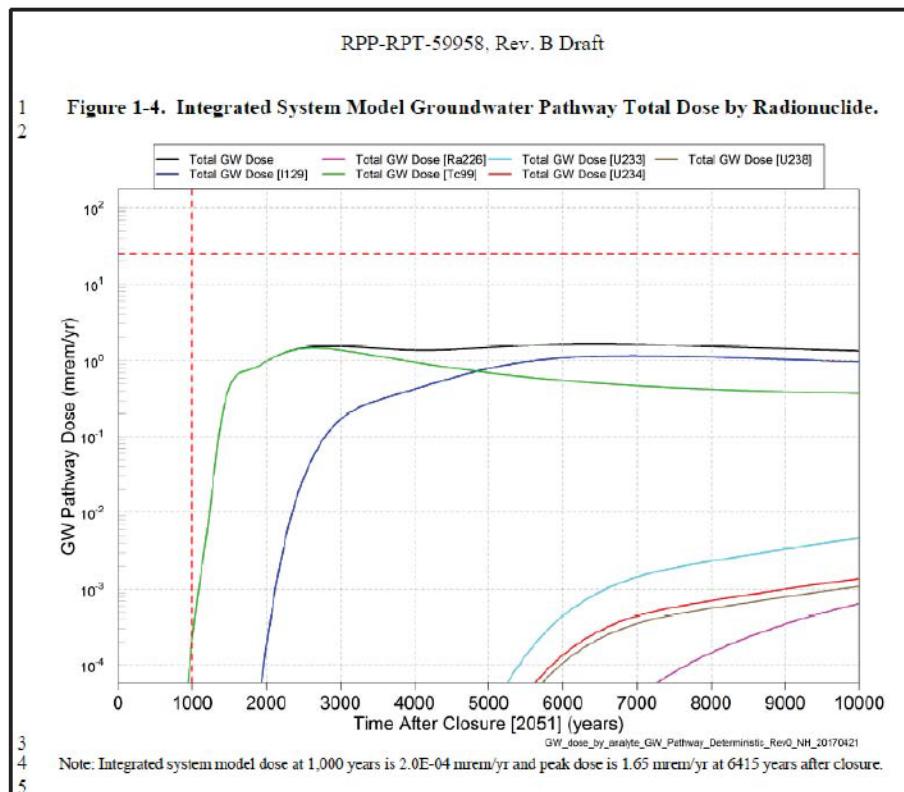


Figure A-5. Figure 1-4 of the IDF PA

A2 Effects of pH and Ionic Strength on Selenium K_d

PNNL-11964, *Effects of High-pH and High-Ionic Strength Groundwater on Iodide, Perchlorate, and Selenate Sorption to Hanford Sediments*, and PNNL-14325, *The Influence of Glass Leachate on the Hydraulic, Physical, Mineralogical and Sorptive Properties of Hanford Sediment*, looked at the effects of increased pH and high ionic strength due to leachate from the waste. Table 4 in PNNL-11964 shows that increasing ionic strength, while maintaining pH at approximately 7.7, did not have a large impact on the selenium K_d , showing a small increase with higher ionic strength. Increasing pH from 8.1 to 11.9 sharply decreased selenium K_d from 5.78 ± 0.28 to 0.04 ± 0.00 mL/g, with most of the drop occurring between pH 8.1 and pH 9.9 (Table 7 in PNNL-11964).

The results of the PNNL-14325 batch sorption study (Table 3.14) show Se-79 K_d values for time 0 that increased with increasing ionic strength. Table 3.14 also shows that there was no sorption after time 0 (10, 90, 180, and 360 days). However, Figure 3.42 in PNNL-14325 shows that, after time 0, the lowest measured pH was about 9.5. This appears to be in agreement with the PNNL-11964 results. PNNL-14325 notes that “as the pH of the glass leachate is neutralized by reactions with the vadose zone sediments, or certainly by the time vadose zone pore water reaches the water table, there would appear to be some adsorption potential for selenate (including ^{79}Se).”

