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San Onofre PWR Data for Code Validation of MOX Fuel Depletion Analyses — Revision 1

O. W. Hermann



Fissile Materials Disposition Program

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ABSTRACT

The isotopic composition of mixed-oxide fuel (fabricated with both uranium and plutonium isotopes) discharged from reactors is of interest to the Fissile Material Disposition Program. The validation of depletion codes used to predict isotopic compositions of MOX fuel, similar to studies concerning uranium-only fueled reactors, thus, is very important. The EEI-Westinghouse Plutonium Recycle Demonstration Program was conducted to examine the use of MOX fuel in the San Onofre PWR, Unit I, during cycles 2 and 3. The data, usually required as input to depletion codes, either one-dimensional or lattice codes, were taken from various sources and compiled into this report. Where data were either lacking or determined inadequate, the appropriate data were supplied from other references. The scope of the reactor operations and design data, in addition to the isotopic analyses, was considered to be of sufficient quality for depletion code validation.

1. INTRODUCTION

One goal of the U.S. Department of Energy's Fissile Materials Disposition Program¹ (FMDP) is to certify the capability to predict the characteristics of mixed-oxide (MOX) spent fuel. Further, any fuel-depletion code for this purpose should be evaluated to determine how the accuracies of the computations compare with those of the more common low-enriched-uranium (LEU)-fueled lightwater reactors (LWRs). Validation studies have been performed for the SCALE code system fuel-depletion analyses of both pressurized-water reactors². (PWRs) and boiling-water reactors⁴ (BWRs). During the operation of these reactors, the plutonium gradually increases in the fuel that initially contained uranium as the only actinide. However, there is a much more dominant influence on the flux and depletion characteristics from the plutonium in the MOX fuel because the plutonium concentration is several times greater than that of spent LEU fuel and it has a greater sensitivity to the thermal spectrum.

The EEI-Westinghouse Plutonium Recycle Demonstration Program, sponsored by Edison Electric Institute, Westinghouse Electric Corporation, and the Atomic Energy Commission, was conducted from 1968 to 1974. A significant part of this program involved the measurement of isotopic compositions of the uranium and plutonium in depleted MOX fuel withdrawn from the San Onofre PWR, a reactor having a Westinghouse design and operated by Southern California Edison and San Diego Gas & Electric companies. The operating and design data typically required by fuel depletion codes can be found in a large number of sources. ⁵⁻¹² Also, the operating reports in the range from Docket 50206-79 through Docket 50206-201 contain similar data. The objective of this report is the compilation of the required data for fuel depletion codes into a single report. Data assumptions, which are considered to be reasonable, are included where the data are lacking. Even though input data for the SAS2H module of SCALE¹³ are the primary objectives of the compilation, an effort was made to include data required by lattice codes and other depletion codes.

Four MOX fuel assemblies were loaded at the start of cycle 2 of the San Onofre Nuclear Generation Station, Unit I and irradiated for both cycles 2 and 3. Then, isotopic composition analyses were conducted by the Westinghouse Electric Corporation on six sample pellets from four fuel rods of test assembly D51X. The basic parameters describing these six sample cases are listed in Table 1. Note that in all cases natural uranium was used, and three different weight percent values for fissile plutonium (²³⁹Pu + ²⁴¹Pu) were applied in the initial MOX fuel. Also, one may refer to Appendix A of this report for details of the determination of the case burnups listed in Table 1. The design data for uranium-only assemblies and core power distributions of cycles 2 and 3, which may be required in applying lattice codes for depletion calculations, are included in Appendix B.

Table 1. Basic parameters of measured MOX spent fuel samples

			_	Initial enrichment		_	
Case No.	ID of test assembly	Pin ID	Cycle No.	(atom % ²³⁵ U) ^a	(wt % fissile Pu)	Location ht ^b (in.)	Burnup ^c (MWd/MTHM) ^d
1	D51X	067	2	0.72	3.31	53.0	8,167
2	D51X	141	2	0.72	2.84	95.5	6,808
3	D51X	079	2, 3	0.72	3.10	49.0	20,891
4	D51X	167	2, 3	0.72	2.84	16.5	17,447
5	D51X	167	2, 3	0.72	2.84	95.5	18,713
6	D51X	167	2, 3	0.72	2.84	114.0	11,065

^aTaken from ref. 14.

Source: ref. 12, unless otherwise specified.

^bHeight above bottom of active fuel.

^cDetermination from ¹⁴⁸Nd measurements described in the appendix.

^dMegawatt days per metric ton heavy metal (U + Pu).

2. MOX FUEL ASSEMBLY DESIGN DATA

The reactor fuel assembly design and fuel compositions for the four San Onofre MOX assemblies loaded in cycles 2 and 3 are presented in this section. The MOX fuel assembly design data, which would be significant to depletion codes, are listed in Table 2. Most of the data are taken from a Westinghouse report (WCAP-4167-2) that was issued near the start of the demonstration program. Some of the data presented in Table 2 were calculated, as noted, from other data in the table. The average soluble boron, not found in the available sources, was estimated to be 500 ppm because the average for five PWRs in previous studies was 490 ppm. The guide tube dimensions were taken from data for the uranium-only assemblies. Although the tube sizes and boron content could have been different, the change would probably not have a significant influence on results.

The initial isotopic compositions of the uranium and plutonium in the MOX fuel of the San Onofre PWR are given in Table 3. Two deficiencies are noted in the initial plutonium data of this table. First, it is known that the plutonium is recycled plutonium from spent fuel and ²³⁸Pu/²³⁹Pu ratios are usually from 0.02 to 0.04 in typical PWR spent fuel.^{2,3} However, no ²³⁸Pu is listed in the available data, and its absence needs to be considered in evaluating the uncertainty related to predicted analytic results. The other deficiency in the initial Pu specifications is that the decay of the 14.35-year half-life ²⁴¹Pu is not considered. In the fuel-assembly fabrication program, 17 analyses were made on ²³⁹Pu and ²⁴¹Pu in revised analyses (as indicated in Table 3) for accountability. The change in ²⁴¹Pu composition, thus, can be taken into account.

The fractional composition of UO₂ and PuO₂ for the different values of wt % fissile plutonium are presented in Table 4. The weight fraction of UO₂ and PuO₂ in the MOX fuel is useful in determining fuel atomic densities or, more directly, as values input in the arbitrary material data of the fuel input to SAS2H. These weight fractions were calculated from the fuel stack density of Table 2, the initial isotopic atom compositions of Table 3 and the atomic weights¹⁴ of the U and Pu isotopes. The initial Pu isotopic data were used because the revised data are incomplete. However, the decay time for the ²⁴¹Pu to change (5.2 to 4.8%) was calculated to be 1.66 year, or 605 days. The depletion code input should have a 605-day step of decay (or very low power) that precedes the irradiation cycles, accounting for the ²⁴¹Pu change. Although the revised data are incomplete, code input may either be revised to 4.8% for ²⁴¹Pu or include the 605-day step of decay.

