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GEN-1270

U.S. Nuclear Regulatory Commission
Medical, Academic, and Commercial Use
Safety Branch (Mail Stop OWEN-6H3)
ATTN: Mr. Michael Lamastra
Washington, D.C. 20555

Subject: Control No. 019643; Submittal of an Expanded Appendix 5.0 to General Atomics' Application for License to Distribute Irradiated Topaz

Reference: Keith E. Asmussen letter GEN-1266 to U.S. NRC Medical, Academic, and Commercial Use Safety Branch, ATTN: Mr. Michael Lamastra, dated September 9, 1988

Dear Mr. Lamastra:

As you know, General Atomics (GA) recently submitted the referenced application for a license authorizing the distribution of irradiated topaz to persons exempt from regulation. While we were finalizing that submittal, we were also nearing the completion of a test run of our system for assaying large samples of gemstones. This system was described in Appendix 5.0 to our application. Important results from this test run have just become available.

In order to make the information contained in our application as complete as possible, we have expanded Appendix 5.0 to include a summary and discussion of these results. The Appendix 5.0 submitted with our referenced application should be replaced by the complete expanded version which is enclosed.

We apologize for the inconvenience caused by this late change to Appendix 5.0.

If you should have any questions or need any additional information, please do not hesitate to contact me at (619) 455-2823 or Drs. Whittemore or Razvi at (619) 455-3277.

Very truly yours,

Keith E. Asmussen

Keith E. Asmussen, Manager
Licensing, Safety and
Nuclear Compliance

KEA/mtk
Enclosure

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app.

A/S

APPENDIX 5.0

Assay of Large Samples of Gemstones Using High Resolution Semiconductor Detector

The Ge(Li) [or equivalent HpGe] high resolution detector used in the determination of radionuclide concentrations in irradiated gemstones is mounted on a vertical cryostat. The available detectors have gamma sensitivities ranging between 5 and 20 percent. The diameters of the outer container will vary between 2.75 inches and 3.0 inches.

The design of a holder for the assay of a large number of gemstones in Step K1 (Figure 1 of Application) (eg., ~50 1-gram stones) is based on experimental measurements made with a specially assembled source containing approximately 0.08 μ Ci Cs-137 (662 keV) and 0.13 μ Ci Co-60 (1173, 1333 keV). The source holder for these measurements was a small polyethylene cap with inside diameter of about 8 mm and about one-half millimeter thickness between the source and the gemstone holder. A survey was made in four quadrants and at various heights to arrive at the curvature shown in Figure 1.

The relative sensitivity as a function of position is 44% of the sensitivity for a single stone placed in the usual location used for measurements on a single stone. The value (44%) is an average at each location of the sensitivities for the gamma rays from cesium and cobalt.

A test run for 44 gemstones weighing about 50 grams was made using the large sample holder. Tables 1 and 2 present the pertinent Ge(Li) data. In addition to the routine isotope callouts (Table 1) the spectroscopist provided results of his inspection and additional calculations that also identified weak Co-60 and Zn-65 components (Table 2). The absence of additional gamma lines for Eu-154 and Se-75 allowed the spectroscopist to line out these possibilities. A summary of the results was derived by the spectroscopist from Tables 1 and 2 and is given below:

Nuclide	nCi/g	10 CFR30.70 Limits (nCi/g)	Ratios
Ta-182	50×10^{-3}	0.40	0.125
Sc-46	2.7×10^{-3}	0.40	0.0068
Mn-54	3.5×10^{-3}	1.0	0.0035
Co-60	0.7×10^{-3}	0.50	0.0014
Zn-65	1.7×10^{-3}	1.0	0.0017
Cr-51	2.3×10^{-3}	0.09	0.026
Na-22	2.7×10^{-3}	(1.0)*	0.0027

*See discussion below.

$$\Sigma = 64 \times 10^{-3} \frac{\text{nCi}}{\text{g}}$$

$$\Sigma = 0.167 \ll 1.00$$

These data demonstrate that for this batch, the "sum of ratios" is less than unity, with a total specific activity of 0.064 nCi/g. It is also apparent that stones with this same distribution of radioactive components could have specific activity as large as 0.383 nCi/g and still satisfy the release criteria - i.e.,

$$\frac{1.00}{0.167} \times 0.064 = \underline{0.383 \text{ nCi/g}}$$

This maximum specific activity is called L_{max} as used in Equations 5 and 6 on page 27 of the Application.