Um and Serne, 2004, “Sorption and Transport Behavior of Radionuclides in the Proposed Low-Level Radioactive Waste Disposal Facility at the Hanford Site,” compared Se-75 (as an analog for Se-79) K_d values for three different Hanford sediments using uncontaminated Hanford groundwater and simulated glass leachate spiked with individual radionuclides. Table 3 of Um and Serne (2004) shows that K_d values for the three tests with uncontaminated groundwater ranged from 7.14 ± 0.18 to 8.65 ± 0.34 mL/g (pH from 7.6 to 7.7). For the three tests with simulated glass leachate, K_d values ranged from 1.08 ± 0.09 to 2.68 ± 0.12 mL/g (pH from 8.9 to 9.0).

PNNL-13037 includes the following discussion regarding Se-79 K_d values:

“In 1998, little Hanford-specific data existed for the adsorption properties of selenium (as selenate or selenite). For the 1998 ILAW PA, it was, therefore, recommended that the K_d values for ^{79}Se be set at 0 mL/g. Between 1998 and 2001, batch K_d studies (Kaplan et al. 1998c) were completed using several Hanford sediments, including IDF borehole 299-E17-21. The solution used in these measurements was uncontaminated groundwater, and the sediments were dominated by sand-sized particles. Kaplan et al. (1998b)² also studied the adsorption of ^{75}Se , as a surrogate for ^{79}Se , from Hanford groundwaters with pH values that had been adjusted to higher than normal values. The measurements suggest that some significant adsorption of selenate would be expected for both groundwater and higher pH solutions. Thus, for the 2001 ILAW PA, the “most probable” K_d value for selenium was chosen as 4 mL/g.

More recent work by Kaplan et al. (2003) indicates that selenate adsorption to Hanford sediments is nil for highly alkaline solutions. This is consistent with geochemical principles (see discussion in EPA 1999a and references therein) that suggest that anionic species, such as selenite and selenate, should show reduced sorption at greater-than-neutral pH conditions onto any sediment containing minerals with variably charged adsorption surface sites, such as iron and aluminum hydroxide minerals and particle coatings.

² Kaplan 1998b is PNNL-11966, *Radionuclide Distribution Coefficients of Sediments Collected from Borehole 299-E17-21: Final Report for Subtask 1a*.

Um and Serne^(a) used an uncontaminated groundwater and a simulated glass leachate based on the composition for the long-term, steady-state chemical composition of glass leachate and vadose zone pore water predicted by the STORM code for the 2001 ILAW PA (see Table 6.2) to study selenate adsorption onto three samples of Hanford formation sediments from another IDF borehole (299-E24-21 [ILAW borehole #2 – C3177]). The K_d values measured by Um and Serne for selenate are described in Section 3.5.3. These tests also contained a trace amount of stable selenate (few parts per billion) that was not present in the earlier studies by Kaplan et al. (1998b, c). Because these earlier studies used only the carrier-free ^{75}Se isotope (which essentially means the mass of selenium present was infinitesimal), we later became concerned that the K_d results might be biased high by not having some selenium mass present. The most recent results by Um and Serne^(a) corroborate the selenium K_d values obtained by Kaplan et al. (1998c) for natural groundwater and Hanford sediments, but do indicate that selenium K_d values for more alkaline solutions, including simulated glass leachate, are considerably smaller than 4 mL/g, the value recommended in 2001. Thus, for the 2005 IDF PA, we changed (decreased) the K_d value for ^{79}Se for the chemically impacted zones, where the glass leachate forces the pore fluid pH to be elevated above background. No changes were made to the K_d values for selenium for the near field concrete-impacted zone. During preparation of this data package, we determined that for the 2001 data package, the recommended K_d values for the chemically impacted gravel zone had inadvertently not been reduced by the factor of 10 to account for the assumed 90% gravel content. Thus, there is a change (correction) to the recommended 2005 K_d values for this zone.

Um and Serne^(a) measured the K_d for selenate in simulated glass leachate onto IDF borehole sediments (see Section 3.5.3) and consistently found non-zero K_d values for selenium for six tests. Their values for the simulated glass leachate ranged from 1 to 3 mL/g with good precision. At long time periods, we assume that glass weathering products will adsorb some selenium.