The locations of the four fuel pins analyzed from the MOX fuel assembly D51X, listed in Table 1, are shown in Fig. 1. A complete description of the assembly, showing the fissile plutonium enrichment pattern, is presented in Fig. 2. This information would probably be required for depletion lattice codes. The locations of the four MOX fuel assemblies during cycles 2 and 3, are shown in Fig. 3. Lattice depletion code input may use the fact that assembly D51X is adjacent to the reflector region in cycle 2.

Table 2. San Onofre MOX fuel assembly design data

Parameter	Data
Assembly general data	
Designer	Westinghouse Electric
Rod lattice	14×14
Number of assemblies/core ^a	157
MOX assemblies in cycles 2 and 3	4
Total MOX loading, metric ton heavy metal, MTHM ^b	1.335
MOX fuel/assembly, kg U + Pu ^c	333.75
Number of MOX fuel rods	180
Number of instrument tubes ^d	1
Number of guide tubes ^d	15
Equivalent core diameter, cm (in.) ^a	282 (111)
Assembly pitch, cm (in.) ^c	19.941 (7.851)
H ₂ O moderator pressure, psia ^a	2100
Average moderator temperature, K (°F) ^b	576.5 (578)
Average moderator density, g/cm ³ c	0.7179
Average clad temperature, $K (^{\circ}F)^b$	615 (648)
Soluble boron (estimated), ppm (wt) ^e	500
Fuel rod data	
Type of fuel pellet	UO ₂ plus PuO ₂ (or MOX)
Stack density, g/cm ³ ^c	10.2235
Rod pitch, cm (in.)	1.41224 (0.556)
Clad OD, cm (in.)	1.07188 (0.422)
Diametrical gap, cm (in.)	0.01905 (0.0075)
Clad thickness, cm (in.)	0.06172 (0.0243)
Clad ID, cm (in.) ^c	0.94844 (0.3734)
Pellet OD, cm (in.)	0.92939 (0.3659)
Active fuel length, cm (in.)	303.28 (119.4)
Clad material	Zircaloy-4
Guide tubes	
Material	Stainless steel-304
Tube ID, cm (in.) ^f	1.29794 (0.511)
Tube OD, cm (in.) ^f	1.35890 (0.535)

^aTaken from ref. 6.

^bTaken from ref. 5.

^cCalculated from other data in table.

^dTaken from ref. 10 or 11.

 $[^]e$ Assumed cycle 3 average boron concentration equals that of cycle 2.

^fAssumed similar to that of U assemblies, taken from ref. 7.

Source: ref. 9, unless otherwise specified.

Table 3. Initial compositions of the uranium and plutonium

Isotope	Atom % in U or Pu, ppm ²⁴¹ Am						
Uranium ^a							
^{234}U ^{235}U ^{238}U	0.0055 0.7200 99.2745						
Plutonium ^b	Initial	Revised	Decay time, d				
²³⁹ Pu ²⁴⁰ Pu ²⁴¹ Pu ²⁴² Pu	80.6 13.4 5.2 0.8	80.7 - ^c 4.8 - ^c	$-^d$ $-^d$ 605^e $-^d$				
Other							
$^{241}\text{Am}^f$	5000						

^aTaken from ref. 14.

Table 4. Fractional composition of UO_2 and PuO_2 in the MOX fuel

	Pin	No. of same	Wt %	Weight fraction in	$MOX (UO_2 + PuO_2)$
Case	ID	type pins	fissile Pu	of UO_2	of PuO ₂
1	067	24	3.31	0.961423	0.038577
2	141	64	2.84	0.966901	0.033099
3	079	92	3.10	0.963870	0.036130
4, 5, 6	167	64	2.84	0.966901	0.033099

^aComputed from data in Tables 2 and 3 and atomic weights of ref. 14.

^bInitial from ref. 5; revised from ref. 9.

^cNo revised measurement reported.

^dDecay time not calculated.

^eCalculated decay time required to change ²⁴¹Pu from the initial to the revised composition.

^fIn ppm (wt) of Pu, from ref. 9.

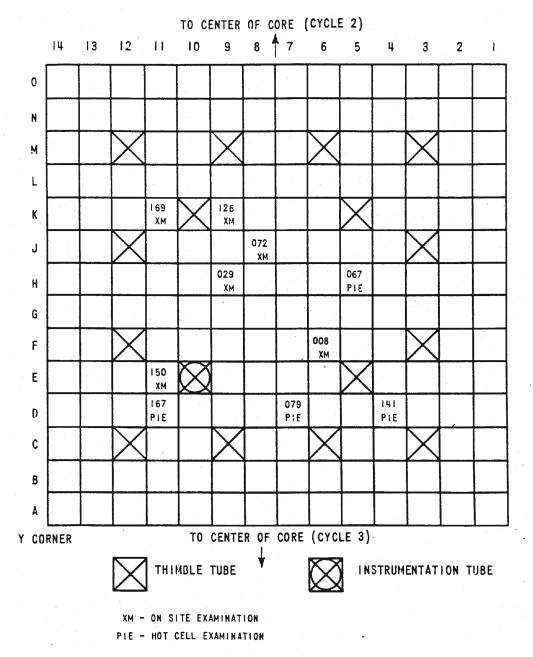


Fig. 1. Location of rods removed from San Onofre Plutonium Demonstration Assembly D51X for post-irradiation examination. *Source:* ref. 11.

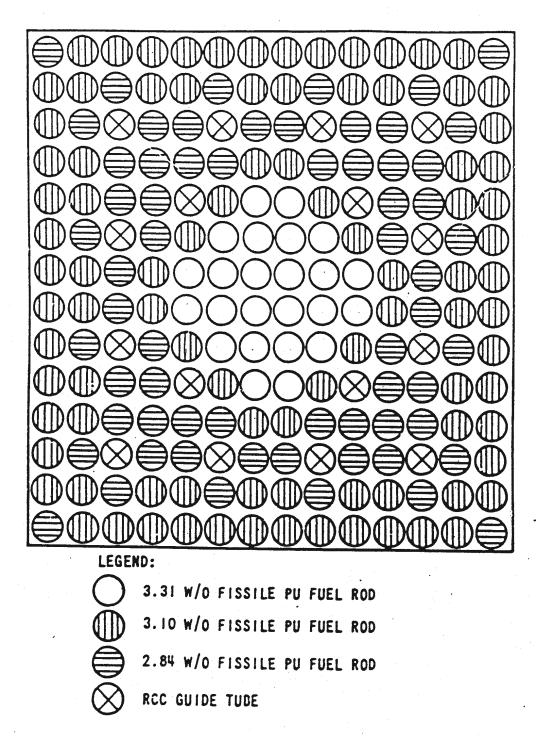


Fig. 2. Enrichment pattern for the four plutonium assemblies. *Source*: refs. 6 and 9.