The assay methodology using a large holder with approximately 50-gram samples of gemstones for the Ge(Li) count (Step K1) has improved the sensitivity of detection level for the minor radioactive components in the

stones. Simultaneously, this has increased the possibility of identifying isotopes (at these very low levels of concentrations) not previously found in assays of one-gram samples. Furthermore, this increases the likelihood of finding an occasional isotope not included in Schedule A of 10CFR30.70. The "catchall" level of 10^{-6} $\mu\text{Ci}/\text{gram}$ is considered inappropriate for this application because it is arbitrarily small. Consider the case of Na-22 which is an isotope not listed in Schedule A and found at a level of 2.7 picocurie/gram in the reported data. In the following we illustrate a suitable approach for handling this and similar cases that may arise in the future.

In the particular case of Na-22, it is omitted from Schedule A even though it can be produced by nuclear reactors. This is well known to workers experienced with sodium cooled reactors. Its threshold for $(n, 2n)$ production is about 12.5 Mev with the cross section as a function of neutron energy given in BNL-325 (1). The fraction of all fast neutrons from a reactor fueled with U-235 above about 13 Mev is 1.103×10^{-4} (2). Since the fast flux at the location of the gemstone irradiation is about 10^{12} nv, the portion of this fast flux suitable for this $(n, 2n)$ reaction is about

$$\phi (E > 13 \text{ Mev}) = 1.1 \times 10^8 \text{ n} \cdot$$

An activation of 2.7×10^{-3} nCi/grams corresponds to an activation of ~ 5.0 dps for the 50 gram sample of stones:

$$\text{Act} = (N\sigma) \phi \left[1 - e^{-0.693t/T} \right]^{1/2} \dots \dots \dots (1)$$

where Act = activation of Na-23 to produce Na-22,
 (N σ) is the number of atoms of Na-23 in the 50 gram sample of stones, and σ is the effective $(n, 2n)$ cross section,

$$\begin{aligned}\sigma &= 1.1 \times 10^8 \text{ nv, and} \\ t &= 4 \text{ weeks or } 0.077 \text{ yr.} \\ T_{1/2} &= 2.6 \text{ yr.}\end{aligned}$$

From equation (1) we see that

$$(N\sigma) = 2.24 \times 10^{-6} \text{ cm}^{-2}.$$

If the average cross section above 12.5 Mev were as small as 1 millibarn, then $N = 2.24 \times 10^{21}$ atoms of Na-23 (~ 90 milligrams). A more reasonable average cross section might be as large as 10 millibarns, for which the mass of Na-23 would be only 9 milligrams. To find such a small admixture of Na-23 (9-90 milligrams) in a 50 gram sample of stones cannot be considered unreasonable, although it is unusual from all past experience.

The question now arises on how to arrive at a reasonable 10CFR30.70 limit for the Na-22 isotope. As demonstrated above, Na-22 can be produced by reactors from Na-23 and therefore, should be listed in Schedule A. To arrive at a suitable limit for this application, we have compared its gamma and beta radiation with other isotopes present and which may have similar half lives and/or emissions. These are compared in the following listing:

Isotope	$T_{1/2}$	γ Energies (keV)	β_{max} (MeV)	30.70 Limits nCi/g
Ta-182	0.32 yr	59, 67, 1121 1189, 1221	0.522	0.40
Sb-125	2.8 yr	176, 428, 601, 636	0.612	1.0
Zn-65	243.8d	1116	β^+ 0.325	1.0
Sc-46	83.3d	889, 1120	0.357	0.4
Na-22	2.6 yr	511, 1275	β^+ 0.545	---

For the case of Na-22 we first note that it cannot be ingested or inhaled because it resides within the completely insoluble topaz crystal. Next, we note from the above listing that on the basis of half life, Na-22 most nearly matches Sb-125 with a release limit of 1.0. We also note that the 1275 keV line of Na-22 matches the 1116 keV line of Zn-65 which also has a release limit of 1.0. Further, the range for positrons with E_{max} of 545 keV that are emitted by Na-22 is very much the same as the betas from the far more prevalent isotope, Ta-182, as well as Sb-125. In other words, even the smallest stones are infinitely thick for the Na-22 positrons. Because of the similarities in the half lives and emitted radiations between Na-22 and Sb-125, we believe a reasonable release limit for this minor component can be taken as that for Sb-125, i.e., 1.0 nCi/g.

