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MPC'S FOR INGESTION OF RARE EARTHS
IN COLUMBIA RIVER WATER

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MPC'S FOR INGESTION OF RARE EARTHS IN COLUMBIA RIVER WATER

by

M. W. McConiga

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MPC'S FOR INGESTION OF RARE EARTHS
IN COLUMBIA RIVER WATER

Introduction

Radioactive rare earths and other radionuclides are discharged into the Columbia River in cooling water from the HAPO reactors. Two analyses of the quantities and relative abundances of nuclides of the rare earth and yttrium analytical group in reactor effluent were started in October, 1956⁽¹⁾. Provisional MPC's for the rare earth group in drinking water derived from the Columbia River were established from the analyses based on individual rare earth MPC's of National Bureau of Standards Handbook 52 and the HAPO Radiation Protection Standards, HW-25457 Rev. 1.

A third analysis was begun in April, 1960⁽²⁾ to determine if relative abundances of rare earth nuclides in reactor effluent had changed with operational changes. MPC's for the rare earth mixture in drinking water were calculated from isotopic concentrations of the 1960 analysis and MPC's of NBS Handbook 69.

Summary

Results of the 1960 analysis plus a recalculation from corrected isotopic concentrations of the 1956 analyses indicate that limits in use since 1956 for rare earths in drinking water (GI tract, 3×10^{-4} $\mu\text{c}/\text{cc}$; bone, 0.5 $\mu\text{c}/\text{cc}$) are more restrictive than necessary and limits nearer the calculated values are recommended as follows: GI tract, 4×10^{-4} $\mu\text{c}/\text{cc}$ and bone, 0.6 $\mu\text{c}/\text{cc}$.

Discussion

The 1960 analysis was made on a composite of effluent from six operating reactors rather than from single reactors as was done previously. Some of the nuclides could not be measured directly but were estimated from cross section and relative

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abundance data using logical irradiation and decay periods. Except for terbium-160, no rare earths heavier than europium were identified and beta decay components were ascribed to the yttrium nuclide with a similar half-life. Since some of the beta components may have been partly caused by heavy rare earths, there is a possibility of error, however not of a magnitude which would significantly alter MPC's. The components concerned were of the order of 10^{-7} $\mu\text{c}/\text{cc}$.

Results

The following listings compare an average of two previous analyses and the most recent analysis as to number of rare earth nuclides detected, their concentrations, and in the latter case, the analyst's estimate of the maximum range of error.

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TABLE I

Average of Reactor Effluent Analyses October 1956		Analysis of Reactor Effluent Completed June 1960	
Isotope	Concentration Times 10^{-6} $\mu\text{c}/\text{cc}$ at 24 Hours After Discharge From Reactor	Concentration Times 10^{-6} $\mu\text{c}/\text{cc}$ at 24 Hours After Discharge From Reactor	Maximum Estimated Error
Y ⁹⁰	2.2	3.5	± 50%
Y ⁹¹	0.35	1.1	+ 25% - 90%
Y ⁹²	0.4	0.55	+ 50% - 100%
Y ⁹³	2.9	2.7	+ 50% - 100%
La ¹⁴⁰	3.4	13.0	± 15%
La ¹⁴¹	0.24	-	-
Ce ¹⁴¹	0.8	5.2*	± 25%
Ce ¹⁴³	1.1	1.4	± 25%
Ce ¹⁴⁴	-	2.6	± 25%
Pr ¹⁴²	0.8	3.6	± 30%
Pr ¹⁴³	0.25	1.8	± 15%
Pr ¹⁴⁵	0.25	0.23	± 30%
Nd ¹⁴⁷	0.37	0.57	± 30%
Pm ¹⁴⁹	0.2	0.13*	± 100%
Pm ¹⁵¹	0.1	-	-
Sm ¹⁵³	6.7	9.9	± 15%
Sm ¹⁵⁶	0.5	-	-
Eu ¹⁵² (9.3 h)	1.9	17.0	± 25%
Eu ¹⁵² (13 y)	0.006	0.76*	± 50%
Eu ¹⁵⁴	0.0003	0.03*	-
Eu ¹⁵⁵	0.015	0.4*	-
Eu ¹⁵⁶	0.25	6.3*	± 30%
Eu ¹⁵⁷	1.3	1.8	± 100%
Tb ¹⁶⁰	-	0.35	± 25%

* Denotes isotopes not measured directly.