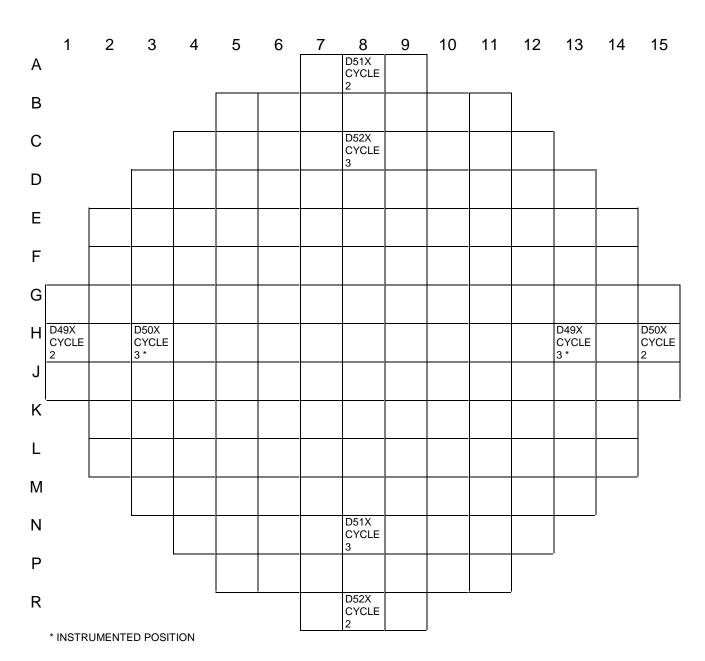


Fig. 3. Location of plutonium demonstration assemblies in San Onofre cycle 2 and cycle 3. *Source*: refs. 10 and 11.

3. THE PWR OPERATIONS DATA AND MODEL OF ASSEMBLIES

The San Onofre PWR, Unit I, operations data, pertaining to the MOX fuel assemblies in cycles 2 and 3 are presented in Table 5. The pellet sample burnups were derived from the ¹⁴⁸Nd measurements, as described in the appendix. The average cycle power experienced by each of the six pellet samples were calculated from the burnup and cycle times.

Portions of the operating conditions were listed in Table 2. The clad and water moderator temperatures⁵ are in the "assembly general data" listed in the table. The moderator density was obtained by interpolation of data in the temperature-pressure-density table¹³ at 578°F and 2100 psia.⁶

The effective fuel temperatures, applied in the resonance treatment, were obtained from the fuel-temperature-vs-rod linear power curve² in Fig. 4. The curve was developed for the Obrigheim PWR. There are similarities in lattice (14×14), pellet OD (<1% difference) and moderator temperature (differ by 4.5 K) between the San Onofre and Obrigheim reactors. This method of estimating fuel temperature was applied in the validation of H. B. Robinson PWR analyses.² The resulting temperatures are given in Table 5.

The unit cell zone geometry data for Path-B of SAS2H is listed in Table 6.

It is recommended that cross sections should be determined during the neutronics calculation for all nuclides that are significant to the results, as recommended in the most current validation study⁴ and a burnup-credit sensitivity study.¹⁵

Table 5. San Onofre operating data, including sample pellet powers and resonance-type fuel temperatures

Operation data type, Pin ID	ruei temp	cratures		Total
(height, in.)	Units	Cycle 2	Cycle 3	burnup
Cycle times ^a			•	
Startup date		11/18/70	3/1/72	
Shutdown date		12/26/71	6/2/73	
Uptime	Days	403	459	
Downtime	Days	66	<u>_</u> b	
Fuel pellet burnups ^c	MWd/MTHM			
067 (53.0)		8,167	_	8,167
141 (95.5)		6,808	_	6,808
079 (49.0)		7,015	13,877	20,891
167 (16.5)		5,999	11,448	17,447
167 (95.5)		6,434	12,279	18,713
167 (114.0)		3,843	7,222	11,065
Fuel pellet powers ^d	MW/MTHM			
067 (53.0)		20.266	_	
141 (95.5)		16.894	_	
079 (49.0)		17.406	30.232	
167 (16.5)		14.885	24.942	
167 (95.5)		15.965	26.751	
167 (114.0)		9.536	15.735	
Effective fuel temperatures ^e	K			
067 (53.0)		744	_	
141 (95.5)		713	_	
079 (49.0)		718	839	
167 (16.5)		695	787	
167 (95.5)		705	805	
167 (114.0)		650	703	
Shutdown to analysis times ^b	Days			
067 (53.0)		717	_	
141 (95.5)		719	_	
079 (49.0)			194	
167 (16.5)			187	
167 (95.5)			187	
167 (114.0)			192	

^aData taken from refs. 10 and 11.

^bTime from cycle shutdown to sample analysis.
^cDetermined in the appendix from ¹⁴⁸Nd measurements.

^dCalculated directly from the burnup- and cycle-time data.

^eEffective fuel temperature for resonance treatment.

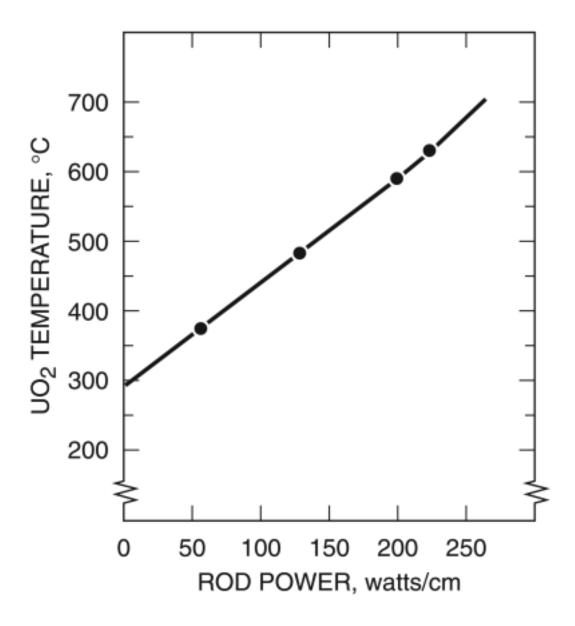


Fig. 4. Fuel-temperature-vs-rod power for Obrigheim. Source: ref. 2.