References:

1. Neutron Cross Sections, BNL-325, Vol. 1, Suppl. 2, Second Edition (May, 1964).
2. H. Etherington, Ed., Nuclear Engineering Handbook, First Edition (1958), p. 7-91.

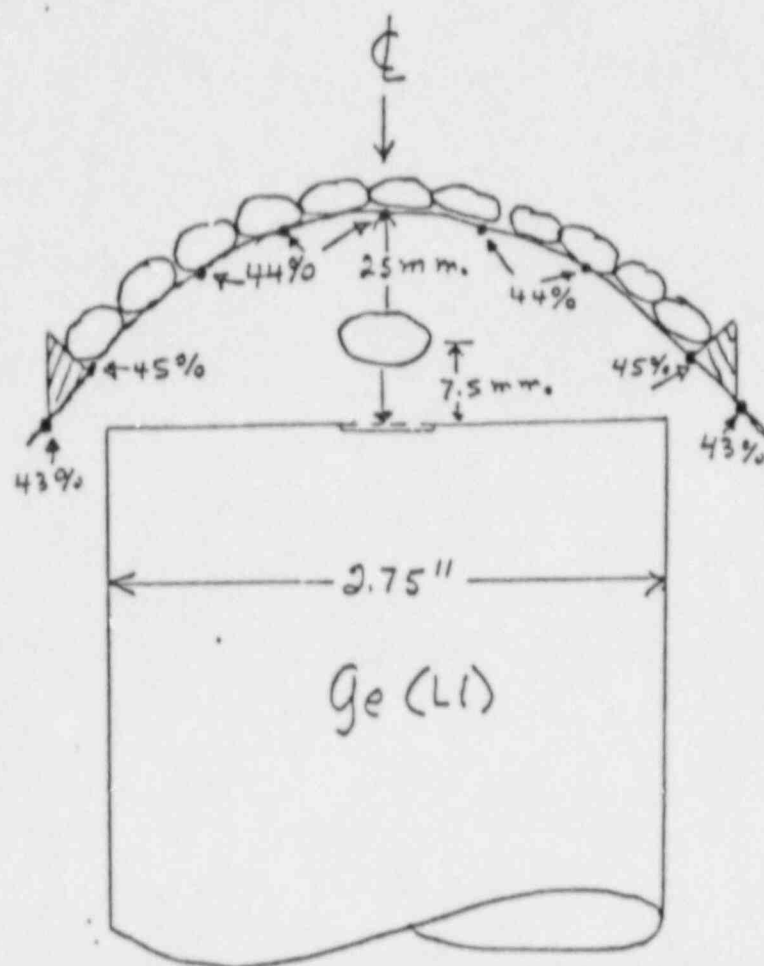


Fig. 1. Sketch roughly to scale showing dome shaped holder. This surface of revolution accommodates about 50 1 gram stones. For this holder the detector has about 44% of the sensitivity for a single stone placed at the usual (7.5 mm) position. Measured sensitivities are shown.

TABLE 1

----- GAMMA RAY SPECTRAL ANALYSIS -----
 PROGRAM VERSION: GA17-109 (4-22-88) DETECTOR: GE(LI)#2

SAMPLE: TRIGA STONES IN PLASTIC HOLDER

REQUESTOR'S NAME: M. WHITTENBOM
 SAMPLE GEOMETRY: STONE HOLDER
 SPECTRUM FILE: REQUEST NUMBER: SPEC 34801 1590

COUNT DATE: 9/ 3/88 START TIME: 8:41:14
 GAIN (KEV/CH): 0.3346937 INTERCEPT (KEV): -0.646
 LIVE TIME (MIN): 119.967 REAL TIME (MIN): 120.009