Therefore, we recommend that a non-zero K_d is appropriate and chose a K_d value of 1 mL/g for selenate for the long-term near-field zone. Based on the results of Um and Serne, we also are more confident that the “most probable” K_d for ^{79}Se for the chemically unaltered pore water/groundwater fluid can be increased from 4 to 7 mL/g (see Table 6.1). We have not tested selenium adsorption on Hanford sediments that contain significant quantities of gravel-sized material. We, therefore, rely on the conservative gravel-correction factor (see Equation 2.6) and assume that the gravel-dominated sequence at the bottom of the vadose zone and at the upper unconfined aquifer has 90% gravel. This effectively reduces the recommended K_d values in gravel zones by a factor of 10 as listed in Table 6.1. To build in further conservatism, the chemically impacted gravel values were reduced further. It is likely that the chemically impacted sand zone controls the travel time of selenium in the PA calculations.”

Based on these studies, it appears that our choice of Se-79 K_d will be dependent on the expected vadose zone pH beneath the source zones.

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Appendix B

Selenium-79 K_d Values

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B1 Introduction

Table B-1 summarizes the Se-79 distribution coefficient (K_d) values found in DOE/EIS-0391, *Final Environmental Impact Statement Tank Closure and Waste Management for the Hanford Site, Richland, Washington*, composite analyses, performance assessments, and related or referenced documents. The table includes only the Se-79 K_d values that were identified as sand size or with no size description; silt-size, gravel corrected, and carbonate-dominated values were not included. Blue shading indicates K_d s that were listed as non-impacted and/or groundwater (assuming little to no impact for groundwater). K_d s that were identified as intermediate impact or chemically impacted far field were shaded green. K_d s that were identified as high impact or near field were shaded tan. Values are arranged in document date order.

The table includes document number, year published, location in the document, waste chemistry, particle size, impact zone, K_d estimates (Conservative, Best, Min, and Max), and any notes included with the K_d estimates (Comments column). The text “---” indicates that information was not included in the source document.

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Table B-1. Documented Selenium K_d Values

| Document | Location | Year | Waste Chemistry | Size | Impact Zone | Conservative | Best | Min | Max | Comments |
|-------------------|-------------|------|------------------------------------|------|--------------------------|--------------|------|-------|-------|---|
| WHC-EP-0645 | Table 3-5 | 1995 | --- | --- | All zones | --- | 0 | --- | --- | |
| WHC-SD-WM-TI-730 | Table 3-6 | 1996 | --- | --- | All zones | --- | 0 | --- | --- | K_d values are reasonably conservative estimates based on many references including Serne and Wood (1990). |