Table 6. Effective SAS2H geometry of the San Onofre MOX assembly model

Cycle	Radial zone	Mixture No.	Composition	Effective radius (cm)
2	1	3	Borated moderator	0.64897
2	2	5	Stainless steel-304	0.67945
2	3	3	Borated moderator ^a	0.79677
2	4	500	Homogenized fuel, clad and borated moderator	2.78870
2	5	3	Borated moderator between assemblies ^b	2.81266
3	1	3	Borated moderator	0.64897
3	2	5	Stainless steel-304	0.67945
3	3	3	Borated moderator ^a	0.84510
3	4	500	Homogenized fuel, clad and borated moderator	2.78870
3	5	3	Borated moderator between assemblies ^b	2.81266

^aRadius different in cycles 2 and 3 to account for the removal of the two pins indicated in Tables 1 and 4.

^bCalculated from assembly pitch and equivalent core diameter in Table 2.

4. ISOTOPIC MEASUREMENTS FOR MOX FUEL SAMPLES

Samples from the San Onofre MOX spent fuel were prepared at the Battelle Memorial Institute hot-cell facility (Columbus, Ohio). These samples were sent to the Westinghouse Waltz Mill Analytical Laboratory for the comprehensive spectrometric analysis of isotopic concentrations. The measured isotopic compositions, as atom ratios, of the six MOX sample pellets are given in Table 7. These results pertain to the listed date of analysis. The results for the uranium and plutonium isotopes adjusted to the end of the irradiation time of the sample are presented in Table 8.

The first 11 isotopic ratios (through ¹⁴⁸Nd/²³⁸U) that are listed in Table 7 were performed by mass spectrometric methods. The remaining five ratios in the table were measured by alpha spectrometry. Although these five cases were analyzed later, they were decay corrected in ref. 12 to the dates given in the table for consistency.

Table 7. Measured isotopic compositions of San Onofre MOX spent fuel in atom ratios at time of analysis

		at thi	ic of analysis			
Pin ID (height, in.)	067 (53.0)	141 (95.5)	079 (49.0)	167 (16.5)	167 (95.5)	167 (114.0)
Date of analysis	12/11/73	12/13/73	12/13/73	12/06/73	12/06/73	12/11/73
Days after shutdown	717	719	194	187	187	192
Burnup,						
MWd,MTHM	8,167	6,808	20,891	17,447	18,713	11,065
$^{234}\text{U/U} \times 10^{-2}$	0.005	0.006	0.005	0.005	0.005	0.005
$^{235}\text{U/U} \times 10^{-2}$	0.628	0.641	0.470	0.483	0.479	0.569
$^{236}\text{U/U} \times 10^{-2}$	0.023	0.018	0.052	0.050	0.051	0.032
$^{238}\text{U/U} \times 10^{-2}$	99.344	99.335	99.473	99.462	99.465	99.394
238 Pu/Pu × 10^{-2}	0.557	0.462	0.989	0.860	0.884	0.642
239 Pu/Pu × 10^{-2}	71.886	73.218	56.998	57.626	57.130	66.193
240 Pu/Pu × 10^{-2}	19.050	18.812	26.422	26.613	26.593	22.401
241 Pu/Pu × 10^{-2}	7.210	6.384	12.530	12.047	12.444	9.088
242 Pu/Pu × 10^{-2}	1.295	1.124	3.061	2.854	2.949	1.678
239 Pu/ 238 U × 10 ⁻²	2.619	2.293	1.741	1.594	1.601	1.965
$^{148}Nd/^{238}U \times 10^{-4}$	1.508	1.250	3.875	3.226	3.460	2.046
241 Am/ 239 Pu × 10^{-2}	_ ^a	<u>_</u> a	<u>_</u> a	6.51	6.83	1.59
243 Am/ 239 Pu × 10-2	_a	<u>_</u> a	<u>_</u> a	1.41	1.55	0.27
236 Pu/ 239 Pu × 10^{-9}	4.04	4.60	17.7	12.4	13.4	6.50
238 Pu/ 239 Pu $\times 10^{-3}$	7.65	6.16	17.0	14.7	15.2	9.43
237 Np/U × 10 ⁻⁵ b	_a	<u>_</u> a	<u>_</u> a	9.7	11.1	5.7

^aNo measurement.

Source: ref. 12.

^bIn place of atom ratio, units are grams ²³⁷Np/g U.

Although there were no indications that uncertainties or standard deviations were developed for the measured data, the reanalysis of three samples "showed excellent agreement between initial and repeat results."

Table 8. Measured isotopic compositions of San Onofre MOX spent fuel in atom ratios adjusted to shutdown time

Pin ID (height, in.)	067 (53.0)	141 (95.5)	079 (49.0)	167 (16.5)	167 (95.5)	167 (114.0)
Burnup, MWd/MTHM	8,167	6,808	20,891	17,447	18,713	11,065
$^{234}\text{U/U} \times 10^{-2}$	0.005	0.006	0.005	0.005	0.005	0.005
$^{235}\text{U/U} \times 10^{-2}$	0.628	0.641	0.470	0.483	0.479	0.569
$^{236}\text{U/U} \times 10^{-2}$	0.023	0.018	0.052	0.050	0.051	0.032
$^{238}\text{U/U} \times 10^{-2}$	99.344	99.335	99.473	99.462	99.465	99.394
238 Pu/Pu $\times 10^{-2}$	0.560	0.465	0.986	0.857	0.880	0.639
239 Pu/Pu × 10^{-2}	71.345	72.729	56.744	57.379	56.877	65.979
240 Pu/Pu × 10^{-2}	18.907	18.686	26.304	26.499	26.495	22.328
241 Pu/Pu \times 10 ⁻²	7.903	7.003	12.919	12.423	12.831	9.382
242 Pu/Pu × 10^{-2}	1.285	1.116	3.047	2.842	2.936	1.673
239 Pu/ 238 U × 10^{-2}	2.619	2.293	1.741	1.594	1.601	1.965

Source: ref. 12.

5. SUMMARY

One of the important functions of the Fissile Materials Disposition Program is to calculate the isotopic composition of mixed-oxide fuel withdrawn from reactors. The validation of depletion codes for this purpose have been conducted²⁻⁴ for uranium-only fueled light-water reactors.

