# NRC RA17-040

# 2016 Annual Radioactive Effluent Release Report

Part 2

Appendix A

Offsite Dose Calculation Manual

**Revision 7** 

ODCM Change Summary Matrix

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Item #	(Old) Rev. 6 page #	(New) Rev. 7 page #	Determination identifier	Description of change

Editorial and grammatical – Determination EG

Changes to the REMP sampling program - Determination REMP

Enhancements due to removal of Plant trip vulnerabilities related to offgas valve 2N62-F057 – Determination OG Changes to the minimum analysis frequency for the Containment Vent and Purge System – Determination PURGE

1	All pages	All pages	EG	Added the ODCM revision effective date (Plant Manager approval date) to the header on all pages for revision 7.	
2	Title Page	Title Page	EG	Updated the ODCM effective date for revision 7.	
3	Table of Contents (TOC)	TOC	EG	Updated reference page numbering for the various sections, subsections, etc. throughout the TOC as necessary.	
4	TOC, page 2	TOC, page 2	EG	Added a dot leader to section 12.5.1 to be consistent with the balance of the TOC.	
5	TOC, page 6	TOC, page 6	EG	Updated section 5.1 with subsections 5.1.4 and 5.1.5 that currently exist in the body of the ODCM.	
6	I-1.1-6	I-1.1-6	EG	Updated the ODCM Equation references within the 10CFR50, Appendix I Regulations (components 1.a. and 2.) and the Technical Specifications Regulations (component 1. and 2.) sections to reference the appropriate ODCM Equations.	
7	1-12.4.1-2	I-12.4.1-2	PURGE	Added superscript "k" in "SAMPLING FREQUENCY" column of row "A. CONTAINMENT VENT AND PURGE SYSTEM" to reference new note "k" on page I- 12.4.1-6 regarding detailed guidance on performing the appropriate sampling, analyses, and purge calculations.	
8	I-12.4.1-2	1-12.4.1-2	PURGE	Split the "MINIMUM ANALYSIS FREQUENCY" column for row "A. CONTAINMENT VENT AND PURGE SYSTEM" into two (2) parts – one (1) part for principal gamma emitters (noble gas samples) and one (1) part for tritium (H-3). The analysis frequency for principal gamma emitters remains unchanged – "Prior to each release Each Purge". However, the minimum analysis frequency for tritium was changed from "Prior to each release Each Purge" to "31 days" to be consistent with NUREG 1302. Regulatory Assurance has verified that there are no regulatory commitments that would be impacted by this change.	
				Since tritium analyses add approximately 3-4 hours to the completion of purge calculations, this change will significantly reduce the time needed to complete purge calculations and will expedite a drywell entry, if required. The time required to complete purge calculations can be reduced from a total of 4 – 6 hours to approximately 1 hour. Drywell tritium results obtained during the previous 31 days will be used to initially perform the purge calculations. Following the completion of the	

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				actual ODCM required tritium purge sampling and analyses, the purge calculations will be updated. New note "k" on page I-12.4.1-6 contains detailed guidance on performing the appropriate sampling, analyses, and purge calculations.
9	I-12.4.1-6	I-12.4.1-6	PURGE	Updated the wording in the last paragraph of Note h to be consistent with the change in the "MINIMUM ANALYSIS FREQUENCY" column in row "A. CONTAINMENT VENT AND PURGE SYSTEM" of Table R12.4.1-1. The original wording stated "both noble gas and tritium analyses must be completed" The new wording states "both noble gas and tritium sampling along with the appropriate purge calculations must be completed" The fact that the appropriate analyses are required is now understood based on the wording of the sentence. That is, the appropriate analyses must be completed in order to perform the purge calculations.
10	I-12.4.1-6	I-12.4.1-6	PURGE	Added notes "j" and "k". Note "j" states: "Not used." This was done to avoid future confusion between "j" and "i" when viewing the ODCM where poor document resolution may exist. Note "k" adds detailed guidance to perform the appropriate sampling, analyses, and purge calculations.
11	I-12.5.1-6 to 1-12.5.1-10	1-12.5.1-6 to 1-12.5.1-11	EG	Updated the page numbering on all pages in Table R12.5.1-1 from "(Page X of 5)" to "(Page X of 6)" due to new page 5 being added to the Table (Page I-12.5.1-10).
12	I-12.5.1-6	I-12.5.1-6	REMP	In the "SAMPLING AND COLLECTION FREQUENCY" column for the airborne radioiodine and particulates exposure pathway (ODCM Table R12.5.1-1, Section 1.), updated the particulate sample collection frequency from "once per 7 days" to "weekly" to be consistent with the current ODCM Table 6-1 and NUREG-1302. Also, updated radioiodine canister collection frequency from "once per 14 days." to "weekly" to be consistent with NUREG-1302 Table 3.12-1 as identified by the NRC (Reference ATI 2541452-02).
13	I-12.5.1-6	I-12.5.1-6	REMP	In the "TYPE AND FREQUENCY OF ANALYSIS" column for the airborne radioiodine and particulates exposure pathway (ODCM Table R12.5.1-1, Section 1.), updated the radioiodine canister analysis frequency from "once per 14 days" to "weekly" to be consistent with NUREG-1302 Table 3.12-1 as identified by the NRC (Reference IR 2541452-02). Updated the particulate gross beta analysis frequency from "once per seven day" to "weekly" Also, updated the particulate gamma isotopic analysis frequency fromonce per 92 days" to "quarterly" The particulate frequency changes were done to be to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
14	I-12.5.1-7	-12.5.1-7	REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the direct radiation exposure pathway (ODCM Table R12.5.1-1, Section 2.) was updated from "92 days" to "Quarterly" to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.