SENSITIVITY: 3.5 TOTAL CPM: 1060
 CALCULATION MODE: AI LIBRARY SEARCH: 4
 CHANNEL RANGE: 90 - 8192 # OF PEAKS FOUND: 23

----- PEAK INTEGRATION DATA -----

NUCLIDE	PEAK CHANNEL	ENERGY KEV	CPM	SIGMA %	HN KEY	TN/HN	ERROR KEY
TA-132	177	88.78	11.7	1.1	1	1	1
TA-132	200	97.41	11.7	1.1	1	1	1
TA-132	223	106.04	11.7	1.1	1	1	1
TA-132	246	114.67	11.7	1.1	1	1	1
TA-132	269	123.30	11.7	1.1	1	1	1
TA-132	292	131.93	11.7	1.1	1	1	1
TA-132	315	140.56	11.7	1.1	1	1	1
TA-132	338	149.19	11.7	1.1	1	1	1
TA-132	361	157.82	11.7	1.1	1	1	1
TA-132	384	166.45	11.7	1.1	1	1	1
TA-132	407	175.08	11.7	1.1	1	1	1
TA-132	430	183.71	11.7	1.1	1	1	1
TA-132	453	192.34	11.7	1.1	1	1	1
TA-132	476	200.97	11.7	1.1	1	1	1
TA-132	499	209.60	11.7	1.1	1	1	1
TA-132	522	218.23	11.7	1.1	1	1	1
TA-132	545	226.86	11.7	1.1	1	1	1
TA-132	568	235.49	11.7	1.1	1	1	1
TA-132	591	244.12	11.7	1.1	1	1	1
TA-132	614	252.75	11.7	1.1	1	1	1
TA-132	637	261.38	11.7	1.1	1	1	1
TA-132	660	270.01	11.7	1.1	1	1	1
TA-132	683	278.64	11.7	1.1	1	1	1
TA-132	706	287.27	11.7	1.1	1	1	1
TA-132	729	295.90	11.7	1.1	1	1	1
TA-132	752	304.53	11.7	1.1	1	1	1
TA-132	775	313.16	11.7	1.1	1	1	1
TA-132	798	321.79	11.7	1.1	1	1	1
TA-132	821	330.42	11.7	1.1	1	1	1
TA-132	844	339.05	11.7	1.1	1	1	1
TA-132	867	347.68	11.7	1.1	1	1	1
TA-132	890	356.31	11.7	1.1	1	1	1
TA-132	913	364.94	11.7	1.1	1	1	1
TA-132	936	373.57	11.7	1.1	1	1	1
TA-132	959	382.20	11.7	1.1	1	1	1
TA-132	982	390.83	11.7	1.1	1	1	1
TA-132	1005	399.46	11.7	1.1	1	1	1
TA-132	1028	408.09	11.7	1.1	1	1	1
TA-132	1051	416.72	11.7	1.1	1	1	1
TA-132	1074	425.35	11.7	1.1	1	1	1
TA-132	1097	433.98	11.7	1.1	1	1	1
TA-132	1120	442.61	11.7	1.1	1	1	1
TA-132	1143	451.24	11.7	1.1	1	1	1
TA-132	1166	459.87	11.7	1.1	1	1	1
TA-132	1189	468.50	11.7	1.1	1	1	1
TA-132	1212	477.13	11.7	1.1	1	1	1
TA-132	1235	485.76	11.7	1.1	1	1	1
TA-132	1258	494.39	11.7	1.1	1	1	1
TA-132	1281	503.02	11.7	1.1	1	1	1
TA-132	1304	511.65	11.7	1.1	1	1	1
TA-132	1327	520.28	11.7	1.1	1	1	1
TA-132	1350	528.91	11.7	1.1	1	1	1
TA-132	1373	537.54	11.7	1.1	1	1	1
TA-132	1396	546.17	11.7	1.1	1	1	1
TA-132	1419	554.80	11.7	1.1	1	1	1
TA-132	1442	563.43	11.7	1.1	1	1	1
TA-132	1465	572.06	11.7	1.1	1	1	1
TA-132	1488	580.69	11.7	1.1	1	1	1
TA-132	1511	589.32	11.7	1.1	1	1	1