During the period from 1968 to 1974 the EEI-Westinghouse Plutonium Recycle Demonstration Program was established to examine the use of MOX fuel. One of the reactors used in the program was the San Onofre PWR Unit I. Many reports of the preliminary plans, progress and summaries were written during the program. The main purpose of this report is to compile the assembly design and cycle operating data of the MOX fuel assemblies into a single report. It is intended that the latest and required data for most depletion codes is included in this compilation. One of the codes in which the data may be applied is the SAS2H¹³ control module of the SCALE system. Also, data which may be required for lattice codes are included (Figs. 1, 2 and 3). The results of both mass and alpha spectrometer analyses, producing 16 atom ratios of significant isotopes, are presented in Table 7.

An effort was made to have the correct reference given for each item of data. When the item was calculated from other data, a reference was given to the proper tables. In a few cases, where data were lacking, the method of estimation was explained. The burnup data reported, ¹² having been calculated in 1974, was not considered accurate. A more universally applied method, as covered in Appendix A, was used for converting ¹⁴⁸Nd data to burnup.

Some depletion codes require design and operating data for fuel assemblies that do not contain MOX fuel. Appendix B contains design data of the uranium-only fuel assemblies and available core power distributions, that may be required for lattice-type depletion codes.

It was concluded that the comprehensive design and operating data and the isotopic MOX spent fuel measurements for the San Onofre PWR were of adequate quality for the proper validation of depletion codes. Calculations that were performed with SAS2H for the six samples using the data of this report indicated that the data were adequate.

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APPENDIX A SAMPLE BURNUP DETERMINATIONS

APPENDIX A

SAMPLE BURNUP DETERMINATIONS

This appendix provides an expanded explanation regarding the sample burnup data listed in Table 5. First, the reasons are given why the available burnup data^{A,1} are considered inadequate, probably due to the use of methods that have become obsolete. Then, the method applied for converting the ¹⁴⁸Nd measurement to burnup is referenced and discussed. Finally, the total burnup results and the separate cycle data are shown.

There is a high degree of confidence in the ANSI/ASTM standards for converting ¹⁴⁸Nd measurements to burnup in nuclear research concerning uranium-only fueled LWRs. A total of 68 spent fuel samples from eight different reactors have been used in comparisons of measured and calculated isotopic results in previous validation studies. A 148 The referenced burnups were, except for one reactor, derived from the ANSI/ASTM analysis. In 40 of the sample cases, from five reactors, the ANSI/ASTM measured result was given and compared with the SAS2H calculation that applied the ANSI/ASTM derived burnup. The computed/measured ANSI/ASTM differences in the 40 cases ranged form -2.4 to 1.7%, with an average of -0.32%. The averages for the five reactors were -1.4, 1.7, 0.4, -2.0 and 0.3%, with an average of 0.20%. Or, the average of the absolute values of the reactor averages was 1.16%.

The preceding comparisons of analyses and SAS2H results validate that the methods of converting ¹⁴⁸Nd measurements to burnups are adequate. However, preliminary computations, using SAS2H, for the MOX fuel cases gave ¹⁴⁸Nd differences as large as 4% for 1-cycle cases and 12% for 2-cycle cases. Although the reports giving data that were used in the validation study did not usually give the conversion factors used in determining burnup, the data^{A.5} for the Turkey Point PWR did contain the required conversion factors. Thus, the decision was made to apply the conversion factors used with the Turkey Point data in deriving burnups for the MOX fuel for the following reasons. First, applying these factors showed that the referenced burnups^{A.1} were too high by approximately the same differences indicated by the ¹⁴⁸Nd code-to-measured-results comparisons. Using the same factors from the Turkey Point PWR on data for the Calvert Cliffs PWR, the Cooper BWR and the JPDR BWR of 18 cases gave values within about 1% of the reported burnups. The discussion on the Turkey Point PWR referred to the ANSI/ASTM Standards E321-75 and E267, and that on the Calvert Cliffs PWR and Cooper BWR referred to the ANSI/ASTM Standard E321-79 and E219.

One more question should be considered before using the above ¹⁴⁸Nd-burnup conversion method (i.e., factors applied in Turkey Point PWR cases) for MOX spent fuel. Higher concentrations of Pu isotopes and lower concentrations of U isotopes are noted in the average MOX fuel than in the average uranium-only fuel during irradiation. There are two factors that this can change: the average fission fraction of ¹⁴⁸Nd production, and the average recoverable energy per fission. The percent fission fractions (based on thermal reactors) for ¹⁴⁸Nd (or mass 148) listed in ENDF/B-V^{A.6} are 1.670 for ²³⁵U, 2.081 for ²³⁸U (fast), 1.635 for ²³⁹Pu, and 1.990 for ²⁴¹Pu. An approximate average fission fraction for a given case may be calculated by weighting these four fission fractions by the average isotopic density times the average microscopic fission cross section of the corresponding isotope (or, rather, the isotopic average macroscopic fission cross section). In two SAS2H trial

cases, average macroscopic fission cross sections of the isotopes were estimated from atomic densities and cross sections listed in the case outputs. An estimate of the sum of the weighted fission fractions indicated a 1 to 2.5% increase in the average fission fraction from MOX fuel compared with that from typical uranium-only fuel. The ORIGEN-S code computes and writes the total recoverable energy per fission for each time step. The values of the average energy per fission of MOX fueled reactors compared with uranium-only reactors indicate an increase of approximately 1 to 1.5%. The fission fraction is in the denominator, and the energy per fission is in the numerator in converting the ¹⁴⁸Nd/U atom ratio to burnup. Thus, it is estimated that the burnup could be no greater than 1.5% less for MOX fuel than that produced by the conversion method used for uranium-only fuel cases.

The detailed procedure for converting the measured atom ratio of 148 Nd/ 238 U to burnup by using the same conversion factors, C and F_{148} , applied in deriving the Turkey Point data, $^{A.5}$ is the following:

$$B = RCAF_{u}(1-D_{238}) / F_{148} , \qquad (A.1)$$

where

B = calculated burnup, MWd/MTHM,

R = atom ratio of (final ¹⁴⁸Nd atoms)/(final ²³⁸U atoms),

 F_{148} = fission fraction for mass 148 = 0.0168, A.5

C = 9600 (MWd/MTHM)/at. % of fuel which fissions, A.5

A = 238 U at. % in natural U, or 99.2745,

 F_u = weight fraction of UO_2 in the MOX from Table 4,

 D_{238} = ^{238}U fractional depletion, 0.007 per cycle estimated.

 F_u should be the atom fraction of U in the U + Pu of the MOX, because each term of the equation is always the ratio of atoms. For the sample compositions, the atom ratio of U to U + Pu is not significantly different than the weight ratio of UO_2 to UO_2 + PuO_2 . The estimate of D_{238} is 0.007 ± 0.002 , which should cause an error in B no greater than 0.2%. The method in ref. A.5 applied the same factors C and F_{148} in converting R to B, but equivalent measurements were used instead of AF_u (1 - D_{238}).