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Rare earth group MPC's were calculated from the 1956 analyses as 3.6×10^{-4} $\mu\text{c}/\text{cc}$ for the GI tract and $0.65 \mu\text{c}/\text{cc}$ for bone. A lower set of MPC's, recommended to allow for variations in isotopic ratios, etc., and used since 1956, were 3.0×10^{-4} $\mu\text{c}/\text{cc}$ for the GI tract, and $0.5 \mu\text{c}/\text{cc}$ for bone.

The Y^{90} concentration reported on the analyses was later found to be four times too high. When recalculated, the MPC_{Bone} changed only slightly, but the MPC_{GI} changed from 3.6×10^{-4} $\mu\text{c}/\text{cc}$ to 4.6×10^{-4} $\mu\text{c}/\text{cc}$.

Rare earth group MPC's calculated in Table 2 from the 1960 analyses are 3.8×10^{-4} $\mu\text{c}/\text{cc}$ for the GI tract and $0.56 \mu\text{c}/\text{cc}$ for bone. MPC's for other organs of possible interest, kidney and total body, are also shown in Table 2.

The comparative analyses in Table I and the decay curves in Figure 1 indicate that there has been no great change in the concentrations of rare earth nuclides in the effluent from the HAP0 reactors between 1956 and 1960. No change in limits is necessary on that basis. However, it is recommended that the MPC's adopted and in use since 1956 be changed to coincide more nearly with calculated limits. The recommended MPC's for rare earths in drinking water derived from the Columbia River

are: GI tract: 4×10^{-4} $\mu\text{c}/\text{cc}$
bone: $0.6 \mu\text{c}/\text{cc}$

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TABLE 2

CALCULATED DRINKING WATER MPC'S FOR THE RARE EARTH GROUP IN REACTOR EFFLUENT AGED 24 HOURS

Rare Earth Isotope	NBS Handbook 69 MPC's x 10 ⁻⁶ µc/cc		Isotopic Congen. x 10 ⁻⁶ µc/cc	MPCGI		MPCBone		MPCTotal Body		MPCKidney		
	GI	Bone		Total Body	Kidney	C	ΣC	C	ΣC	C	ΣC	C
Y90	200	4,000,000	30,000,000	--	3.5	.04799	240.	.01199	.00159	--	--	--
Y91	300	300,000	2,000,000	--	1.1	.01508	50.3	.05026	.00754	--	--	--
Y92	600	40,000,000	300,000,000	--	0.55	.00754	12.6	.00018	.00002	--	--	--
Y93	300	20,000,000	90,000,000	--	2.7	.03702	123.	.000185	.00041	--	--	--
La140	200	20,000,000	20,000,000	--	13.0	.1783	892.	.00891	.00891	--	--	--
Ce141	900	5,000,000	10,000,000	7,000,000	5.2	.07131	79.	.01426	.00713	--	--	.01018
Ce143	400	20,000,000	50,000,000	40,000,000	1.4	.01919	48.	.00095	.00038	--	--	.00047
Ce144	100	80,000	300,000	200,000	2.6	.03565	357.	.44187	.11883	--	--	.17825
Pr142	300	30,000,000	90,000,000	60,000,000	3.6	.04936	164.5	.00164	.00054	--	--	.00082
Pr143	500	4,000,000	10,000,000	9,000,000	1.8	.02468	49.4	.00617	.00246	--	--	.00274
Pr145	---	238,000	---	---	0.23	.00315	--	.01323	--	--	--	--
Nd147	600	6,000,000	10,000,000	5,000,000	0.57	.00781	13.	.00130	.00078	--	--	.00156
Pm149	400	20,000,000	50,000,000	40,000,000	0.13	.00178	4.45	.00008	.00003	--	--	.00004
Sm153	800	50,000,000	100,000,000	80,000,000	9.9	.1358	170.	.00271	.00136	--	--	.00169
Eu152(9.3 h)	600	90,000,000	200,000,000	100,000,000	17.0	.2331	389.	.00259	.00116	--	--	.00233
Eu152(13 y)	800	200,000	200,000	90,000	0.76	.0104	13.	.05200	.05200	--	--	.11555
Eu154	200	30,000	80,000	30,000	0.038	.00052	2.6	.01733	.00650	--	--	.01733
Eu155	2000	800,000	1,000,000	700,000	0.4	.00548	2.74	.00685	.00548	--	--	.00782
Eu156	--	66,666	--	--	6.3	.0864	--	1.1296	--	--	--	--
Eu157	--	2,000,000	--	--	1.8	.0247	--	.0124	--	--	--	--
Tb160	400	800,000	1,000,000	1,000,000	0.35	.0048	12.	.0060	.00480	--	--	.00480
					ΣC = 72.92		2610.6	1.78	.2199			.34358

