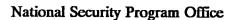
MARTIN MARIETTA



ANALYSIS OF HEU SAMPLES FROM THE ULBA METALLURGICAL PLANT

E. H. Gift
National Security Programs Office
Martin Marietta Energy Systems, Inc.
Oak Ridge, Tennessee

Initially Issued July 1994

Revised by A. W. Riedy

May 1995

MANAGED BY
MARTIN MARIETTA ENERGY SYSTEMS, INC.
FOR THE UNITED STATES
DEPARTMENT OF ENERGY

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National Security Program Office

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Prepared for the
U.S. DEPARTMENT OF ENERGY
OFFICE OF ARMS CONTROL AND NONPROLIFERATION

Prepared by
MARTIN MARIETTA ENERGY SYSTEMS, INC.
Oak Ridge, Tennessee 37831
under Contract No. DE-AC05-84OR21400
with the
U.S. DEPARTMENT OF ENERGY

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SUMMARY

In early March 1994, eight highly enriched uranium (HEU) samples were collected from materials stored at the Ulba Metallurgical Plant in Oskamen (formerly Ust Kamenogorsk), Kazakhstan. While at the plant site, portions of four of these samples were dissolved and analyzed by mass spectrograph at the Ulba analytical laboratory by Ulba analysts. Three of these mass spectrograph solutions and the eight HEU samples were subsequently delivered to the Oak Ridge Y-12 Plant for complete chemical and isotopic analyses. The chemical forms of the eight samples were uranium metal chips, UO₂ powder, uranium/beryllium oxide powder, and uranium/beryllium alloy rods. All were declared, by the Ulba plant, to have a uranium assay of ~ 90 wt % ²³⁵U. The uranium/beryllium powder and alloy samples were also declared to range from about 8 to 28 wt % uranium.

The chemical and uranium isotopic analyses done at the Y-12 Plant confirm the Ulba plant declarations. In addition, all samples appear to have been enriched using some reprocessed uranium, probably from recovery of uranium from plutonium production reactors. As a result, all samples contain some ²³⁶U and ²³²U and have small, but measurable quantities of plutonium. This plutonium could be the result of either contamination carried over from the enrichment process or cross-contamination from weapons material. It is not the result of direct reactor exposure. Neither the ²³²U nor the plutonium concentrations are sufficiently high to provide a significant industrial health hazard. Both are well within established or proposed acceptance criteria for storage at the Y-12 Plant.

The trace metal analyses showed that, with the exception of beryllium, there are no trace metals in any of these HEU samples that pose a significant health hazard.

BACKGROUND AND OBJECTIVES OF ANALYSIS

Eleven uranium-containing samples collected in Kazakhstan arrived at the Y-12 Plant in early April 1994. Eight of the samples were in solid form and three were solutions. All were contained in small glass vials ~ 1 cm in diameter and 4 cm long. The total uranium content was less than 15 grams. Visual examination of the eight solid samples showed that one was apparently metal chips or filings, four were powders, and three were small chunks of metal. All had been exposed to air and all exterior surfaces were fully oxidized. The analyses requested were:

- 1. complete uranium isotopics analyses, including ²³²U;
- 2. determination of uranium content;
- 3. transuranic alpha activity (primarily plutonium); and
- 4. complete trace metal analyses.

The objective of this sample program was to completely characterize the material and to determine the suitability for eventual storage of ton quantities of similar material in Y-12's HEU storage vaults.

DISCUSSION OF RESULTS

The complete uranium isotopics analyses, the uranium content, and the transuranic alpha activity, are reported in Table 1. The trace metal analyses for all samples except one are reported in Table 2.

The uranium isotopics, excluding ²³²U, were done using thermal ionization mass spectrometry. The ²³²U concentration was measured by counting the characteristic decay alpha spectrum from both ²³²U and ²³⁴U and computing the ²³²U concentration by ratio with the measured ²³⁴U concentration. The determination of the weight percent uranium in the sample was done primarily by isotope dilution using ²³³U as the spike. Some were done by X-ray analysis using internal standards. The transuranic alpha activity, primarily plutonium, was measured by counting the characteristic alpha spectrum from ²³⁹Pu and ²³⁸Pu. The trace metal analyses were done using spark source mass spectrometry. Specific approved procedures, which are available, were followed for each of these analyses.