As an example of using Eq. (A.1), consider the 2-cycle case for pin ID 079, where the value of F_u from Table 4 is 0.963870 and $R = 3.875 \times 10^{-4}$:

$$B = 3.875 \times 10^{-4} \times 9600 \times 99.2745 \times 0.96387 \times 0.986 / 0.0168$$

= 20,891.4 MWd / MTHM. (A.2)

The reported burnup of 23,500 MWd/MTHM is 12.5% greater than that determined in Eq. (A.2).

Burnups of the six cases computed by Eq. (A.1) are listed in Table A.1. The revised values are given in Table 5. It is seen that the old, or reported, burnups^{A.1} are significantly greater than the revised burnups.

The calculation of the burnup for each cycle of the samples irradiated in both cycles 2 and 3 are given in Table A.2. The use of the reported^{A.1} linear power for both cycles of the samples permitted the calculation of linear burnups, the fraction of total burnup per cycle, and the final burnup by cycle. These resulting burnups are listed in Table 5, in addition to the computed cycle powers required in depletion code input.

Table A.1. Data applied in determining recommended burnup^a

	No. of			% fissile	% fissile		B, MWd/MTHM	
Case	R	cycles	1-D ₂₃₈	Pu	F_{u}	revised	old	% diff. ^b
1	1.508×10^{-4}	1	0.993	3.31	0.961423	8,167	8,700	6.5
2	$1.250\times10^{\text{-4}}$	1	0.993	2.84	0.966901	6,808	7,200	5.8
3	3.875×10^{-4}	2	0.986	3.10	0.963870	20,891	23,500	12.5
4	$3.226\times10^{\text{-4}}$	2	0.986	2.84	0.966901	17,447	19,800	13.5
5	3.460×10^{-4}	2	0.986	2.84	0.966901	18,713	21,200	13.3
6	$2.046\times10^{\text{-4}}$	2	0.986	2.84	0.966901	11,065	12,500	13.0

^aDerived B by applying Eq. (A.1), the constants F_{148} , C and A, in addition to data in this table. $^b(B_{old}/B_{revised}-1)$ 100%.

Table A.2. Calculation of burnup for each cycle for 2-cycle cases

	P _{linear}	kw/ft	Uptii	me, d	B _{linear} , kwd/ft		Fraction B _{total}		B _{cycle} , MWd/MTHM	
	Су	cle	Су	cle	Cycle		Cycle		Cycle	
Case	2	3	2	3	2	3	2	3	2	3
3	3.8	6.6	403	459	1,531.4	3,029.4	0.335774	0.664226	7,015	13,877
4	3.7	6.2	403	459	1,491.1	2,845.8	0.343817	0.656183	5,999	11,448
5	3.7	6.2	403	459	1,491.1	2,845.8	0.343817	0.656183	6,434	12,279
6	2.0	3.3	403	459	806.0	1,514.7	0.347309	0.652691	3,843	7,222

 $^{^{}a}P_{linear}$ is linear power at the sample height in ref. A.1.

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APPENDIX B

URANIUM-ONLY FUEL ASSEMBLY DESIGN DATA AND CORE POWER DISTRIBUTIONS

APPENDIX B

URANIUM-ONLY FUEL ASSEMBLY DESIGN DATA AND CORE POWER DISTRIBUTIONS

The data included here, essentially, are the design data of the uranium-only fuel assemblies and core power distributions during the cycles in which MOX fuel was irradiated. Also, the calculated axial buckling and $k_{\it eff}$ versus cycle 2 burnup is included. There were significantly less data available for cycle 3. Two of the analyzed samples were taken from fuel rods removed at the end of cycle 2 and the other four samples were from rods irradiated during cycle 3, also. The data here may be used with that previously given in this report to allow modeling of the San Onofre reactor core (MOX and U-only fuel assemblies) if desired.

B.1 Uranium-Only San Onofre PWR Fuel Assemblies

Although there were differences in the fuel for the different regions in the core, the basic design data^{B.1} of all uranium-only fuel assemblies remained the same. The assembly design data for the uranium-only fuel are presented in Table B.1. Although some of the design data were the same as that for the MOX assemblies, the data were repeated for completeness.

The uranium-only fuel assembly data that varies by region in core are shown in Table B.2. The data includes the ²³⁵U enrichments, pellet densities and the uniform stack densities of the fuel by region. The pellet dishing, or the volume lost from rounding off the ends of the pellets, was used to convert to the final stack density.

Table B.1. San Onofre uranium-only fuel assembly design data

Parameter	Data
Assembly general data	
Rod lattice	14×14
Number of fuel rods	180
Number of instrument tubes ^a	1
Number of guide tubes ^a	15
Assembly pitch, cm (in.) b	19.941 (7.851)
H ₂ O moderator pressure, psia ^c	2100
Average moderator temperature, $K (^{\circ}F)^d$	576.5 (578)
Average moderator density, g/cm ³ b	0.7179
Average clad temperature, K (°F) ^d	615 (648)
Soluble boron, ppm (wt) ^{d,e}	500
Fuel rod data	
Type of fuel pellet	UO_2
Stack density	Variable (Table B.2)
Rod pitch, cm (in.)	1.41224 (0.556)
Clad OD, cm (in.)	1.07188 (0.422)
Diametrical gap, cm (in.)	0.01397 (0.0055)
Clad thickness, cm (in.)	0.04191 (0.0165)
Clad ID, cm (in.) ^f	0.98806 (0.3890)
Pellet OD, cm (in.)	0.97409 (0.3835)
Active fuel length, cm (in.)	304.8 (120)
Clad material	Stainless steel-304
²³⁵ U enrichment	Variable (Table B.2)
Guide tubes	
Material	Stainless steel-304
Tube ID, cm (in.)	1.29794 (0.511)
Tube OD, cm (in.)	1.35890 (0.535)

^aTaken from ref. B.2 or B.3.

Source: Ref B.1, unless otherwise specified.

^bTaken from Table 2.

^cTaken from ref. B.4.

^dTaken from ref. B.5.

^eAssumed cycle 3 average boron concentration equal to that of cycle 2.

^fCalculated from other data in table.