MPCGI is $\frac{1}{2611} = 3.8 \times 10^{-4} \mu\text{c/cc}$

MPCTotal Body is $\frac{1}{.2199} = 4.5 \mu\text{c/cc}$

MPCBone is $\frac{1}{1.78} = 0.56 \mu\text{c/cc}$

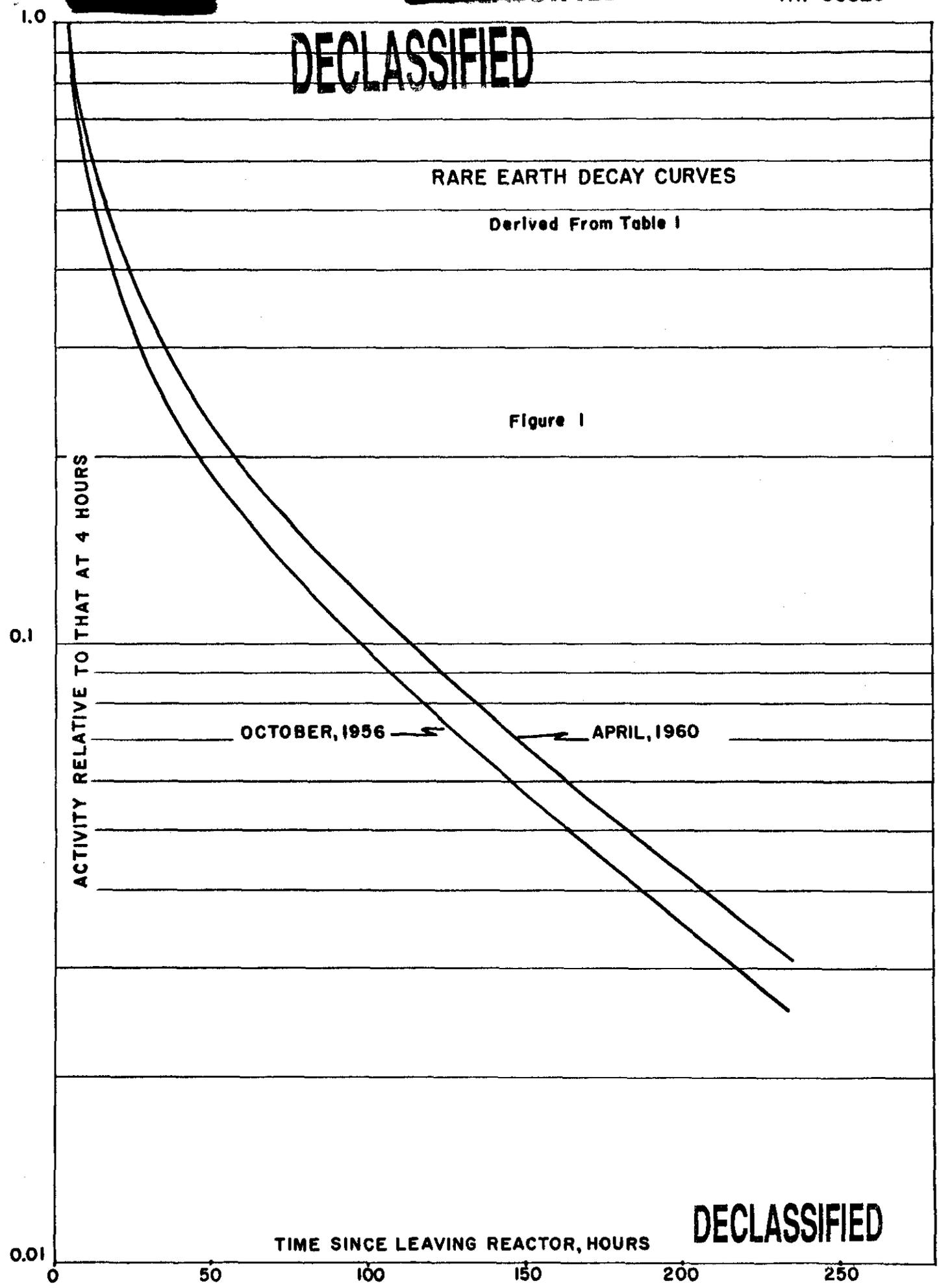
MPCKidney is $\frac{1}{.344} = 2.9 \mu\text{c/cc}$

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RARE EARTH DECAY CURVES

Derived From Table I

Figure 1



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1. Silker, W. B., and L. J. Kirby - Separation and Determination of Radioactive Rare Earths in Reactor Effluent Water. HW-48021 (Declassified), January 7, 1957.
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