TA-132	1534	597.95	11.7	1.1	1	1	1
TA-132	1557	606.58	11.7	1.1	1	1	1
TA-132	1580	615.21	11.7	1.1	1	1	1
TA-132	1603	623.84	11.7	1.1	1	1	1
TA-132	1626	632.47	11.7	1.1	1	1	1
TA-132	1649	641.10	11.7	1.1	1	1	1
TA-132	1672	649.73	11.7	1.1	1	1	1
TA-132	1695	658.36	11.7	1.1	1	1	1
TA-132	1718	666.99	11.7	1.1	1	1	1
TA-132	1741	675.62	11.7	1.1	1	1	1
TA-132	1764	684.25	11.7	1.1	1	1	1
TA-132	1787	692.88	11.7	1.1	1	1	1
TA-132	1810	701.51	11.7	1.1	1	1	1
TA-132	1833	710.14	11.7	1.1	1	1	1
TA-132	1856	718.77	11.7	1.1	1	1	1
TA-132	1879	727.40	11.7	1.1	1	1	1
TA-132	1902	736.03	11.7	1.1	1	1	1
TA-132	1925	744.66	11.7	1.1	1	1	1
TA-132	1948	753.29	11.7	1.1	1	1	1
TA-132	1971	761.92	11.7	1.1	1	1	1
TA-132	1994	770.55	11.7	1.1	1	1	1
TA-132	2017	779.18	11.7	1.1	1	1	1
TA-132	2040	787.81	11.7	1.1	1	1	1
TA-132	2063	796.44	11.7	1.1	1	1	1
TA-132	2086	805.07	11.7	1.1	1	1	1
TA-132	2109	813.70	11.7	1.1	1	1	1
TA-132	2132	822.33	11.7	1.1	1	1	1
TA-132	2155	830.96	11.7	1.1	1	1	1
TA-132	2178	839.59	11.7	1.1	1	1	1
TA-132	2201	848.22	11.7	1.1	1	1	1
TA-132	2224	856.85	11.7	1.1	1	1	1
TA-132	2247	865.48	11.7	1.1	1	1	1
TA-132	2270	874.11	11.7	1.1	1	1	1
TA-132	2293	882.74	11.7	1.1	1	1	1
TA-132	2316	891.37	11.7	1.1	1	1	1
TA-132	2339	899.00	11.7	1.1	1	1	1
TA-132	2362	907.63	11.7	1.1	1	1	1
TA-132	2385	916.26	11.7	1.1	1	1	1
TA-132	2408	924.89	11.7	1.1	1	1	1
TA-132	2431	933.52	11.7	1.1	1	1	1
TA-132	2454	942.15	11.7	1.1	1	1	1
TA-132	2477	950.78	11.7	1.1	1	1	1
TA-132	2500	959.41	11.7	1.1	1	1	1
TA-132	2523	968.04	11.7	1.1	1	1	1
TA-132	2546	976.67	11.7	1.1	1	1	1
TA-132	2569	985.30	11.7	1.1	1	1	1
TA-132	2592	993.93	11.7	1.1	1	1	1
TA-132	2615	1002.56	11.7	1.1	1	1	1
TA-132	2638	1011.19	11.7	1.1	1	1	1
TA-132	2661	1019.82	11.7	1.1	1	1	1
TA-132	2684	1028.45	11.7	1.1	1	1	1
TA-132	2707	1037.08	11.7	1.1	1	1	1
TA-132	2730	1045.71	11.7	1.1	1	1	1
TA-132	2753	1054.34	11.7	1.1	1	1	1
TA-132	2776	1062.97	11.7	1.1	1	1	1
TA-132	2799	1071.60	11.7	1.1	1	1	1
TA-132	2822	1080.23	11.7	1.1	1	1	1
TA-132	2845	1088.86	11.7	1.1	1	1	1
TA-132	2868	1097.49	11.7	1.1	1	1	1
TA-132	2891	1106.12	11.7	1.1	1	1	1
TA-132	2914	1114.75	11.7	1.1	1	1	1
TA-132	2937	1123.38	11.7	1.1	1	1	1
TA-132	2960	1132.01	11.7	1.1	1	1	1
TA-132	2983	1140.64	11.7	1.1	1	1	1