Table B.2. San Onofre uranium-only fuel assembly data that varies by region in core

	Data by Region							
Parameter	1	2	3	4U	4Pu	5		
Enrichment, wt-% ²³⁵ U	3.16	3.40	3.86	4.00	0.71	4.00		
Pellet density, % Theoretical	93.8	94.0	92.7	93.2	91.0	92.9		
Pellet dishings, % of cylindrical volume ^a	6.87	6.87	6.87	6.87	6.87	6.87		
Stack density, b g/cm ³	9.574	9.595	9.462	9.513	10.224^{c}	9.482		
Number in cycle 2 ^d	1	52	52	48	4	0		
Number in cycle 3 ^e	1	0	52	48	4	52		

^aCalculated for MOX fuel from fuel loadings, volume and % theoretical density and assumed equal for uranium fuel.

Source: Ref. B.6, unless otherwise specified.

^bApplied, for uranium-only assemblies, the pellet density, the % dishing and a UO₂ theoretical density of 10.96 g/cm³.

^cTaken from Table 2.

^dTaken from ref. B.5.

^eTaken from ref. B.1.

B.2 San Onofre Core Power Distributions

The quality of quarter-core power distributions determined for cycle 2 of the San Onofre PWR are significantly better than those available for cycle 3. This is true because the cycle 2 power distributions applied data^{B.5} available after the shutdown of cycle 2, whereas the power distributions for cycle 3 were preliminary estimations^{B.1} made prior to the loading of cycle 2 fuel.

There were three methods of deriving quarter core power distributions of cycle 2. First, there was the preliminary estimation method^{B.1} made prior to cycle 2. Second, there was a partly experimental method^{B.5} using flux levels determined from detectors located in the reactor core. The flux measurements were processed using the INCORE code, a Westinghouse proprietary code. The third method^{B.5} used the PDQ-7/HARMONY code package for two-dimensional depletion calculations with input cross sections obtained from zero-dimensional computations by the LASER-M code.

The study^{B.5} utilizing the third method of computing quarter core power distributions chose the second method as a reference because it was partly experimental and more accurate than the first method. The calculated values of axial buckling^{B.7} and $k_{\it eff}$ as a function of cycle time and burnup, applied in the study^{B.5} using the third method of deriving power distributions, are presented in Table B.3.

Quarter core power distributions^{B.5} derived by the third method and compared with the reference (method 2) for different burnups of cycle 2 are shown in Figs. B.1–B.4, inclusive. Fig. B.1 contains the core region numbers, the reference burnups, and the quarter core power distributions of the fuel assemblies for the beginning of cycle 2. Fig. B.2 shows the calculated versus reference power distributions for the beginning of cycle 2. Figs. B.3 and B.4 show the calculated versus reference power distributions at cycle 2 burnups of 3342 and 6045 MWd/MTM, respectively.

As stated above, the quarter-core power distributions available for cycle 3 were preliminary calculations made prior to loading of cycle 2 fuel. These were less accurate because both the fuel assembly burnups and the flux levels in the core were not measured prior to the calculation. These preliminary quarter-core power distribution calculations^{B.1} for the beginning of cycle 3 are shown in Fig. B.5. No other calculations for cycle 3 were reported. It may be noted that the location of the two MOX assemblies are inward by one position from the actual location in Fig. 3. In order to help determine the quality (or error) of cycle 3 calculations, it might be useful to note the similarly derived power distributions for the beginning of cycle 2 shown in Fig. B.6.

Table B.3. Calculated axial buckling and $k_{\it eff}$ versus cycle 2 time and burnup

Time (hours)	Cycle Burnup (MWd/MTM)	Accumulated Burnup (MWd/MTM)	Axial Buckling ^a (cm ⁻²)	$k_{\it eff}$
0	0.0	8828.2	3.16×10^{-2}	1.0065
100	88.5	8916.7	3.16×10^{-2}	1.0051
1910	1691.3	10519.7	3.32×10^{-2}	1.0043
3755	3300.4	12128.6	3.44×10^{-2}	1.0046
5620	4905.1	13733.3	3.56×10^{-2}	1.0040
6792	5905.2	14733.4	3.64×10^{-2}	1.0031
9110	7858.0	16686.0	3.72×10^{-2}	1.0012

^aTaken from ref.B.7.

Source: Ref. B.5 unless otherwise specified.

	Core Center Line	9	10	11	12	13	14	15	
Н	18131 0.924	2 14560 1.039	3 8996 1.148	2 18464 0.974	2 17446 0.952	2 17490 0.919	2 17733 0.883	4Pu 0 0.740	
J	2	3 7563 1.174	2 16930 1.016	3 14537 1.087	3 8028 1.166	2 16283 1.016	4 0 1.110	4 0 0.620	
K	3	2	2 13525 1.056	3 9141 1.186	2 18264 1.044	3 11768 1.134	4 0 0.946		
L	2	3	3	2 18161 1.034	3 14364 1.110	4 0 1.166	4 0 0.673		
M	2	3	2	3	3 7575 1.050	4 0 0.748	Key	, ,	
N	2	2	3	4	4		Region No. Ref. Burnup (MWd/	* MTM)	
P	2	4	4	4			Ref. Assemb	oly Power	
R	4Pu	4		*See 7	Γable B.2 for	r enrichmen	t		

Fig. B.1. Reference quarter core power distribution and assembly burnups for beginning of cycle 2.

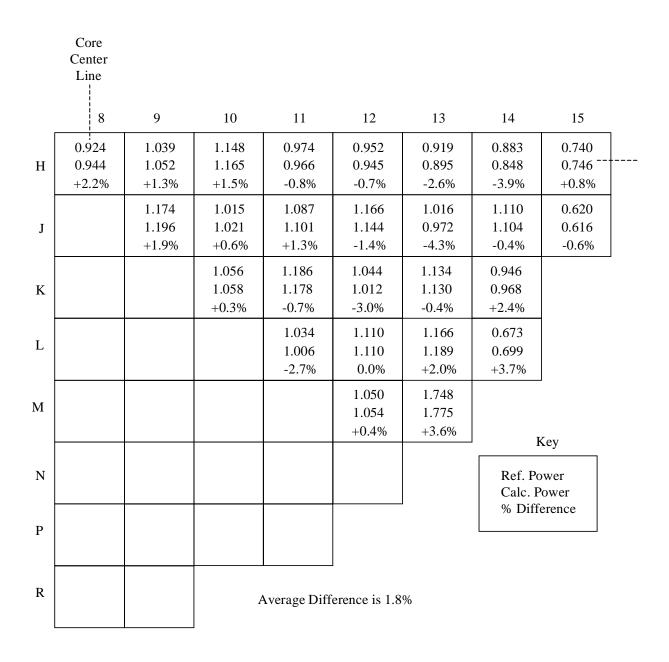


Fig. B.2. Calculated versus reference powers for beginning of cycle 2.