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-	New) Rev. Determination

pathway (ODCM Table R12.5.1-1, Section 2.) was updated from to "quarterly." to be consistent with the current ODCM Table 6- Table 3.12-1.           16         I-12.5.1-8         I-12.5.1-8         REMP         The "SAMPLING AND COLLECTION FREQUENCY" for the water water exposure pathway (ODCM Table R12.5.1-1, Section 3.a.) days" to "Quarterly" to be consistent with the current ODCM Table 1302 Table 3.12-1.           17         I-12.5.1-8         I-12.5.1-8         REMP         The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn exposure pathway (ODCM Table R12.5.1-1, Section 3.a.) was up per 92 days." to "quarterly." to be consistent with the current O NUREG-1302 Table 3.12-1.           18         I-12.5.1-8         I-12.5.1-8         REMP         The "SAMPLING AND COLLECTION FREQUENCY" for the waterborn exposure pathway (ODCM Table R12.5.1-1, Section 3.t.) up per 92 days." to "quarterly." to be consistent with the current O NUREG-1302 Table 3.12-1.           18         I-12.5.1-8         I-12.5.1-8         REMP         The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.t. gamma isotopic analyses was updated from "once per 92 day The charges were done to be to be consistent with the 6-1 and NUREG-1302 Table 3.12-1.           20         I-12.5.1-8         I-12.5.1-8         EG, REMP         The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.d. "once per 7 days." to "weekly." to be consistent with the current NUREG-1302 Table 3.12-1.           20         I-12.5.1-8         I-12.5.1-8		•			
water exposure pathway (ODCM Table R12.5.1-1, Section 3.a.) 1         days" to "Quarterly" to be consistent with the current ODCM Table 31302 Table 3.12-1.         17       I-12.5.1-8       I-12.5.1-8         18       I-12.5.1-8       I-12.5.1-8         19       I-12.5.1-8       REMP         19       I-12.5.1-8       REMP         19       I-12.5.1-8       I-12.5.1-8         19       I-12.5.1-8       I-12.5.1-8         19       I-12.5.1-8       I-12.5.1-8         19       I-12.5.1-8       REMP       The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.0. and years was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 31 day. Also, the					
exposure pathway (ODCM Table R12.5.1-1, Section 3.a.) was up per 92 days." to "quarterly." to be consistent with the current O NUREG-1302 Table 3.12-1.         18       I-12.5.1-8       I-12.5.1-8       REMP       The "SAMPLING AND COLLECTION FREQUENCY" for the wate (3.b.) indicator exposure pathway (ODCM Table R12.5.1-1, Secti from "once per 7 days." to "weekly." to be consistent with the 6-1 and NUREG-1302 Table 3.12-1."         19       I-12.5.1-8       I-12.5.1-8       REMP       The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.1 gamma isotopic analyses was updated from "once per 92 day The changes were done to be to be consistent with the current O NUREG-1302 Table 3.12-1."         20       I-12.5.1-8       I-12.5.1-8       EG, REMP       Added reference to note "(7)" after "Indicator" in Table R12.5.1-1 the "NUMBER OF REPRESENTATIVE SAMPLES AND SAMPLI modified verbiage following "Indicator" in Section 3.c. a. to clarify where the indicator sample is obtained.         21       I-12.5.1-8       I-12.5.1-8       REMP       The "SAMPLING AND COLLECTION FREQUENCY" for the wate indicator sample is obtained.         22       I-12.5.1-8       I-12.5.1-8       REMP       The "SAMPLING AND COLLECTION FREQUENCY" for the wate indicator sample is obtained.         21       I-12.5.1-8       I-12.5.1-8       REMP       The "SAMPLING AND COLLECTION FREQUENCY" for the wate indicator sample is obtained.         22       I-12.5.1-8       I-12.5.1-8       REMP       The "SAMPLING AND COLLE	16	I-12.5.1-8	1-12.5.1-8	REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the waterborne ground/well water exposure pathway (ODCM Table R12.5.1-1, Section 3.a.) was updated from "92 days" to "Quarterly" to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
(3.b.) indicator exposure pathway (ODCM Table R12.5.1-1, Secti from "once per 7 days." to "weekly." to be consistent with the 6-1 and NUREG-1302 Table 3.12-1."19I-12.5.1-8I-12.5.1-8REMPThe "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.t gamma isotopic analyses was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 92 day The changes were done to be to be consistent with the current O NUREG-1302 Table 3.12-1.20I-12.5.1-8I-12.5.1-8EG, REMPAdded reference to note "(7)" after "Indicator" in Table R12.5.1-1 the "NUMBER OF REPRESENTATIVE SAMPLES AND SAMPLI modified verbiage following "Indicator" in Section 3.c.a. to clarify where the indicator sample is obtained.21I-12.5.1-8I-12.5.1-8REMPThe "SAMPLING AND COLLECTION FREQUENCY" for the water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a to clarify where the indicator sample is obtained.22I-12.5.1-8I-12.5.1-8REMPThe "SAMPLING AND COLLECTION FREQUENCY" for the water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a to caper 7 days." to "weekly." to be consistent with the currer and NUREG-1302 Table 3.12-1.22I-12.5.1-8I-12.5.1-8REMPThe "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. and gamma isotopic analyses was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, th	17	l-12.5.1-8	I-12.5.1-8	REMP	The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborne ground/well exposure pathway (ODCM Table R12.5.1-1, Section 3.a.) was updated from "once per 92 days." to "quarterly." to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.1 gamma isotopic analyses was updated from "once per 31 day. Also, the tritium analysis was updated from "once per 92 day The changes were done to be to be consistent with the current O NUREG-1302 Table 3.12-1.20I-12.5.1-8I-12.5.1-8EG, REMPAdded reference to note "(7)" after "Indicator" in Table R12.5.1-1 the "NUMBER OF REPRESENTATIVE SAMPLES AND SAMPLI modified verbiage following "Indicator" in Section 3.c.a. to clarify where the indicator sample is obtained.21I-12.5.1-8I-12.5.1-8REMPThe "SAMPLING AND COLLECTION FREQUENCY" for the water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.0 "once per 7 days." to "weekly." to be consistent with the curr and NUREG-1302 Table 3.12-1.22I-12.5.1-8I-12.5.1-8REMPThe "SAMPLING AND COLLECTION FREQUENCY" for the water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.0 "once per 7 days." to "weekly." to be consistent with the curr and NUREG-1302 Table 3.12-1.22I-12.5.1-8I-12.5.1-8REMPThe "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.0 and gamma isotopic analyses was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once mer 31 monthly" Also, the tritium analysis was updated from "	18	I-12.5.1-8	I-12.5.1-8	REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the waterborne drinking water (3.b.) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.b.) was updated from "once per 7 days." to "weekly." to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1."
21       I-12.5.1-8       I-12.5.1-8       REMP       The "SAMPLING AND COLLECTION FREQUENCY" for the water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c. a) indicator exposure pathway (Section 2.c. a) indicator exposure pathway (Section 3.c. a) i	19	l-12.5.1-8	1-12.5.1-8	REMP	The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborne drinking water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.b.) for gross beta and gamma isotopic analyses was updated from "once per 31 day" tomonthly" Also, the tritium analysis was updated from "once per 92 day" to "quarterly" The changes were done to be to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
22       I-12.5.1-8       I-12.5.1-8       REMP       The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborn indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.0 and gamma isotopic analyses was updated from "once per 31monthly" Also, the tritium analysis was updated from "once per 31	20	I-12.5.1-8	1-12.5.1-8	EG, REMP	Added reference to note "(7)" after "Indicator" in Table R12.5.1-1, Section 3.c.a. under the "NUMBER OF REPRESENTATIVE SAMPLES AND SAMPLE LOCATIONS" and modified verbiage following "Indicator" in Section 3.c.a. to clarify the general area where the indicator sample is obtained.
indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.0 and gamma isotopic analyses was updated from "once per 31 monthly" Also, the tritium analysis was updated from "once	21	I-12.5.1-8	I-12.5.1-8	REMP	
Table 6-1 and NUREG-1302 Table 3.12-1.	22	-12.5.1-8	I-12.5.1-8	REMP	The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborne surface water indicator exposure pathway (ODCM Table R12.5.1-1, Section 3.c.a.) for gross beta and gamma isotopic analyses was updated from "once per 31 day" to monthly" Also, the tritium analysis was updated from "once per 92 day" to "quarterly" The changes were done to be to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.

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23	I-12.5.1-8	I-12.5.1-8	EG, REMP	Removed line separating row 3.c. from 3.d. in ODCM Table R12.5.1-1 and combined row 3.d. with row 3.c. Removed "d. Control Sample" from the "EXPOSURE PATHWAY AND / OR SAMPLE" column, as the "Control Sample" is part of surface water exposure pathway (ODCM Table R12.5.1-1, Section 3.c.). Changed "a. Control" to "b. Control" to maintain sample location order, added reference to note "(7)" after "Control", and clarified the general location of the indicator sample in Table R12.5.1-1, Section 3.d.a. under the "NUMBER OF REPRESENTATIVE SAMPLES AND SAMPLE LOCATIONS". Removed duplicate information in the "SAMPLING AND COLLECTION FREQUENCY" and "TYPE AND FREQUENCY OF ANALYSIS" columns.
24	I-12.5.1-8	I-12.5.1-8	EG, REMP	Changed "e. Sediment" to "d. Sediment" in the "EXPOSURE PATHWAY AND / OR SAMPLE" column to maintain exposure pathway sample order (ODCM Table R12.5.1-1, Section 3.) following removal of "d. Control Sample".
25	I-12.5.1-8	I-12.5.1-8	EG, REMP	Clarified the general location of the indicator sample for waterborne sediment in Table R12.5.1-1, Section 3. under the "NUMBER OF REPRESENTATIVE SAMPLES AND SAMPLE LOCATIONS".
26	I-12.5.1-8	1-12.5.1-8	REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the waterborne sediment exposure pathway (ODCM Table R12.5.1-1, Section 3.) was updated from "184 days" to "Semiannually" to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
27	I-12.5.1-8	1-12.5.1-8	REMP	The "TYPE AND FREQUENCY OF ANALYSIS" for the waterborne sediment exposure pathway (ODCM Table R12.5.1-1, Section 3.) was updated from "once per 184 days." to "semiannually." to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
28	I-12.5.1-9	I-12.5.1-9	EG, REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the ingestion milk exposure pathway (ODCM Table R12.5.1-1, Section 4.) was updated from "Once per 14 days" to "Biweekly" and "once per 31 days" tomonthly" to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
29	I-12.5.1-9	I-12.5.1-9	EG, REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the ingestion fish exposure pathway (ODCM Table R12.5.1-1, Section 4.) was updated from "Twice per 12 months." to "Two times annually." to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
30	I-12.5.1-9	I-12.5.1-9	EG, REMP	The "SAMPLING AND COLLECTION FREQUENCY" for the ingestion food products exposure pathway (ODCM Table R12.5.1-1, Section 4.) was updated from "12 months" to "Annually" to be consistent with the current ODCM Table 6-1 and NUREG-1302 Table 3.12-1.
31	I-12.5.1-9	I-12.5.1-9	EG, REMP	Added "continued" to bottom of page as a new page containing the ingestion vegetation exposure pathway was added to ODCM Table R12.5.1-1, Section 4.