Uranium Isotopic Measurements

The average 235 U assay of all samples was 89.62 ± 0.6 wt %. This agrees with both the Kazakhstan declarations (i.e., within ± 0.7 wt %) and with the mass spectrographic analysis (i.e., within ± 0.06 wt %) done on four of the samples on the day of the collections.

All samples had significant quantities of ²³⁶U, the range being from 0.255 to 0.497 wt %. This amount of ²³⁶U in the HEU is consistent with enrichment using reprocessed uranium from plutonium production reactors as a portion of the feed. The ²³⁴U content of the HEU samples ranged from 0.813 to 1.122 wt %. This level is consistent with the enrichment of near natural uranium as the feed material. These concentrations, for both ²³⁴U and ²³⁶U, are well within the range of HEU stored at Y-12 and that used in U.S. nuclear weapons.

The 232 U concentration in the samples ranged from 0 to 0.83 ppb in the sample. The current allowable 232 U limit in ASTM specification C-787-90 is 0.01 ppb in commercial UF₆ made entirely from natural uranium. In enriched UF₆ made from reprocessed uranium, the allowable limit is raised to 5 ppb. The analysis of the Ulba samples is well within this reprocessed uranium standard.

				Table 1			
Y-12	Analys	es of HEU	Sample	s from	Ulba M	etallurgical l	Plant
			Y-12	Y-12	Y-12	Ulba	Ulba MS
	Original	Material	U-234	U-235	U-236	U-235*	U-235**
Sample No.	Can No.	Туре	wt %	wt %	wt %	wt %	wt %
79776	140	U metal chips	0.967	89.997	0.497	90.01	90.06
78777	588	UO2 powder	1.014	89.808	0.437	89.84	89.86
78778	140	solution	0.966	90.021	0.497	90.01	
79779	116	solution	1.122	89.057	0.298	89	
79780	129	solution	0.813	89.52	0.266	89	
79781	63	U/Be powder	1.063	89.555	0.401	89	
78782	129	U/Be powder	0.812	89.562	0.265	89	89.58
79783	116	U/Be rods	1.122	89.064	0.297	89	89.12
79784	12	U/Be powder	0.928	90.045	0.484	90.01	
79785	455	U/Be rods	0.956	89.714	0.282	89	
79786	70	U/Be rods	0.999	89.439	0.255	89	
						* Provided by Ulbs in analyses	** These mass spectrogrphi
	 					list for all cans containing	analyses done at Ulba on the
	<u> </u>					HEU.	day that the samples were
	<u> </u>						collected
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				Table 1 (cont.)		• • • • • • • • • • • • • • • • • • •
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	Y-12	Ulba				- Annahariti sati ka sa kaki an a ay a ama yamandanga 1908 da saka kabibbik
- down	wt %	wt %	pCi/gm of sample	pCi/gm of sample	Bq/gm of sample*	Bq/gm of sample*
Sample No.	Uranium	Uranium*	Pu-238	Pu-239	Pu-238	Pu-239
79776	90.81(see note)	99.94	976.20	43.34	36.12	1.60
78777	86.98	87.49	18.40	171.10	0.68	6.33
78778	na	na	107.70	3.72	3.98	0.14
79779	na	na	0.37	27.17	0.01	1.01
79780	na	na	0.02	1.35	0.00	0.05
79781	26.25	18.3	149.20	14050.00	5.52	519.85
78782	8.55	8.67	3.51	242.70	0.13	8.98
79783	19.35	19.83	60.23	4449.00	2.23	164.61
79784	20.25	21.7	1.09	7.60	0.04	0.28
79785	8.85	9.02	5,46	121.40	0.20	4.49
79786	26.29	27.65	129.60	10440.00	4.80	386.28
	Note In the process of	• Provided by Ulba in			* Defined as (pCi per gm of sample)	* Defined as (pCi per gm of sample)
	collecting the sample from the	analyses list for all cans			times (0.037 Bq/pCi).	times (0 037 Bq/pCi)
	original metal slug, the chips	containing HEU				Control of the second of the s
	were partially oxidized by the		A =0.		** *	
	air. As a result the sample		··· ··· - · · · · · · · · · · · · · · ·		THE PERSON NAMED IN COLUMN TWO IS ADDRESS OF THE PERSON OF	Mariana (1986-1988 M. Herry P. 198-16.2 Magnet P.S. Calabathan
	analyzed at Y-12 looks more				THE PARTY AND A STATE OF THE COLUMN ASSESSMENT	