| PNNL-11800 | Table E.5 | 1998 | High Organic/Very Acidic | --- | High Impact | 0 | 0 | 0 | 0 | K_d values were estimated. |
| PNNL-11800 | Table E.6 | 1998 | High Organic/Very Acidic | --- | Intermediate Impact | 0 | 0 | 1 | 1 | Selenium is a soluble anion. The K_d value was estimated. |
| PNNL-11800 | Table E.10 | 1998 | High Organic/Very Acidic | --- | Groundwater | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.6 | 1998 | High Organic/Near Neutral | --- | High Impact | 0 | 0 | 1 | 1 | Selenium is a soluble anion. The K_d value was estimated. |
| PNNL-11800 | Table E.6 | 1998 | High Organic/Near Neutral | --- | Intermediate Impact | 0 | 0 | 1 | 1 | Selenium is a soluble anion. The K_d value was estimated. |
| PNNL-11800 | Table E.10 | 1998 | High Organic/Near Neutral | --- | Groundwater | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.8 | 1998 | Very High Salt/Very Basic | --- | High Impact | 0 | 0 | 0 | 0.2 | Technetium, carbon, iodine, selenium, and chlorine are anionic... |
| PNNL-11800 | Table E.9 | 1998 | Very High Salt/Very Basic | --- | Intermediate Impact | 0 | 0 | 0 | 4 | Selenium is anionic. The K_d values were estimated. |
| PNNL-11800 | Table E.10 | 1998 | Very High Salt/Very Basic | --- | Groundwater | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.11 | 1998 | Chelates/High Salts | --- | High Impact | 0 | 0 | 0 | 0.5 | Technetium, iodine, selenium, had chlorine are anions... |
| PNNL-11800 | Table E.11 | 1998 | Chelates/High Salts | --- | Intermediate Impact | 0 | 0 | 0 | 0.5 | Technetium, iodine, selenium, and chlorine are anions... |
| PNNL-11800 | Table E.12 | 1998 | Low Organic/Low Salts/Acidic | --- | High Impact | 0.1 | 0.2 | 0.1 | 2 | Anions sorb to iron oxides and kaolinite at lower pH levels. |
| PNNL-11800 | Table E.10 | 1998 | Low Organic/Low Salts/Acidic | --- | Intermediate Impact | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.10 | 1998 | Low Organic/Low Salts/Acidic | --- | Groundwater | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.10 | 1998 | Low Organic/Low Salts/Near Neutral | --- | High Impact | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.10 | 1998 | Low Organic/Low Salts/Near Neutral | --- | Intermediate Impact | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11800 | Table E.10 | 1998 | Low Organic/Low Salts/Near Neutral | --- | Groundwater | 0 | 0 | -3.44 | 0.78 | In the Hanford groundwater/sediment system, the K_d values range from -3.44 to 0.78 mL/g (Serne et al., 1993). |
| PNNL-11966 | Table 4 | 1998 | Non-impacted | --- | Groundwater | 3.8 | 6.7 | 3.75 | 10.85 | |
| DOE/ORP-2000-24 | Table 3-5 | 2001 | chemically impacted | Sand | far field | --- | 4 | --- | --- | |
| PNNL-13895 | Section 5.3 | 2003 | --- | --- | Trace Se concentrations | --- | --- | 3 | 10 | |
| PNNL-13895 | Section 5.3 | 2003 | --- | --- | Higher Se concentrations | --- | --- | 0 | 3 | (10) measured the K_d for selenate in synthetic glass leachate onto IDF borehole sediments and found nonzero values consistently for six tests. Values ranged from 1 to 3 mL/g with good precision. |
| PNNL-13037 Rev. 2 | Table 5.1 | 2004 | Vitrified Waste | --- | Near Field | 0 | 1 | 0 | 3 | |