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32	N/A	I-12.5.1-10	REMP	Inserted new section to ODCM Table R12.5.1-1, Section 4. The new ingestion exposure pathway is identified as "d. Vegetation" and was inserted to be consistent with NUREG-1302 Table 3.12-1 as identified by the NRC (Reference IR 2541452-02).	
33	I-12.5.1-10	I-12.5.1-11	REMP	Updated note (1) by adding "(both food products and vegetation exposure pathways)" to include newly added vegetation exposure pathway.	
34	I-12.5.1-10	I-12.5.1-11	REMP	Enhanced note (8) by adding "See the vegetation exposure pathway for additional sampling requirements." to give more robust guidance in the absence of milk sampling.	
35	N/A	II.1-16	OG	Added references #110 and #111 per ATI 2459034-01.	
36	II.1-20	II.1-20	EG	Updated title of table from "Table 1 – (Cont'd)" to "Table 1 – 4 (Cont'd)".	
37	11.2-9	11.2-9	EG, OG	Clarified the second paragraph under section 2.5.4. Reference 2459034-01.	
38	II.2-10	II.2-10	OG	Added discussion to section 2.5.4 detailing the changes to U2 Offgas valve 2N62-F057 based on EC #387237. Reference ATI 2459034-01.	
39	II.2-12	II.2-13	EG	Corrected the editorial errors identified in equation 2-5: " $F_i$ " was corrected to " $f_i$ ". and " $S_i$ " was corrected to " $\overline{S}_i$ " The two (2) editorial errors occurred when creating CY-LA- 170-301 (ODCM), Revision 0 in Aug 2005 from Revision 8 (that includes Revision 7 of Chapter 10 dated September 2003) of the LaSalle Annex. The " $\leq$ " was corrected to "<". This error occurred when creating Revision 1.9 of the LaSalle Annex (Chapter 10) in March 1998 from Revision 1.8 (November 1996). Reference IR 2544411 and 2591990.	
40	II.2-12	II.2-13	EG	Corrected the editorial error in equation 2-6. " $\overline{L}_i$ " referred to the beta skin dose factor in the generic ODCM prior to revision 4 (See Table D-17). Following that revision, the beta skin dose factor was referred to as " $L_i$ " (See ODCM Table 4-13). Reference IR 2544411 and 2591990.	
41	II.2-12	11.2-13	EG	Added information to more robustly define $\mathbf{f}_i$ .	
42	II.2-12	II.2-13 and II.2-14	EG	Updated the "fancy cross" reference symbol in the variable definition section immediately following the equations to match the symbol used in equation 2-6 for clarity. Added definitions for additional variables listed in equations 2-5 and 2-6 that were not defined. Removed paragraph immediately following the variables section as it no longer applies. Updated second paragraph following the variables with the correct reference information.	
43	11.4-2	11.4-2	EG	Added equation number "(4-1)".	
44	11.4-3	11.4-3	EG	Added equation number "(4-2)".	
45	11.4-4	11.4-4	EG	Added equation number "(4-3)".	
46	11.4-5	11.4-5	EG	Added equation number "(4-4)".	
47	11.4-7	11.4-7	EG	Added equation number "(4-5)".	
48	11.4-8	11.4-8	EG	Added equation number "(4-6)".	
49	11.4-9	11.4-9	EG	Added equation number "(4-7)".	
50	II.4-10	11.4-10	EG	Added equation number "(4-8)".	
51	11.4-11	11.4-11	EG	Added equation number "(4-9)".	

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52	11.4-12	11.4-12	EG	Added equation number "(4-10)".	
53	II.4-12 and II.4-13	11.4-13	EG	Added equation number "(4-11a)", "(4-11b)", and "(4-11c)".	
54	11.4-14	11.4-14	EG	Added equation number "(4-12a)", "(4-12b)", and "(4-12c)".	
55	II.4-16	11.4-16	EG	Added equation number "(4-13a)", "(4-13b)", and "(4-13c)".	
56	II.6-2 to II.6-8	II.6-2 to II.6-9	EG, REMP	Updated the page numbering on Table 6-1 from "(Page X of 7)" to "(Page X of 8)" due to new page 7 being added to the Table (Page II.6-8)	
57	11.6-2	11.6-2	REMP	In the "Sampling or Collection Frequency" column for the airborne radioiodine and particulates exposure pathway (ODCM Table 6-1, Section 1), updated the radioiodine canister collection frequency from "biweekly." to "weekly." to be consistent with NUREG-1302 Table 3.12-1 as identified by the NRC (Reference ATI 2541452-02).	
58	II.6-2	11.6-2	REMP	In the "Type and Frequency of Analysis" column for the airborne radioiodine and particulates exposure pathway (ODCM Table 6-1, Section 1.), updated the radioiodine canister analysis frequency from "biweekly" to "weekly" to be consistent with NUREG-1302 Table 3.12-1 as identified by the NRC (Reference ATI 2541452-02).	
59	11.6-2	11.6-2	REMP	Installed a new air monitoring location identified as L-11A to replace the existing L-11 air monitoring location. L-11A is located 165 feet west of L-11. The new air monitoring location has no obstructions at this time and will replace L-11 following the approval of this ODCM revision. L-11 has significant obstructions and is required to be replaced per CY-AA-170-1000, Radiological Environmental Monitoring Program and Meteorological Program Implementation. The coordinates for L-11A are N 41° 09.405', W 88° 39.565'. Location L-11 has been replaced with L-11A in the "Sample or Monitoring Location" column under "2. Indicators – Far Field" for the airborne radioiodine and particulates exposure pathway (ODCM Table 6-1, Section 1.). Reference ATI 1494410-10 and 2430660-02.	
60	11.6-5	11.6-5	EG	Page II.6-5 is a continuation of the previous page. Therefore, the "Sampling or Collection Frequency" and "Type and Frequency of Analysis" columns were enhanced by including information from the same columns on the previous page.	
61	11.6-7	11.6-7	EG	Added metric equivalent distances to the indicator and control sample locations for the ingestion food product exposure pathway (ODCM Table 6-1, Section 4).	
62	N/A	11.6-8	REMP	Inserted new section to ODCM Table 6-1, Section 4. The new ingestion exposure pathway is identified as "d. Vegetation" and was inserted to be consistent with NUREG-1302 Table 3.12-1 as identified by the NRC (Reference ATI 2541452-02).	
63	11.6-9	II.6-10	REMP	Updated air monitoring location identifier at the Ransom location from L-11 to L-11A.	

Appendix B

Offsite Dose Calculation Manual

Part I RECS

Part II ODCM

LaSalle Station Units 1 and 2

**Revision 7** 

# **OFFSITE DOSE CALCULATION MANUAL**

# **PART I RECS**

# **PART II ODCM**

LaSalle Station Units 1 and 2

Effective: January 2016

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# RADIOLOGICAL EFFLUENT CONTROLS

# LASALLE STATION Units 1 and 2

### 1.0 USE AND APPLICATION

### 1.1 DEFINITIONS

-----NOTE-----

The defined terms of this section appear in capitalized type and are applicable throughout these Offsite Dose Calculation Manual (ODCM) Controls and Bases.