	like oxide than metal				v value (1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	AT TO THE ENGLISH WHEN THE
		A	# 11 <u>-11 -11 -</u> , 114/1 - 1		t commence and the commence of	-
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			Table 1 (cont.)		
			Micrograms of Pu-238	Micrograms of Pu-239	Total
	Bq/gm of sample,	Pu	per gram of sample	per gram of sample	micrograms of Pu
Sample No.	Total plutonium	dpm/gm	(see note 1)	(see note 2)	<u>per gm of sample</u>
79776	37.72	2263.38	5.71E-05	6.99E-04	7.56E-04
78777	7.01	420.69	1.08E-06	2.76E-03	2.76E-03
78778	4.12	247.35	6.30E-06	6.00E-05	6.63E-05
79779	1.02	61.14	2.17E-08	4.38E-04	4.38E-04
79780	0.05	3.03	1.07E-09	2.17E-05	2.17E-05
79781	525.37	31522.22	8.73E-06	2.27E-01	2.27E-01
78782	9.11	546.59	2.05E-07	3.91E-03	3.91E-03
79783	166.84	10010.49	3.52E-06	7.18E-02	7.18E-02
79784	0.32	19.29	6.37E-08	1.23E-04	1.23E-04
79785	4.69	281.63	3.19E-07	1.96E-03	1.96E-03
79786	391.08	23464.51	7.58E-06	1.68E-01	1.68E-01
			Note 1: Defined as: (pCi per gm of sample)/	Note 2: Defined as (pCi per gm of sample)/	The second section of the second section is a second section of the second section in the second section is a second section of the second section in the second section is a second section of the second section in the second section is a second section of the second section in the second section is a second section of the second section in the second section is a second section of the second section is a second section of the second section is a second section in the second section is a second section of the second section is a second section of the second section is a second section in the section is a section in the section is a section in the section is a section in the section in the section is a section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section is a section in the section in the section in the section in the section is a section in the section in the section in the section is a section in the section in t
			(1.71x10E7 pCi per microgram)	(6.2x10E4_pCl per microgram)	
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		Table 1 (cont.)	
	Total Uranium alpha,	% Plutonium alpha in	U-232 concentration,
Sample No.	dpm/gr of sample	transuranics**	ppb U-232 in sample*
79776	1.264E+08	0.00179	0.83
78777	1.269E+08	0.00033	0.37
78778	1.943E+07	0.00127	Insufficient Sample
79779	2.070E+05	0.02953	0
79780	4.583E+04	0.00662	5.8E-05
79781	3.999E+07	0.07876	0.2
78782	1.334E+07	0.00410	0.051
79783	3.103E+07	0.03225	Insufficient Sample
79784	2.710E+07	0.00007	0.16
79785	1.093E+07	0.00258	Insufficient Sample
79786	3.767E+07	0.06225	Insufficient Sample
		** Y-12 has proposed specifications for	* The current U-232 allowable limit in ASTM
		recycle material that may go into	specification C-787-90 is 0.01 ppb in Commercial
		effect soon. They are based on DOE	UF6 from natural uranium and is 5 ppb in
		Order 5480 11 (II.A 1) and Federal	UF6 from reprocessed uranium All of the
		Guidance Report No. 11, 1988	above samples meet the reprocessed
		The proposed specification is that the	uranium specification.
		ratio of total plutonium to total uranium	Y-12 is storing some HEU having up to 40
		activity (as Ci/g of uranium) be less than	pprn U-232.
		or equal to 0.0075 or 0.75%.	
	and the second section of the section o	All of the above samples are well below	
		that recommendation.	
		Y-12 is storing some HEU having	
		plutonium concentrations up to 1 ppm	
		Based on nominal compositions for	
		weapns grade plutonium and US HEU the	
		activity ratios would be 0 0082 or 0.82%	