Table B-1. Documented Selenium K_d Values

| Document | Location | Year | Waste Chemistry | Size | Impact Zone | Conservative | Best | Min | Max | Comments |
|-------------------|------------|------|--|------|----------------------------------|--------------|------|-----|-----|--|
| PNNL-13037 Rev. 2 | Table 5.3 | 2004 | Cementitious Secondary Wastes - Young Concrete | -- | Near Field | 1 | 2 | 1 | 800 | Estimated. Dominant species for Se and Ru were assumed to be SeO_4^{2-} and RuO_4^{2-} , respectively [19]. Sulfate may be used as an analog for selenite chemical behavior in concrete. Sulfate (or sulfite) is often included in concrete mixes, and therefore it would be expected to be retained strongly by concrete, primarily by coprecipitation constraints. Selenite adsorption, independent of precipitation processes, would be expected to be rather large. |
| PNNL-13037 Rev. 2 | Table 5.3 | 2004 | Cementitious Secondary Wastes - Moderately Aged Concrete | -- | Near Field | 1 | 2 | 1 | 100 | --- |
| PNNL-13037 Rev. 2 | Table 5.3 | 2004 | Cementitious Secondary Wastes - Aged Concrete | -- | Near Field | 0 | 1 | 0 | 300 | --- |
| PNNL-13037 Rev. 2 | Table 5.5 | 2004 | Chemically Impacted | Sand | Far Field | 1 | 2 | 0 | 10 | Anionic. K_d measured at the ILAW/IDF site had K_d values of $6.7 \pm 0.4 \text{ mL/g}$ [14]. Results of a Se sorption experiment to Hanford sediments in high ionic strength (NaOH and NaOCl) indicate $\text{Se} K_d$ values range from 0 to 18 mL/g ; but values for 0.03 NaOH are 0 mL/g and are beyond the causticity of probable glass leachates [16]. K_d values will be chosen from recent tests on IDF borehole sediments with synthetic glass leachate that yielded K_d values which ranged from 1 to 3 mL/g [15]. |
| PNNL-13037 Rev. 2 | Table 5.6 | 2004 | Natural Recharge (no impact from wastes) | Sand | Far Field | 3 | 7 | 3 | 15 | Hanford groundwater/sediment system: -3.44 to 0.78 mL/g [12]. Most recent data using ILAW borehole sediment [299-E17-21] yielded K_d values ranging from 3.75 to 10.85 mL/g and had an average of $6.7 \pm 1.9 \text{ mL/g}$ [18]. More recent data for ILAW borehole 299-E24-21 yielded a K_d range from 7.1 to 8.65 for six measurements in Hanford groundwater. [19]. The latter two studies are in excellent agreement. Carr et al. 2003 [21] recommends a range of 0 to 3 and 3 to 10 mL/g for Se for "higher" and "low/trace" concentrations of Se for SAC stochastic predictions. Our range is slightly larger but the best and reasonable conservative values we recommend for the IDF deterministic PA activities fit within the range chosen for trace concentrations of Se. |
| PNNL-13037 Rev. 2 | Table 5.9 | 2004 | -- | -- | Unconfined Far Field Aquifer | 3 | 7 | 3 | 15 | Hanford groundwater/sediment system: -3.44 to 0.78 mL/g [12]. Most recent data using ILAW borehole sediment [299-E17-21] yielded K_d values ranging from 3.75 to 10.85 mL/g and had an average of $6.7 \pm 1.9 \text{ mL/g}$ [18]. More recent data for ILAW borehole 299-E24-21 yielded a K_d range from 7.1 to 8.65 for six measurements in Hanford groundwater. [19]. The latter two studies are in excellent agreement. Carr et al. 2003 [21] recommends a range of 0 to 3 and 3 to 10 mL/g for Se for "higher" and "low/trace" concentrations of Se for SAC stochastic predictions. Our range is slightly larger but the best and reasonable conservative values we recommend for the IDF deterministic PA activities fit within the range chosen for trace concentrations of Se. |
| PNNL-13037 Rev. 2 | Table 6.1 | 2004 | -- | -- | 1998 ILAW PA - All zones | 0 | 0 | -- | -- | |
| PNNL-13037 Rev. 2 | Table 6.1 | 2004 | Glass | -- | 1998 ILAW PA - Near field | 0 | 0 | -- | -- | |
| PNNL-13037 Rev. 2 | Table 6.1 | 2004 | Concrete | -- | 1999 ILAW PA - Near field | 0 | 1 | -- | -- | |
| PNNL-13037 Rev. 2 | Table 6.1 | 2004 | Chemically Impacted | Sand | 2000 ILAW PA | 2 | 4 | -- | -- | |
| PNNL-13037 Rev. 2 | Table 6.1 | 2004 | Non-impacted | Sand | 2001 ILAW PA | 3 | 7 | -- | -- | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 1: Very Acidic | -- | High Impact (1H) | -- | 5 | 3 | 10 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 1: Very Acidic | Sand | Intermediate Impact - Sand (III) | -- | 5 | 3 | 10 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 2: Very High Salt/Very Basic | -- | High Impact (2H) | -- | 0 | 0 | 0.1 | |