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Term	Definition
ACTION	ACTION shall be that part of a control that prescribes remedial measures required under designated conditions.
CHANNEL CALIBRATION	A CHANNEL CALIBRATION shall be the adjustment, as necessary, of the channel output such that it responds within the necessary range and accuracy to known values of the parameter that the channel monitors. The CHANNEL CALIBRATION shall encompass the entire channel, including the required sensor, alarm, display, and trip functions, and shall include the CHANNEL FUNCTIONAL TEST. Calibration of instrument channels with resistance temperature detector (RTD) or thermocouple sensors may consist of an in-place qualitative assessment of sensor behavior and normal calibration of the remaining adjustable devices in the channel. The CHANNEL CALIBRATION may be performed by means of any series of sequential, overlapping, or total channel steps so that the entire channel is calibrated.
	For specific calibration requirements refer to surveillance requirements section for the applicable instrumentation.
CHANNEL CHECK	A CHANNEL CHECK shall be a qualitative assessment, by observation, of channel behavior during operation. This determination shall include, where possible, comparison of the channel indication and status to other indications or status derived from independent instrument channels measuring the same parameter.
CHANNEL FUNCTIONAL TEST	A CHANNEL FUNCTIONAL TEST shall be the injection of a simulated or actual signal into the channel as close to the sensor as practicable to verify OPERABILITY, including required alarm, interlock, display, and trip functions, and channel failure trips. The CHANNEL FUNCTIONAL TEST may be performed by means of any series of sequential, overlapping, or total channel steps so that the entire channel is tested.

## 1.1 DEFINITIONS (continued)

CONTINUOUS SAMPLING	Uninterrupted sampling with the exception of sampling interruptions of short duration for required surveillances.
DOSE EQUIVALENT I-131	That concentration of I-131 (microcuries/gram) that alone would produce the same thyroid dose as the quantity and isotopic mixture of I-131, I-132, I-133, I-134, and I-135 actually present. The thyroid dose conversion factors used for this calculation shall be those listed in Table III of TID -14844, AEC, 1962, "Calculation of Distance Factors for Power and Test Reactor Sites"; Table E-7 of Regulatory Guide 1.109, Rev. 1, NRC, 1977; or ICRP 30, Supplement to Part 1, pages 192-212, Table titled, "Committed Dose Equivalent in Target Organs or Tissues per Intake of Unit Activity."
GASEOUS RADWASTE TREATMENT SYSTEM	Any system designed and installed to reduce radioactive gaseous effluents by collecting primary coolant system offgases from the primary system and providing for delay or holdup for the purpose of reducing the total radioactivity prior to release to the environment.
MEMBERS OF THE PUBLIC	Any individual, except when that individual is receiving an occupational dose.
MODE	A MODE shall correspond to any one inclusive combination of mode switch position, average reactor coolant temperature, and reactor vessel head closure bolt tensioning specified in Technical Specifications with fuel in the reactor vessel.
OCCUPATIONAL DOSE	The dose received by an individual in the course of employment in which the individual's assigned duties involve exposure to radiation and/or to radioactive material from licensed and unlicensed sources of radiation, whether in the possession of the licensee or other person. Occupational dose does not include dose from background radiation, as a patient from medical practices, from voluntary participation in medical research programs, or as a member of the public.

# 1.1 DEFINITIONS (continued)

OFFSITE DOSE CALCULATION MANUAL (ODCM)	The ODCM shall contain the methodology and parameters used in the calculation of offsite doses resulting from radioactive gaseous and liquid effluents, in the calculation of gaseous and liquid effluent monitoring Alarm/Trip Setpoints, and in the conduct of the Radiological Environmental Monitoring Program. The ODCM shall also contain (1) the Radioactive Effluent Controls and Radiological Environmental Monitoring Program and (2) descriptions of the information that should be included in the Annual Radiological Environmental Operating and Radioactive Effluent Release Reports.
OPERABLE - OPERABILITY	A system, subsystem, division, component, or device shall be OPERABLE or have OPERABILITY when it is capable of performing its specified function(s) and when all necessary attendant instrumentation, controls, normal or emergency electrical power, cooling and seal water, lubrication, and other auxiliary equipment that are required for the system, subsystem, division, component, or device to perform its specified function(s) are also capable of performing their related support function(s).
POSITION INDICATION VERIFICATION	POSITION INDICATION VERIFICATION shall be the comparison of the physical position of the Blowdown Flow Control Valve (0WL005) actuator shaft in percent open to the remote (0ZI- WL002A) Blowdown Flow Control Valve position indication in percent open.
PROCESS CONTROL PROGRAM (PCP)	The PCP shall contain the current formulas, sampling, analyses, test, and determinations to be made to ensure that processing and packaging of solid radioactive wastes based on demonstrated processing of actual or simulated wet solid wastes shall be accomplished in such a way as to assure compliance with 10 CFR Parts 20, 61, and 71, State regulations, burial ground requirements, and other requirements governing the disposal of solid radioactive waste.
PURGE – PURGING	PURGE or PURGING shall be the controlled process of discharging air or gas from a confinement to maintain temperature, pressure, humidity, concentration or other operating condition, in such a manner that replacement air or gas is required to purify the confinement.
RATED THERMAL POWER (RTP)	The applicable unit's RTP shall be a total reactor core heat transfer rate to the reactor coolant as defined in Technical Specifications.

## 1.1 DEFINITIONS (continued)

RADIOLOGICAL EFFLUENT CONTROL STANDARDS (RECS)	A compilation of the various regulatory requirements, surveillance and bases, commitments and/or components of the radiological effluent and environmental monitoring programs for LaSalle Station. To assist in the understanding of the relationship between effluent regulations, ODCM equations, RECS and related Technical Specification requirements, Table 1-1 provides a matrix that relates these various components, as well as the Radiological Environmental Monitoring Program fundamental requirements.
SITE BOUNDARY	That line beyond which the land is not owned, leased, or otherwise controlled by licensee as defined in ODCM Part II Figure 1-3.
SOLIDIFICATION	SOLIDIFICATION shall be the conversion of radioactive wastes from liquid systems to a homogeneous (uniformly distributed), monolithic, immobilized solid with definite volume and shape, bounded by a stable surface of distinct outline on all sides (free- standing).
SOURCE CHECK	A SOURCE CHECK shall be the qualitative assessment of channel response when the channel sensor is exposed to a radioactive source (This could be an external source or known radioactive process stream).
THERMAL POWER	THERMAL POWER shall be the total reactor core heat transfer rate to the reactor coolant.
UNRESTRICTED AREA	UNRESTRICTED AREA means an area, access to which is neither limited nor controlled by the licensee.
VENTILATION EXHAUST TREAT- MENT SYSTEM	A VENTILATION EXHAUST TREATMENT SYSTEM shall be any system designed and installed to reduce gaseous radioiodine or radioactive material in particulate form in effluents by passing ventilation or vent exhaust gases through charcoal adsorbers and/or HEPA filters for the purpose of removing iodines or particulates from the gaseous exhaust system prior to the release to the environment (such a system is not considered to have any effect on noble gas effluents). Engineered Safety Feature (ESF) atmospheric cleanup systems are not considered to be VENTILATION EXHAUST TREATMENT SYSTEM components.

VENTING VENTING shall be the controlled process of discharging air or gas from a confinement to maintain temperature, pressure, humidity, concentration or other operating condition, in such a manner that replacement air or gas is not provided or required during VENTING. Vent, used in system names, does not imply a VENTING process.

DEFINITIONS PECULIAR TO ESTIMATING DOSE TO MEMBERS OF THE PUBLIC USING THE ODCM COMPUTER PROGRAM:

- a. ACTUAL Refers to using known release data to project the dose to the public for the previous time period. These data are stored in the database and used to demonstrate compliance with the reporting requirements of RECS.
- b. PROJECTED Refers to using known release data from the previous time period or estimated release data to forecast a future dose to the public. This data is <u>NOT</u> incorporated into the database.