Table 2

Trace Metal Analyses of HEU Samples From the Ulba Metallurgical Plant

Parts per Million (ppm) of Sample

	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.	Sample No.
Element	79776	79777	79 77 8	79 779	79780	79781	79782	79783	79784	79785	79788
śe		7.00	1.00	> 1000	> 1000	> 0.5 %	> 0.5 %	> 1000	> 1000	> 1000	>1000
B	This sample	0.04	0.08	0.30	0.30	0.04	0.04	0.50	0.06	2.00 0.07	2.00 0.40
Na Na	too smeli for data	0.50	0.20	0.10 5.00	1.00	9.00	0. 50	1.00	0.80	0.01	- 040
Mg	anelyses.	150.00	1.00	14.00	1.00	6.00	2.00	94.00	26.00	190.00	29.00
Al		6.00	1.00	0.05	0.20	18.00	72.00	3.00	10.00	1.00	5.00
Si	L	130.00	3.00	1.00	1.00	740.00	60.00	11.00	64.00	2.00	22.00
P		3.00	6.00	0.09	0.09	2.00	4.00	2.00	3.00	0.30	0.90
S Ci		25.00 16.00	2.00	46.00	15.00 4.00	68.00 16.00	20.00 19.00	46.00 7.00	9.00	30.00	31.00 0.90
K	······································	30.00	0.60	3.00	5.00	4.00	4.00		1.00		
Ca		29.00	5.00	€ 820	E 380	370.00	140.00	160.00	23.00	> 1000	49 00
Sc		0.30	0.20	0.08	0.08	0.40	0.40	0.05	0.50	0.05	0.20
		0.10	0.08	0.40	0.04	22.00	3.00	0.40	0.20	0.03	0.08
Cr		4 00	0.08 3.00	0.03	0.03 2.00	0.50 560.00	0.10 25.00	0.02 4.00	0.20 22.00	0.02 2.00	0.06 2.00
Mn		— 1 00	0.70	0.40	1.00	49.00	52.00	2.00	7.00	11.00	3.00
Fe		85.00	7.00	5.00	12.00	690.00	230.00	130.00	110.00	E 1500	100.00
co		0.20	0.20	0.04	0.04	2.00	0.20	9.00	0.20	76.00	7.00
Ni Cu		18.00 7.00	1.00	2.00	3.00	92.00	8.00	340.00	10.00	> 1000	340.00
Zn		11.00	1.00	2.00 3.00	2.00 8.00	8.00 6.00	10.00	140.00 9.00	11.00 5.00	>1000 14.00	170.00 2.00
Ga		0.20	0.10	0.20	0.07	0.30	0.30	0.04	0.40	0.04	0.10
Ge		0.40	0.20	0.10	0.10	0.50	0.30	0.08	0.70	0.08	0.20
As		0.20	0.09	0.04	0.04	0.20	0.20	0.40	0.30	1.00	0.10
Se Br		0.30	0.20	0.00	0.09	0.40	0.40	0.06	0.60	0.06	0.20
Rb		0.70	0.20 1.00	0.09	0.20	0.40	0.40	0.06	0.80 1.00	0.06 0.10	0.20
Sr		0.20	0.10	0.06	0.06	0.30	0.30	0.04	0.40	0.04	0.10
Y		0.20	0.10	0.05	0.05	0.20	0.20	0.03	0.30	0.03	0.10
<u>Zr</u>		0.40	0.20	0.10	0.40	2.00	2.00	0.09	0.60	0.20	0.20