Table B-1. Documented Selenium K_d Values

| Document | Location | Year | Waste Chemistry | Size | Impact Zone | Conservative | Best | Min | Max | Comments |
|-------------------|------------|------|--|------|---------------------------------|--------------|------|-----|-----|---|
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 2: Very High Salt/Very Basic | Sand | Intermediate Impact - Sand (21) | --- | 0 | 0 | 1 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 3: Chelates/High Salts | --- | High Impact (3H) | --- | 0 | 0 | 0.1 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 3: Chelates/High Salts | Sand | Intermediate Impact - Sand (31) | --- | 0 | 0 | 1 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 4: Low Organic/Low Salt/Near Neutral | --- | High Impact (4H) | --- | 5 | 3 | 10 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 4: Low Organic/Low Salt/Near Neutral | Sand | Intermediate Impact - Sand (41) | --- | 5 | 3 | 10 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 4: Low Organic/Low Salt/Near Neutral | --- | Groundwater (4G) | --- | 5 | 3 | 10 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 5: IDF Virified Waste | --- | High Impact (5H) | --- | 1 | 0 | 3 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 5: IDF Virified Waste | Sand | Intermediate Impact - Sand (51) | --- | 2 | 0 | 10 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 6: IDF Cementitious Waste | --- | High Impact (6H) | --- | 1 | 0 | 300 | |
| PNNL-14702 Rev. 1 | Table 4.11 | 2006 | Source Category 6: IDF Cementitious Waste | Sand | Intermediate Impact - Sand (61) | --- | 7 | 3 | 15 | |
| PNNL-17154 | Table 3.3 | 2008 | Waste Management Area A-AX | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - ^{79}Se not studied at SST WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.3 | 2008 | Waste Management Area A-AX | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | " " |
| PNNL-17154 | Table 3.4 | 2008 | Natural Pore Waters/Groundwater | Sand | Not Impacted | --- | 5 | 3 | 10 | Appendix A - Adequate Hanford data; Last et al. (2006) |
| PNNL-17154 | Table 3.7 | 2008 | Waste Management Area B-BX-BY | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - ^{79}Se not studied at SST WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.7 | 2008 | Waste Management Area B-BX-BY | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | " " |
| PNNL-17154 | Table 3.9 | 2008 | Waste Management Area C | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - ^{79}Se not studied at SST WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.9 | 2008 | Waste Management Area C | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | " " |

Table B-1. Documented Selenium K_d Values

| Document | Location | Year | Waste Chemistry | Size | Impact Zone | Conservative | Best | Min | Max | Comments |
|-----------------------|--------------------|------|---|------------------------|---------------------|--------------|------|-----|-----------------------|---|
| PNNL-17154 | Table 3.13 | 2008 | Waste Management Area S-SX | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - 79Se not studied at SST WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.13 | 2008 | Waste Management Area S-SX | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | --- |
| PNNL-17154 | Table 3.17 | 2008 | Waste Management Areas T and TX-TY | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - 79Se not studied at T or TX WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.17 | 2008 | Waste Management Areas T and TX-TY | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | --- |
| PNNL-17154 | Table 3.21 | 2008 | Waste Management Area U | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - 79Se not studied at T or TX WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.21 | 2008 | Waste Management Area U | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | --- |
| PNNL-17154 | Table 3.23 | 2008 | BC Cribs | Sand | High Impact | --- | 0 | 0 | 3 | Appendix A - 79Se not studied at SST WMAs; estimated no sorption in high impact zones, but allowed some sorption in intermediate zones based on Kaplan sorption versus pH work on IDF sediments |
| PNNL-17154 | Table 3.23 | 2008 | BC Cribs | Sand | Intermediate Impact | --- | 0.1 | 0 | 3 | --- |
| WCH-515 | Table 25 | 2013 | Low-Organic/Low-Salt/Near-Neutral Waste Chemistry | Sand not Impacted Sand | --- | 5 | 3 | 10 | Reference: PNNL-17154 | |
| RPP-ENV-58782 | Table 6-11 | 2016 | low-salt, near-neutral | < 2mm | --- | --- | 0.1 | --- | --- | Reference: PNNL-17154 |
| RPP-CALC-61032 | Table A-2 | 2017 | Chemically Impacted | Sand | Zone 2a - Far Field | 1 | 2 | --- | --- | Reference: PNNL-13037, Table 5.5 |
| RPP-CALC-61032 | Table A-2 | 2017 | Natural Recharge (no impact from wastes) | Sand | Zone 2b - Far Field | 3 | 7 | --- | --- | Reference: PNNL-13037, Table 5.6 |
| RPP-RPT-59958, Rev. B | Table 4-33 (Draft) | 2017 | Chemically Impacted | Sand | Far Field | 1 | 2 | --- | --- | Reference: PNNL-13037, Table 5.5 |
| RPP-RPT-59958, Rev. B | Table 4-33 (Draft) | 2017 | Natural Recharge (no impact from wastes) | Sand | Far Field | 3 | 7 | 3 | 10 | Reference: PNNL-13037, Table 5.6 |

High Impact or Near Field

Intermediate Impact or Far Field Chemically Impacted

Not Impacted

Not Provided in the Referenced Document