## Table 1-1 (Page 1 of 2)

## COMPLIANCE MATRIX

Regulation		Dose Component Limit	ODCM Equation	RECS	Technical Specification
10 CFR 50 Appendix I	1.	Gamma air dose and beta air dose due to airborne radioactivity in effluent plume.	4-4 4-5	12.4.2	5.5.4.h
		a. Whole body and skin dose due to airborne radioactivity in effluent plume are reported only if certain gamma and beta air dose criteria are exceeded.	4-6 4-7	N/A	N/A
		b Projected doses due to gaseous release, when averaged over 31 days, exceed 0.3 mrem to any organ.	N/A	12.4.5	5.5.4.f
		c Projected doses due to liquid release, when averaged over 31 days, exceed 0.06 mrem to the total body or 0.2 mrem to any organ.	N/A	12.3.3	5.5.4.f
	2.	CDE for all organs and all four age groups due to iodines and particulates in effluent plume. All pathways are considered.	4-8	12.4.3	5.5.4.i
	3.	CDE for all organs and all four age groups due to radioactivity in liquid effluents.	3-3	12.3.2	5.5.4.d
10 CFR 20	1.	TEDE, totaling all deep dose equivalent components (direct, ground and plume shine) and CDE (all pathways, both airborne and liquid-borne). CDE evaluation is made for adult only using FGR 11 database.	5-3	12.4.9	5.5.4.c
40 CFR 190 (now by	1.	Whole body dose (DDE) due to direct dose, ground and plume shine from all sources at a station.	5-2	12.4.7	5.5.4.j
reference, also part of 10 CFR 20)	2.	Organ doses (CDE) to an adult due to all pathways.	3-3 4-8		
Technical Specifications	1.	"Instantaneous" whole body (DDE), skin (SDE), and thyroid (CDE) dose rates due to radioactivity in airborne effluents. For the thyroid dose, only inhalation is considered.	4-1 4-2 4-3	12.4.1	5.5.4.g
	2.	"Instantaneous" concentration limits for liquid effluents.	3-1	12.3.1	5.5.4.b
	3.	Radioactive Effluent Release Report	N/A	12.6.2	5.6.3

## Table 1-1 (Page 2 of 2)

### COMPLIANCE MATRIX

Regulation		Dose Component Limit	ODCM Equation	RECS	Technical Specification
10CFR50 Appendix I Section IV.B.2	1.	Implement environmental monitoring program.	N/A	12.5.1	N/A
10CFR50 Appendix I Section IV.B.3	1.	Land Use Census	N/A	12.5.2	N/A
10CFR50 Appendix I Section IV.B.2	1.	Interlaboratory Comparison Program	N/A	12.5.3	N/A
10CFR50 Appendix I Section IV.B.2 and Technical Specifications	1.	Annual Radiological Environmental Operating Report	N/A	12.6.1	5.6.2

### 1.0 USE AND APPLICATION

### 1.2 Logical Connectors

PURPOSE	The purpose of this section is to explain the meaning of logical connectors. Logical connectors are used in ODCM to discriminate between, and yet connect, discrete Conditions, Required Actions, Completion Times, Surveillances, and Frequencies. The only logical connectors that appear in ODCM are <u>AND</u> and <u>OR</u> . The physical arrangement of these connectors constitutes logical conventions with specific meanings.
BACKGROUND	Several levels of logic may be used to state Required Actions. These levels are identified by the placement (or nesting) of the logical connectors and by the number assigned to each Required Action. The first level of logic is identified by the first digit of the number assigned to a Required Action and the placement of the logical connector in the first level of nesting (i.e., left justified with the number of the Required Action). The successive levels of logic are identified by additional digits of the Required Action number and by successive indentations of the logical connectors. When logical connectors are used to state a Condition, Completion Time, Surveillance, or Frequency, only the first level of logic is used, and the logical connector is left justified with the statement of the Condition, Completion Time, Surveillance, or Frequency.
EXAMPLES	The following examples illustrate the use of logical connectors.
	(continued)

# EXAMPLES (continued) EXAMPLE 1.2-1

ACTIONS

CON	IDITION	REQUIRED ACTION	COMPLETION
A. Control not met.		A.1 Verify	
		AND	
		A.2 Restore	

In this example, the logical connector <u>AND</u> is used to indicate that, when in Condition A, both Required Actions A.1 and A.2 must be completed.

#### EXAMPLES (continued) EXAMPLE 1.2-2

ACTI	ACTIONS						
CONDITION		REQUIRED ACTION		COMPLETION TIME			
A.	Control not met.	A.1	Trip				
		<u>OR</u>					
		A.2.1	Verify				
		AND					
		A.2.2.1	Reduce				
			OR				
		A.2.2.2	Perform				
		OR					
		A.3	Align				

This example represents a more complicated use of logical connectors. Required Actions A.1, A.2 and A.3 are alternate choices, only one of which must be performed as indicated by the use of the logical connector <u>OR</u> and the left justified placement. Any one of these three Action may be chosen. If A.2 is chose, then both A.2.1 and A.2.2 must be performed as indicated by the logical connector <u>AND</u>. Required Action A.2.2 is met by performing A.2.2.1 or A.2.2.2. The indented position of the logical connector <u>OR</u> indicates that A.2.2.1 and A.2.2.2 are alternative choices, only one of which must be performed.

### 1.0 USE AND APPLICATION

### 1.3 Completion Times

PURPOSE	The purpose of this section is to establish the Completion Time convention and to provide guidance for its use.
BACKGROUND	ODCM Radiological Effluent Controls (RECs) specify minimum requirements for ensuring safe operation of the unit. The ACTIONS associated with a REC state Conditions that typically describe the ways in which the requirements of the REC can fail to be met. Specified with each stated Condition are Required Action(s) and Completion Times.
DESCRIPTION	The Completion Time is the amount of time allowed for completing a Required Action. It is referenced to the time of discovery of a situation (e.g., inoperable equipment or variable not within limits) that requires entering an ACTIONS Condition unless otherwise specified, providing the unit is in a MODE or specified condition stated in the Applicability of the REC. Required Actions must be completed prior to the expiration of the specified Completion Time. An ACTIONS Condition remains in effect and the Required Actions apply until the Condition no longer exists or the unit is not within the REC Applicability.
	If situations are discovered that require entry into more than one Condition at a time within a single REC (multiple Conditions), the Required Actions for each Condition must be performed within the associated Completion Time. When in multiple Conditions, separate Completion Times are tracked for each Condition starting from the time of discovery of the situation that required entry into the Condition.
	Once a Condition has been entered, subsequent divisions, subsystem, components or variables expressed in the Condition, discovered to be inoperable or not within limits, will <u>not</u> result in separate entry into the Condition unless specifically stated. The Required Actions of the Condition continue to apply to each additional failure, with Completion Times based on initial entry into the Condition.
	(continued)

DESCRIPTION (continued)	expres limits,	ver, when a <u>subsequent</u> division, subsystem, component, or variable sed in the Condition is discovered to be inoperable or not within the Completion Time(s) may be extended. To apply this Completion extension, two criteria must first be met. The subsequent ability:		
	a.	Must exist concurrent with the first inoperability; and		
	<ul> <li>Must remain inoperable or not within limits after the first inoperability is resolved.</li> </ul>			
		al Completion Time allowed for completing a Required Action to address osequent inoperability shall be limited to the more restrictive of either:		
	a.	The stated Completion Time, as measured from the initial entry into the Condition, plus an additional 24 hours; or		
	b.	The stated Completion Time as measured from discovery of the subsequent inoperability.		
	have e (for ea Condit	pove Completion Time extension does not apply to those RECs that exceptions that allow completely separate re-entry into the Condition ach division, subsystem, component, or variable expressed in the tion) and separate tracking of Completion Times based on this re- These exceptions are stated in individual RECs.		
Time with a modified "time zero." This modified "time expressed as a repetitive time (i.e., "once per 8 hou Completion Time is referenced from a previous com Action versus the time of Condition entry) or as a time phrase "from discovery" Example 1.3-3 illustrated		bove Completion Time extension does not apply to a Completion with a modified "time zero." This modified "time zero" may be seed as a repetitive time (i.e., "once per 8 hours," where the letion Time is referenced from a previous completion of the Required versus the time of Condition entry) or as a time modified by the e "from discovery" Example 1.3-3 illustrates one use of this type npletion Time. The 10 day Completion Time specified for Condition B in Example 1.3-3 may not be extended.		
EXAMPLES	The following examples illustrate the use of Completion Times with different types of Conditions and changing Conditions.			