Nb Mo		0.20	0.10	0.06	0.07	0.30	1.00	0.04	0.30	0.04	0.10
Ru		0.90 0.70	0.50	0.20	0.80	36.00 0.90	1.00 0.90	0.20	1.00	0.20	0.50
Rh		0.20	0.10	0.08	0.06	0.30	0.30	0.10	0.40	0.04	0.10
Pd		0.80	0.50	0.20	0.20	1 00	1.00	0.20	1.00	0.20	0.50
Ag		0.50	0.80	0.10	0.10	0.60	0.60	0 40	0.70	0.20	0.30
Cd In		0.80	0.50	0.20	0.20	1.00	1.00	0.20	1.00	0.20	0.50
Sn		2.00	0.10	0.07	0.50	2.00	2.00	0.05	3.00	0. 05	0.10
Sb		0.50	0.30	0.10	0.10	0.80	0.80	0.08	0.80	0.08	0.30
Te		0.80	0.40	0.20	0.20	1 00	1.00	0.10	1 00	0.10	0.40
		0.30	0.20	0.08	0.08	0.30	0.30	0.05	0.50	0.05	0.20
Cs Ba		0.30	0.20	0.08	0.08	0 40	0.40	0.05	0.50	0.05	0.20
<u> </u>		0.30	0.20	0.10 0.08	0.10	0.50	0.50	0.07	0.70	0.07	0.20 0.20
Co		0.30	0.20	0.09	0.09	0.40	0.40	0.06	0.60	0.06	0.20
Pr		0.30	0.20	0.08	0.08	0.40	0.40	0.05	0.30	0.05	0.20
- Nd Sm		1.00	0.70	0.40	0 40	2.00	2.00	0.20	2.00	0.20	0.70
Eu		0.60	0.70	0.30 0.20	0.30	2.00 0.80	2.00 0.80	0.20	1.00	0.20	0.70
Gd		2.00	0.90	0.20	0.20	2.00	2.00	0.30	3.00	0.10	0.30
ТЪ		0.30	0.20	0.10	0 10	0 40	0.40	0.06	0.60	0.06	0.20
Dy		1.00	0.70	0.30	0.30	2.00	2.00	0.20	2.00	0.20	0.79
Ho Er		1.00	0.20	0.10	0.10	2.00	1.00	0.06	2.00	0.08	0.20
Tm		0.40	0.20	0.10	0.10	0.40	0.50	0.20	0.60	0.20	0.20
Yb		1.00	0.60	0.30	0.30	1.00	1.00	0.20	2.00	0.20	0.60
Lu		0.40	0.20	0.10	0.10	0.50	0.50	0.07	0.60	0.07	0.20
Hf Ta		1.00	0.80	0 40	0 40	2.00	2.00	0.30	2.00	0.30	0.80
- 'a		1.00	2.00 0.70	0.30	0.30	1.00	2.00	0.70	6.00	0.70	1.00 .
Re		0.60	0.70	0.20	0.40	2.00 0.80	2.00 0.80	0. 20 0. 10	2.00 1.00	0. 20 0. 10	0.70
Os		1.00	0.60	0.30	0.30	2.00	1.00	0.20	2.00	0.20	0.60
lr .		0.70	0.40	0.20	0.20	2.00	0.80	0.10	1 00	0 10	0 40
Pt Au		1.00	0.70	0.30	0.30	2.00	2.00	0.20	2.00	0.20	0.70
Au Hg		1.00	0.70	0.40	0.40	0.80	2.00	0.20	2.00	0.20	0.70
T)		0.60	0.80	0.40	0.40	1.00	2.00	0.30	2.00	0.30	0.80
Pb		0.80	0.50	0.20	0.20	0. 80 1.00	0.80 1.00	0.10	1.00	0.10 0.20	0.30 0.50
8-		0.40	0.20	0.10	0.10	0.60	0.60	0.08	0.70	0.08	0.20
Th		0.30	0.30	0.10	0.10	0.60	0.80	0.09	0.80	0.09	0.30
											