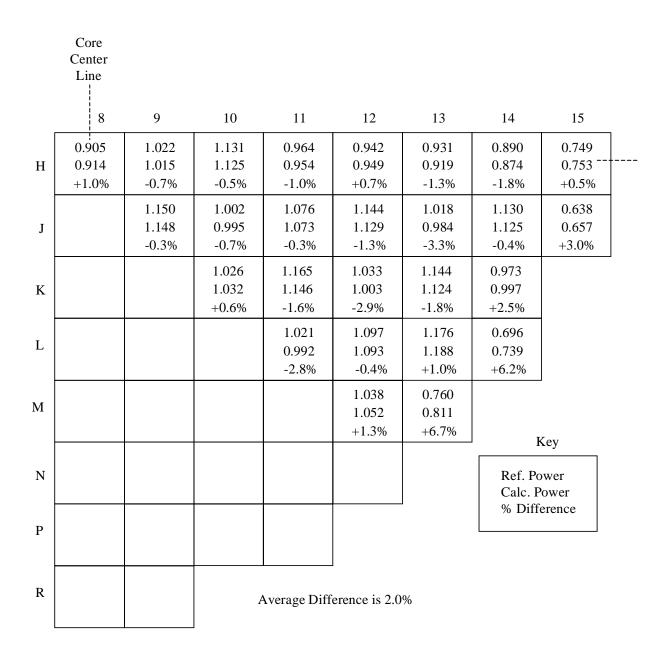


Fig. B.3. Calculated versus reference powers at 3342 MWd/MTM.

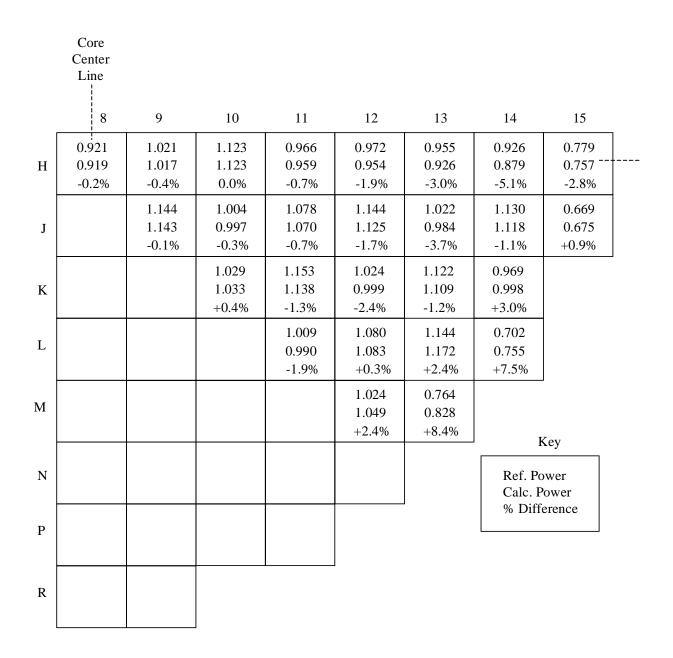


Fig. B.4. Calculated versus reference powers at 6045 MWd/MTM.

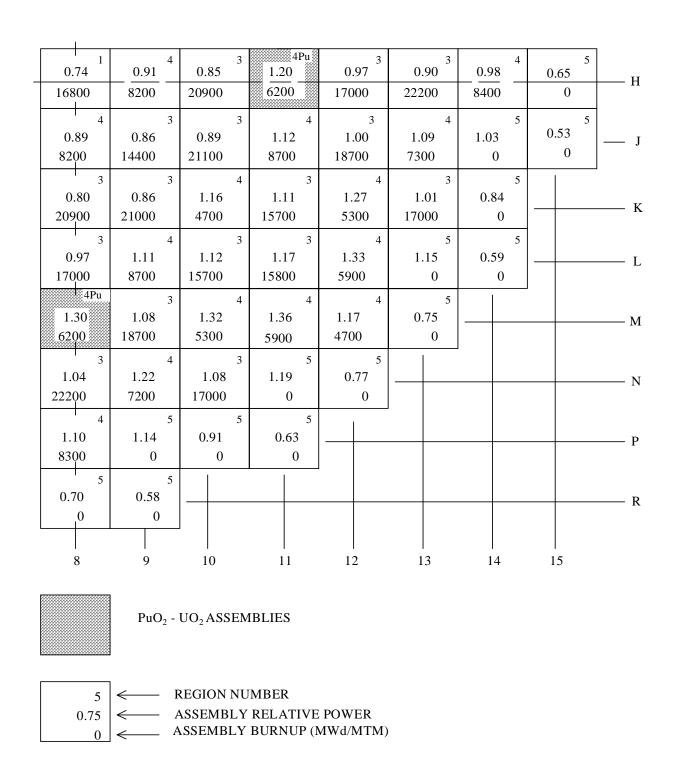


Fig. B.5. Power and burnup distributions San Onofre cycle 3, BOL.

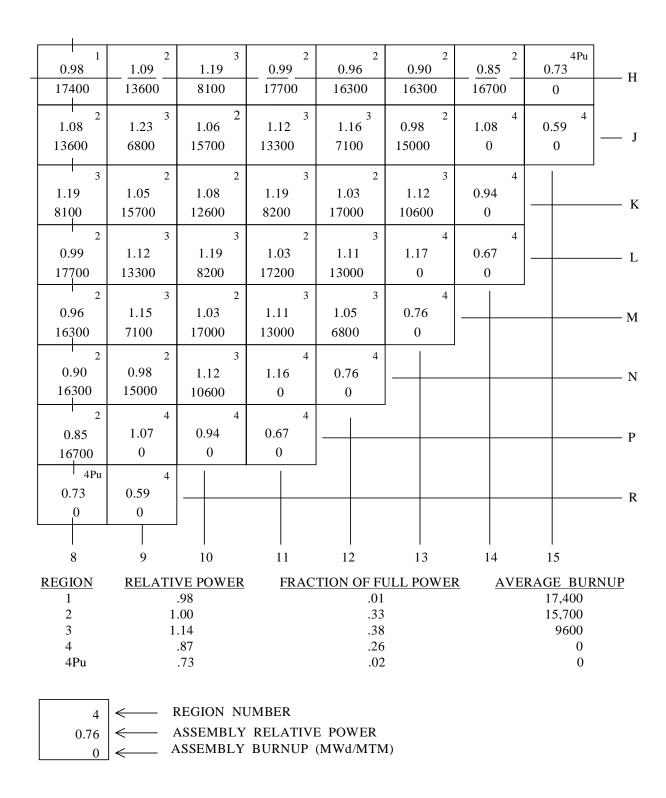


Fig. B.6. Power and burnup distributions San Onofre cycle 2, BOL.

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