# EXAMPLES (continued) EXAMPLE 1.3-1

### ACTIONS

CONDITION		I	REQUIRED ACTION	COMPLETION TIME
B. Required Action and associated Completion Time not met.		B.1	Be in MODE 3.	12 hours
		AND		36 hours
		B.2	Be in MODE 4.	

Condition B has two Required Actions. Each Required Action has its own separate Completion Time. Each Completion Time is referenced to the time that Condition B is entered.

The Required Actions of Condition B are in to be in MODE 3 within 12 hours <u>AND</u> in MODE 4 within 36 hours. A total of 12 hours is allowed for reaching MODE 3 and a total of 36 hours (not 48 hours) is allowed for reaching MODE 4 from the time that Condition B was entered. If MODE 3 is reached within 6 hours, the time allowed for reaching MODE 4 is the next 30 hours because the total time allowed for reaching Mode 4 is 36 hours.

If Condition B is entered while in MODE 3, the time allowed for reaching MODE 4 is the next 36 hours.

#### 1.3 Completion Times

#### EXAMPLES (continued) EXAMPLE 1.3-2

ACTIONS

	CONDITION	I	REQUIRED ACTION	COMPLETION TIME
A.	One monitor inoperable.	A.1	Restore monitor to OPERABLE status.	7 days
В.	B. Required Action and associated Completion Time not met.		Be in MODE 3.	12 hours
	nino not mot.	AND		
		B.2	Be in MODE 4.	36 hours

When a monitor is declared inoperable, Condition A is entered. If the monitor is not restored to OPERABLE status within 7 days, Condition B is also entered and the Completion Time clocks for Required Action B.1 and B.2 start. If the inoperable monitor is restored to OPERABLE status after Condition B is entered, Condition A and B are exited, and therefore, the Required Actions of Condition B may be terminated.

When a monitor pump is declared inoperable while the first monitor is still inoperable, Condition A is not re-entered for the second monitor. REC 12.0.3 is entered, since the ACTIONS do not include a Condition from more than one inoperable monitor. The Completion Time clock for Condition A does not stop after REC 12.0.3 is entered, but continues to be tracked from the time Condition A was initially entered.

While in REC 12.0.3, if one of the inoperable monitors is restored to OPERABLE status and the Completion Time for Condition A has not expired, REC 12.0.3 may be exited and operation continued in accordance with Condition A.

### 1.3 Completion Times

### EXAMPLES <u>EXAMPLE 1.3-2</u> (continued)

While in REC 12.0.3, if one of the inoperable monitors is restored to OPERABLE status and the Completion Time for Condition A has expired, REC 12.0.3 may be exited and operation continued in accordance with Condition B. The Completion Time for Condition B is tracked from the time the Condition A Completion Time expired.

On restoring one of the monitors to OPERABLE status, the Condition A Completion Time is not reset, but continues from the time the first monitor was declared inoperable. This Completion Time may be extended if the monitor restored to OPERABLE status was the first inoperable monitor. A 24 hour extension to the stated 7 days is allowed, provided this does not result in the second monitor being inoperable for > 7 days.

#### **EXAMPLES** EXAMPLE 1.3-3

(continued)

## ACTIONS

	CONDITION		REQUIRED ACTION	COMPLETION TIME
Α.	One Function X subsystem inoperable.	A.1	Restore Function X subsystem to OPERABLE status.	7 days <u>AND</u> 10 days from discovery of failure to meet the Control
В.	One Function Y subsystem inoperable.	B.1	Restore Function Y subsystem to OPERABLE status.	72 hours <u>AND</u> 10 days from discovery to meet Control
C.	One Function X subsystem inoperable. <u>AND</u> One Function Y subsystem inoperable.	C.1 <u>OR</u> C.2	Restore Function X subsystem to OPERABLE status. Restore Function Y subsystem to OPERABLE status.	72 hours 72 hours

## EXAMPLES <u>EXAMPLE 1.3-3</u> (continued)

When one Function X subsystem and one Function Y subsystem are inoperable, Condition A and Condition B are concurrently applicable. The Completion Times for Condition A and Condition B are tracked separately for each subsystem, starting from the time each subsystem was declared inoperable and the Condition was entered. A separate Completion Time is established for Condition C and tracked from the time the second subsystem was declared inoperable (i.e., the time the situation described in Condition C was discovered).

If Required Action C.2 is completed within the specified Completion Time, Conditions B and C are exited. If the Completion Time for Required Action A.1 has not expired, operation may continue in accordance with Condition A. The remaining Completion Time in Condition A is measured from the time the affected subsystem was declared inoperable (i.e., initial entry into Condition A).

The Completion Times of Conditions A and B are modified by a logical connector, with a separate 10 day Completion Time measured from the time it was discovered the REC was not met. In this example, without the separate Completion Time, it would be possible to alternate between Conditions A, B, and C in such a manner that operation could continue indefinitely without ever restoring systems to meet the REC. The separate Completion Time modified by the phrase "from discovery of failure to meet the Control" is designed to prevent indefinite continued operation while not meeting the REC. This Completion Time allows for an exception to the normal "time zero" for beginning the Completion Time "clock." In this instance, the Completion Time "time zero" is specified as commencing at the time the associated Condition was entered.

#### EXAMPLES (continued) EXAMPLE 1.3-4

ACTIONS

Construction of the local division of the lo				
	CONDITION	F	REQUIRED ACTION	COMPLETION TIME
A.	One or more required instruments inoperable.	A.1	Restore instruments(s) to OPERABLE status.	4 hours
В.	Required Action and associated Completion	B.1	Be in MODE 3.	12 hours
Time not met.		AND		36 hours
		B.2	Be in MODE 4.	

A single Completion Time is used for any number of instruments inoperable at the same time. The Completion Time associated with Condition A is based on the initial entry into Condition A and is not tracked on a per instrument basis. Declaring subsequent instruments inoperable, while Condition A is still in effect, does not trigger the tracking of separate Completion Times.

Once one of the instruments has been restored to OPERABLE status, the Condition A Completion Time is not reset, but continues from the time the first instrument was declared inoperable. The Completion Time may be extended if the instrument restored to OPERABLE status was the first inoperable instrument. The Condition A Completion Time may be extended for up to 4 hours provided this does not result in any subsequent instrument being inoperable for > 4 hours.

If the Completion Time of 4 hours (plus the extension) expires while one or more instruments are still inoperable, Condition B is entered.

## EXAMPLES <u>EXAMPLE 1.3-5</u> (continued)

ACTIONS

	CONDITION	1	REQUIRED ACTION	COMPLETION TIME
Α.	One or more instruments inoperable.	A.1	Restore instrument(s) to OPERABLE status.	4 hours
В.	Required Action and associated Completion	B.1	Be in MODE 3.	12 hours
	Time not met.	AND		36 hours
		B.2	Be in MODE 4.	

The Note above the ACTIONS Table is a method of modifying how the Completion Time is tracked. If this method of modifying how the Completion Time is tracked was applicable only to a specific Condition, the Note would appear in that Condition rather than at the top of the ACTIONS Table.

The Note allows Condition A to be entered separately for each inoperable instrument, and Completion Times tracked on a per instrument basis. When an instrument is declared inoperable, Condition A is entered and its Completion Time starts. If subsequent instruments are declared inoperable, Condition A is entered for each instrument and separate Completion Times start and are tracked for each instrument.

If the Completion Time associated with an instrument in Condition A expires, Condition B is entered for that instrument. If the Completion Times associated with subsequent instruments in Condition A expire, Condition B is entered separately for each instrument and separate Completion Times start and are tracked for each instrument. If a instrument that caused entry into Condition B is restored to OPERABLE status, Condition B is exited for that instrument.