Currently, Y-12 does not have an official ²³²U acceptance standard. However, HEU containing up to 40 ppb ²³²U has been accepted and is currently being stored. Some studies have been made in which acceptance of HEU containing up to 100 ppb ²³²U was considered. However, no firm decision was made in the studies.

Plutonium Contamination

Alpha counting for plutonium isotopes revealed the presence of both ²³⁹Pu/²⁴⁰ Pu and ²³⁸Pu in all of the samples. (The activity reported for ²³⁹Pu is the sum of that for ²³⁹ Pu and ²⁴⁰Pu since the alpha energies for the two isotopes are quite close.) Although plutonium is present on the samples, the quantities are all well within Y-12 proposed specifications for material acceptance criteria. The proposed specification is that the ratio of total plutonium alpha to total uranium alpha activity is less than or equal to 0.0075 or 0.75%. Table 1 shows that these samples are at least a factor of 10 less than the proposed specification.

Y-12 is currently storing some uranium having plutonium concentrations up to 1 ppm. Based on nominal compositions for weapons grade plutonium and nominal U.S. weapons HEU composition, the activity ratio for this material would be about 0.0082 or 0.82%.

Based on the ratio of measured alpha activity (and the derived atom ratios) for the ²³⁸ Pu and ²³⁹Pu sources, there appears to be at least two different sources for the plutonium contamination in these samples. The high ²³⁸Pu contamination in the uranium metal samples may indicate that this material may have been used for a different application than the uranium/beryllium fuel rods. The ²³⁸Pu atom ratio (0.082 to 0.11) in the plutonium appears to be too low to indicate a ²³⁸Pu heat source application (which would be about 0.80), but is too high to be from either a normal light-water reactor or a liquid metal fast breeder reactor (which would be about 0.01 to 0.04). However, some preliminary calculations indicate that the irradiation of either HEU or low-enriched uranium (LEU) containing small amounts of ²³⁷Np (for HEU, ²³⁷Np concentrations of 0.015 to 0.025 atom %, and for LEU concentrations of 0.2 to 0.3 atom %) can make plutonium having ²³⁸Pu in the observed concentration range. There is no information indicating why this contamination should be present in the HEU metal sample. In the remaining samples, the plutonium contamination and the ²³⁸Pu concentration could be attributed to cross-contamination of the HEU with weapons grade plutonium either in the weapon or during dismantlement/disassembly operations.

Trace Metal Analyses

The trace metal analyses done for this project are summarized in Table 2. First, note that with the exception of those element concentrations labeled '> xxxx', all other concentrations are maximum values and they could very well be less than the reported values.

As expected, all the samples showed high concentrations of beryllium. Otherwise, all the high concentrations noted were in the standard common elements Mg, Si, S, Cl, K, Ca, Cr, Mn, Fe, Ni, and Cu. Even in the concentrations noted, none of these pose a health hazard. All the samples had significant quantities of Zn, the maximum concentration being 14 ppm. The remaining elements, with the exception of Mo in Sample No. 79781, are generally below 2 ppm, well below any potential health hazard.

With the exception of beryllium, there are no trace metals in these HEU samples that are of any significant health hazard.

ACKNOWLEDGMENTS

I would like to acknowledge the good cooperation and prompt analysis of these samples by the Quality Service Division of the Y-12 Analytical Laboratories. Especially helpful were J. H. Hamilton, E. E. Dukes, and J. B. Wilson.

DISTRIBUTION

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12-16. U.S. Department of Energy

Oak Ridge Operations

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Enclosure 1

Letter: Bostock to Spence

Dated May 15, 1995

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"Project Sapphire Data Summary in Response to DOE/ORO-DOE/HQ Teleconference on December 21, 1994"

The following data are provided per agreement with DOE/ORO and DOE/HQ to provide further information concerning the material and nuclear properties of the Project Sapphire material that is now safely stored at the Y-12 Plant in Oak Ridge, Tennessee. These data provide detailed analyses of samples evaluated at the Y-12 Plant Laboratory (see Enclosure 2), a summary of Health Physics measurements of the DOT 6M-2R containers and some of the inner stainless steel containers, a description of the nondestructive assay (NDA) measurement techniques used to evaluate the U-235 assay and uranium content of a majority of the stainless steel cans, and finally hydrogen analysis data.

Enclosure 2 describes the analytical results of 11 samples (representing eight different Kazakhstani containers) of Project Sapphire material that were analyzed at the Y-12 Plant from April to July 1994. These samples provided the initial data on the following material forms: uranium metal, UO₂ powder, uranium-beryllium oxide powder (machining scrap), and uranium-beryllium alloy

Summary of Health Physics Data

The following information was extracted from the Health Physics Container Survey records for each DOT 6M-2R container. These records specifically list the readings taken for Alpha and Beta/Gamma surface contamination, as well as Beta/Gamma and Neutron dose rates at the container surfaces and at a distance of 1 meter. Due to the urgency to provide this information to DOE/HQ on a timely basis it is being provided in summary form. In summary, the data are as follows:

Alpha (dpm/100 cm²)

Max. Fixed Plus Removable - < 250 (Removable) Low - 0 (Removable) High - 24

Beta/Gamma (dpm/100 cm²) (Removable) - < 120

Beta/Gamma (mR/hr)