Since the Note in this example allows multiple Condition entry and tracking of separate Completion Times, Completion Time extensions do not apply.

#### EXAMPLES (continued) EXAMPLE 1.3-6

ACTIONS

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	CONDITION	REQUIRED ACTION	COMPLETION TIME
A. One channel inoperable.		Perform RSR 12.x.x.x.	Once per 8 hours
		OR	-
		Reduce THERMAL POWER to ≤ 50% RTP.	8 hours
В.	Required Action and associated Completion Time not met.	B.1 Be in MODE 3.	12 hours

Entry into Condition A offers a choice between Required Action A.1 or A.2. Required Action A.1 has a "once per" Completion Time, which qualifies for the 25% extension, per RSR 12.0.2 to each performance after the initial performance. The initial 8 hour interval of Required Action A.1 begins when Condition A is entered and the initial performance of Required Action A.1 must be completed within the first 8 hour interval. If Required Action A.1 is followed and the Required Action is not met within the Completion Time (plus the extension allowed by RSR 12.0.2), Condition B is entered. If Required Action A.2 is followed and the Completion Time of 8 hours is not met, Condition B is entered.

If after entry into Condition B, Required Action A.1 or A.2 is met, Condition B is exited and operation may then continue in Condition A.

#### EXAMPLES (continued) EXAMPLE 1.3-7

ACTIONS

0113			
CONDITION	F	REQUIRED ACTION	COMPLETION TIME
One subsystem inoperable.	A.1	Verify affected subsystem isolated.	1 hour <u>AND</u> Once per 8 hours thereafter
	<u>AND</u>		
	A.2	Restore subsystem to OPERABLE status.	72 hours
Required Action and associated Completion	B.1	Be in MODE 3.	12 hours
Time not met.	AND		
	B.2	Be in MODE 4.	36 hours
	CONDITION One subsystem inoperable. Required Action and	CONDITION       F         One subsystem inoperable.       A.1         A.1       A.1         AND A.2       A.2         Required Action and associated Completion Time not met.       B.1         AND A.2	CONDITIONREQUIRED ACTIONOne subsystem inoperable.A.1Verify affected subsystem isolated.A.1Verify affected subsystem isolated.AND A.2AND Restore subsystem to OPERABLE status.Required Action and associated Completion Time not met.B.1B.1Be in MODE 3. AND

Required Action A.1 has two Completion Times. The 1 hour Completion Time begins at the time the Condition is entered and each "Once per 8 hours thereafter" interval begins upon performance of Required Action A.1.

If after Condition A is entered, Required Action A.1 is not met within either the initial 1 hour or any subsequent 8 hour interval from the previous performance (plus the extension allowed by RSR 12.0.2), Condition B is entered. The Completion Time clock for Condition A does not stop after Condition B is entered, but continues from the time Condition A was initially entered. If Required Action A.1 is met after Condition B is entered, Condition B is exited and operation may continue in accordance with Condition A, provided the Completion Time for Required Action A.2 has not expired.

IMMEDIATE<br/>COMPLETIONWhen "Immediately" is used as a Completion Time, the Required Action should be<br/>pursued without delay and in a controlled manner.TIME

## 1.4 Frequency

# PURPOSE The purpose of this section is to define the proper use and application of Frequency requirements.

DESCRIPTION Each ODCM Radiological Effluent Surveillance Requirement (RSR) has a specified Frequency in which the Surveillance must be met in order to meet the associated ODCM REC. An understanding of the correct application of the specified Frequency is necessary for compliance with the RSR.

The "specified Frequency" is referred to throughout this section and each of the Requirements of Section 12.0, ODCM Surveillance Requirement (RSR) Applicability. The "specified Frequency" consists of the requirements of the Frequency column of each RSR, as well as certain Notes in the Surveillance column that modify performance requirements.

Sometimes special situations dictate when the requirements of a Surveillance are to be met. They are "otherwise stated" conditions allowed by RSR 12.0.1. They may be stated as clarifying Notes in the Surveillance, as part of the Surveillance, or both. Example 1.4-4 discusses these special situations.

Situations where a Surveillance could be required (i.e., its Frequency could expire), but where it is not possible or not desired that it be performed until sometime after the associated REC is within its Applicability, represent potential RSR 12.0.4 conflicts. To avoid these conflicts, the RSR (i.e., the Surveillance or the Frequency) is stated such that it is only "required" when it can be and should be performed. With a RSR satisfied, RSR 12.0.4 imposes no restriction.

The use of "met" or "performed" in these instances conveys specified meanings. A Surveillance is "met" only when the acceptance criteria are satisfied. Known failure of the requirements of a Surveillance, even without a Surveillance specifically being "performed," constitutes a Surveillance not "met." "Performance" refers only to the requirement to

## 1.4 Frequency

DESCRIPTION (continued)	specifically determine the ability to meet the acceptance criteria. RSR 12.0.4 restrictions would not apply if both the following conditions are satisfied:		
	a. The Surveillance is not required to be perform	ned; and	
	<ul> <li>The Surveillance is not required to be met or, even if required to be met, is not known to be failed.</li> </ul>		
EXAMPLES	The following examples illustrate the various ways that Frequencies are specified. In these examples, the Applicability of the REC (REC not shown) is MODES 1, 2, and 3.		
	EXAMPLE 1.4-1		
	SURVEILLANCE REQUIREMENTS		
	SURVEILLANCE	FREQUENCY	
Perform CHANNEL CHECK		12 hours	

Example 1.4-1 contains the type of RSR most often encountered in the ODCM. The Frequency specifies an interval (12 hours) during which the associated Surveillance must be performed at least one time. Performance of the Surveillance initiates the subsequent interval. Although the Frequency is stated as 12 hours, an extension of the time interval to 1.25 times the interval specified in the Frequency is allowed by RSR 12.0.2 for operational flexibility. The measurement of this interval continues at all times, event when the RSR is not required to be met per RSR 12.0.1 (such as when the equipment is inoperable, a variable is outside specified limits, or the unit is outside the Applicability of the REC). If the interval specified by RSR 12.0.2 is exceeded while the unit is in a MODE or other specified condition in the Applicability of the REC,

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## 1.0 USE AND APPLICATION

## 1.4 Frequency

## EXAMPLES <u>EXAMPLE 1.4-1</u> (continued)

and the performance of the Surveillance is not otherwise modified (refer to Examples 1.4-3 and 1.4-4), then RSR 12.0.3 becomes applicable.

If the interval as specified by RSR 12.0.2 is exceeded while the unit is not in a MODE or other specified condition in the Applicability of the REC for which performance of the RSR is required, the Surveillance must be performed within the Frequency requirements of RSR 12.0.2 prior to entry into the MODE or other specified condition. Failure to do so would result in a violation of RSR 12.0.4.

## EXAMPLE 1.4-2

SURVEILLANCE REQUIREMENTS

SURVEILLANCE	FREQUENCY
Verify flow is within limits.	Once within 12 hours after <u>&gt;</u> 25% RTP
	AND 24 hours thereafter

Example 1.4-2 has two Frequencies. The first is a one time performance Frequency, and the second is of the type shown in Example 1.4-1. The logical connector "<u>AND</u>" indicates that both Frequency requirements must be met. Each time reactor power is increased from a power level < 25% RTP to  $\geq$  25% RTP, the Surveillance must be performed within 12 hours.

## 1.4 Frequency

## EXAMPLES <u>EXAMPLE 1.4-2</u> (continued)

The use of "once" indicates a single performance will satisfy the specified Frequency (assuming no other Frequencies are connected by "<u>AND</u>"). This type of Frequency does not qualify for the extension allowed by RSR 12.0.2.

"Thereafter" indicates future performances must be established per RSR 12.0.2, but only after a specified condition is first met (i.e., the "once" performance in this example). If reactor power decreases to < 25% RTP, the measurement of both intervals stops. New intervals start upon reactor power reaching 25% RTP.