(at contact) Low - 0.0 (at contact) High - 0.7

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(at 1 meter) Low - 0.0 (at 1 meter) High - 0.2

Neutron (mrem/hr)

(at contact) Low - 0

(at contact) High - 1.0

(at 1 meter) Low - 0

(at 1 meter) High - 0.5

Radiation measurements were also taken of some of the stainless steel cans of repackaged Kazakhstani material prior to insertion into DOT 6M-2R containers to establish the dose rates for the various material forms. These dose rates are summarized below:

Maximum Dose Rates
8 mrem/hr
4-6 mrem/hr
3.5 mrem/hr (neutron), 3.5 mrem/hr (gamma)
4.8 mrem/hr (neutron), 4.2 mrem/hr (gamma)
No measurements recorded
No measurements recorded
No measurements recorded

Summary of Nondestructive Assay Measurement Techniques

Two types of NDA assay measurements were performed in Kazakhstan on the materials after they were repackaged into stainless steel, crimp sealed cans: (1) a neutron counting technique and (2) a gamma spectroscopy technique using a Davidson Multichannel Analyzer. The data from the NDA measurements are not tabulated in this document since the results either indicated the mass of uranium per can or the uranium-235 enrichment level for each can and these data would fill dozens of pages and be of limited value for a request for proposals. However, a brief description of each NDA technique used is provided below:

Sodium Iodide (NaI) detector to confirm enrichment. Confirmatory measurements ranged from 80-86% and declared values were on the

order of 89%.

Uranium Oxide Measured with Davidson Portable Multichannel Analyzer (PMCA) and

Sodium Iodide (NaI) detector to confirm enrichment. Confirmatory measurements ranged from 83-86% and declared values were on the

order of 89%.

Uranium-Beryllium Alloy Rods

Uranium-234 is enriched proportionally with Uranium-235. Uranium-234 is a strong alpha emitter, which in the presence of beryllium leads to an alpha-n reaction and an increased neutron flux that can be measured with a SNAP detector. The weight and physical dimensions of several different weight percent rod batches were measured to determine the density of the batches. The density correlates to within about 1% of the uranium-beryllium (U-Be) ratio. Their declared enrichment was assumed to be correct, the U-Be ratio was confirmed as described above, and several "standards" were made by filling one of our cans one-quarter full and measuring it with the SNAP detector, then one-half full and measuring, and then completely full and measuring it, This was repeated over the range of different U-Be ratios. The grams of uranium proved to be proportional to the net neutrons counted by the SNAP. The slope of the line was essentially the same for all U-Be ratios with a corresponding shift in the intercept. The method is capable of measuring the grams of uranium in a can to within 10%, relative.

Uranium Oxide-Beryllium Oxide Rods

Measured with a Davidson Portable Multichannel Analyzer (PMCA) and sodium iodide (NaI) using a previously calibrated point source measurement.

U-Be Residue

The SNAP detector was used to measure the neutrons emitted in the same manner as above, utilizing the same linear equation it can confirm the uranium content of a can to within +/- 10%, relative.

HEU Contaminated Graphite

Measured with a Davidson Portable Multichannel Analyzer (PMCA) and Sodium Iodide (NaI) using a previously calibrated point source measurement.

Laboratory Salvage

The SNAP detector was used to measure the neutrons emitted in the same manner as above.

Hydrogen Analysis

The Sapphire materials were analyzed for hydrogen content on a statistical basis to comply with the regulatory requirements of the U.S. Department of Transportation's Competent Authority Certification for the Type B Fissile Radioactive Materials Package Certificate USA/0002/X (Note: This container is also called the DOT 6M-2R). This certificate requires the shipper to certify that the hydrogen to uranium-235 ratio is less than or equal to three for any

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uranium compounds. Therefore, this applied to all the materials, except the HEU metal. Samples of the uranium compounds, listed previously, were taken on a statistical basis from the Kazakhstani containers and analyzed for hydrogen content using a laboratory induction furnace and hydrogen trapping technique. Only a few cans of material were packaged with an H/X ratio greater than three and these cans were shipped with other cans that had H/X substantially less than three, so the total H/X ratio for any 6M-2R container never exceeded the limit.