## EXAMPLE 1.4-2

SURVEILLANCE REQUIREMENTS	an takan sa
SURVEILLANCE	FREQUENCY
Not required to be performed until 12 hours after $\geq$ 25% RTP.	
Perform channel adjustment.	7 days

The interval continues whether or not the unit operation is < 25% RTP between performances.

As the Note modifies the required <u>performance</u> of the Surveillance, it is construed to be part of the "specified Frequency." Should the 7 day interval be exceeded while operation is < 25% RTP, this Note allows 12 hours after power reaches  $\geq$  25% RTP to perform the Surveillance. The Surveillance is still considered to be within the "specified Frequency." Therefore, if the Surveillance were not performed within the 7 day interval (plus the extension allowed by RSR 12.0.2), but operation was < 25% RTP,

## 1.4 Frequency

## EXAMPLES <u>EXAMPLE 1.4-3</u> (continued)

it would not constitute a failure of the RSR or failure to meet the REC. Also, no violation of RSR 12.0.4 occurs when changing MODES, even with the 7 day Frequency not met, provided operation does not exceed 12 hours with power  $\geq$  25% RTP.

Once the unit reaches 25% RTP, 12 hours would be allowed for completing the Surveillance. If the Surveillance were not performed within this 12 hour interval, there would then be a failure to perform a Surveillance within the specified Frequency, and the provisions of RSR 12.0.3 would apply.

## EXAMPLE 1.4-2

### SURVEILLANCE REQUIREMENTS

SURVEILLANCE	FREQUENCY
Only required to be met in MODE 1.	
Verify leakage rates are within limits.	24 hours

Example 1.4-4 specifies that the requirements of this Surveillance do not have to be met until the unit is in MODE 1. The interval measurement for the Frequency of this Surveillance continues at all times, as described in Example 1.4-1. However, the Note constitutes an "otherwise stated" exception to the Applicability of this Surveillance. Therefore, if the Surveillance were not performed within the 24 hour interval (plus the extension allowed by RSR 12.0.2), but the unit was not in MODE 1, there would be no failure of the RSR nor failure to meet the REC. Therefore, no violation of RSR 12.0.4 occurs when changing MODES, even with the 24 hour Frequency exceeded, provided the MODE change was not made into MODE 1. Prior to entering MODE 1 (assuming again that the 24 hour Frequency were not met), RSR 12.0.4 would require satisfying the RSR.

## 1.5 REC and RSR Implementation

The ODCM provides those limitations upon plant operations which are part of the licensing basis for the station but do not meet the criteria for continued inclusion in the Technical Specifications.

It also provides information which supplements the Technical Specifications by implementing the requirements of Technical Specification Sections 5.5.1, 5.5.4, 5.6.2, and 5.6.3.

RECs and RSRs are implemented the same as Technical Specifications (see 12.0 Applicability). However, RECs and RSRs are treated as plant procedures and are not part of the Technical Specifications. Therefore the following exceptions apply:

- Violations of the Action or Surveillance requirements in a REC are not reportable as conditions prohibited by, or deviations from, the Technical Specifications per 10 CFR 50.72 or 10 CFR 50.73.
- b. Power reduction or plant shutdowns required to comply with the Actions of a REC are not reportable per 10 CFR 50.72 or 10 CFR 50.73.

2.0 through 11.0 NOT USED

## INTENTIONALLY BLANK

Sections 2.0 through 11.0 are not used in the ODCM in order to maintain the Original ODCM numbering convention 12.0 ODCM RADIOLOGICAL EFFLUENT CONTROL (REC) APPLICABILITY

REC 12.0.1	RECs shall be met during the MODES or other specified conditions in the Applicability, except as provided in REC 12.0.2.		
REC 12.0.2	Upon discovery of a failure to meet a REC, the Required Actions of the associated Conditions shall be met, except as provided in REC 12.0.5. If the REC is met or is no longer applicable prior to expiration of the specified Completion Time(s), completion of the Required Action(s) is not required, unless otherwise stated.		
REC 12.0.3	<ul> <li>When a REC is not met and the associated ACTIONS are not met, an associated ACTION is not provided, or if directed by the associated ACTIONS, action shall be initiated within 1 hour to:</li> <li>a. Implement appropriate compensatory actions as needed;</li> <li>b. Verify that the plant is not in an unanalyzed condition or that a required safety function is not compromised by the inoperabilities; and</li> <li>c. Within 12 hours, obtain Shift Operations Superintendent or designee approval of the compensatory actions and the plan for exiting REC 12.0.3.</li> <li>Exceptions to this REC are stated in the individual RECs.</li> <li>Where corrective measures are completed that permit operation in accordance with the REC or ACTIONS, completion of the actions required by REC 12.0.3 is not required.</li> <li>REC 12.0.3 is only applicable in MODES 1, 2, and 3.</li> </ul>		
REC 12.0.4	When a REC is not met, entry into a MODE or other specified condition in the Applicability shall not be made except when the associated ACTIONS to be entered permit continued operation in the MODE or other specified		

## 12.0 REC APPLICABILITY

REC 12.0.4 (continued)	condition in the Applicability for an unlimited period of time. This REC shall not prevent changes in MODES or other specified conditions in the Applicability that are required to comply with ACTIONS or that are part of a shutdown of the unit.
	Exceptions to this REC are stated in the individual RECs.
	REC 12.0.4 is only applicable for entry into a MODE or other specified condition in the Applicability in MODES 1, 2, and 3.
REC 12.0.5	Equipment removed from service or declared inoperable to comply with ACTIONS may be returned to service under administrative control solely to perform testing required to demonstrate its OPERABILITY or the OPERABILITY of other equipment. This is an exception to REC 12.0.2 for the system returned to service under administrative control to perform the testing required to demonstrate OPERABILITY.
REC 12.0.6	RECs, including associated ACTIONS, shall apply to each unit individually, unless otherwise indicated. Whenever the REC refers to a system or component that is shared by both units, the ACTIONS will apply to both units simultaneously.

## 12.0 ODCM RADIOLOGICAL EFFLUENT SURVEILLANCE REQUIREMENT (RSR) APPLICABILITY

RSR 12.0.1 RSRs shall be met during the MODES or other specified conditions in the Applicability for individual RECs, unless otherwise stated in the RSR. Failure to meet a RSR, whether such failure is experienced during the performance of the RSR or between performances of the RSR, shall be failure to meet the REC. Failure to perform a RSR within the specified Frequency shall be failure to meet the REC except as provided in RSR 12.0.3. RSRs do not have to be performed on inoperable equipment or variables outside specified limits.

## RSR 12.0.2 The specified Frequency for each RSR is met if the RSR is performed within 1.25 times the interval specified in the Frequency, as measured from the previous performance or as measured from the time a specified condition of the Frequency is met.

For Frequencies specified as "once," the above interval extension does not apply.

If a Completion Time requires periodic performance on a "once per . . ." basis, the above Frequency extension applies to each performance after the initial performance.

Exceptions to this RSR are stated in the individual RSRs.

RSR 12.0.3 If it is discovered that a RSR was not performed within its specified Frequency, then compliance with the requirement to declare the REC not met may be delayed, from the time of discovery, up to 24 hours or up to the limit of the specified Frequency, whichever is greater. This delay period is permitted to allow performance of the RSR. A risk evaluation shall be performed for any Surveillance delayed greater than 24 hours and the risk impact shall be managed.

If the RSR is not performed within the delay period, the REC must immediately be declared not met, and the applicable Condition(s) must be entered.

When the RSR is performed within the delay period and the RSR is not met, the REC must immediately be declared not met, and the applicable Condition(s) must be entered.

RSR 12.0.4	Entry into a MODE or other specified condition in the Applicability of a REC shall not be made unless the REC's RSRs have been met within their specified Frequency. This provision shall not prevent entry into MODES or other specified conditions in the Applicability that are required to comply with ACTIONS or that are part of a shutdown of the unit.
	RSR 12.0.4 is only applicable for entry into a MODE or other specified condition in the Applicability in MODES 1, 2, and 3.
RSR 12.0.5	RSRs shall apply to each unit individually, unless otherwise indicated.