TA-132	3006	1149.27	11.7	1.1	1	1	1
TA-132	3029	1157.90	11.7	1.1	1	1	1
TA-132	3052	1166.53	11.7	1.1	1	1	1
TA-132	3075	1175.16	11.7	1.1	1	1	1
TA-132	3098	1183.79	11.7	1.1	1	1	1
TA-132	3121	1192.42	11.7	1.1	1	1	1
TA-132	3144	1201.05	11.7	1.1	1	1	1
TA-132	3167	1209.68	11.7	1.1	1	1	1
TA-132	3190	1218.31	11.7	1.1	1	1	1
TA-132	3213	1226.94	11.7	1.1	1	1	1
TA-132	3236	1235.57	11.7	1.1	1	1	1
TA-132	3259	1244.20	11.7	1.1	1	1	1
TA-132	3282	1252.83	11.7	1.1	1	1	1
TA-132	3305	1261.46	11.7	1.1	1	1	1
TA-132	3328	1270.09	11.7	1.1	1	1	1
TA-132	3351	1278.72	11.7	1.1	1	1	1
TA-132	3374	1287.35	11.7	1.1	1	1	1
TA-132	3397	1295.98	11.7	1.1	1	1	1
TA-132	3420	1304.61	11.7	1.1	1	1	1
TA-132	3443	1313.24	11.7	1.1	1	1	1
TA-132	3466	1321.87	11.7	1.1	1	1	1
TA-132	3489	1330.50	11.7	1.1	1	1	1
TA-132	3512	1339.13	11.7	1.1	1	1	1
TA-132	3535	1347.76	11.7	1.1	1	1	1
TA-132	3558	1356.39	11.7	1.1	1	1	1
TA-132	3581	1365.02	11.7	1.1	1	1	1
TA-132	3604	1373.65	11.7	1.1	1	1	1
TA-132	3627	1382.28	11.7	1.1	1	1	1
TA-132	3650	1390.91	11.7	1.1	1	1	1
TA-132	3673	1399.54	11.7	1.1	1	1	1
TA-132	3696	1408.17	11.7	1.1	1	1	1
TA-132	3719	1416.80	11.7	1.1	1	1	1
TA-132	3742	1425.43	11.7	1.1	1	1	1
TA-132	3765	1434.06	11.7	1.1	1	1	1
TA-132	3788	1442.69	11.7	1.1	1	1	1
TA-132	3811	1451.32	11.7	1.1	1	1	1
TA-132	3834	1459.95	11.7	1.1	1	1	1
TA-132	3857	1468.58	11.7	1.1	1	1	1
TA-132	3880	1477.21	11.7	1.1	1	1	1
TA-132	3903	1485.84	11.7	1.1	1	1	1
TA-132	3926	1494.47	11.7	1.1	1	1	1
TA-132	3949	1503.10	11.7	1.1	1	1	1
TA-132	3972	1511.73	11.7	1.1	1	1	1
TA-132	3995	1520.36	11.7	1.1	1	1	1
TA-132	4018	1528.99	11.7	1.1	1	1	1
TA-132	4041	1537					

TABLE 2

***** PEAK INTEGRATION DATA *****

NUCLIDE	PEAK CHANNEL	ENERGY KEV	CPM	SIGMA N	HN KEV	TN/MN	ERROR YEV
Zn-65	1112.00	1115.22	0.418	17.46	1.47	1.35	-0.10
Co-60	1173.00	1173.16	0.169	26.62	0.31	10.00	-0.05
Co-60	1132.00	1132.21	0.267	22.75	0.65	3.05	-0.24

***** DPM COMPUTATION DATA *****

LIBRARY FILE: AHS LONG EFFICIENCY FILE: EFF. STONES
 DIVISOR = 5.06000E 01 GRAM
 DAYS DECAYED = 15.830 8/22/68 12:00:00

NUCLIDE	ENERGY	HALF LIFE	MICROCI PER GRAM	2 SIGMA ERROR	DPM PER GRAM	2 SIGMA ERROR
Co-60	1173.16	5.27 Y	7.119E-07	2.94E-07	1.585E 00	6.54E-01
Co-60	1132.21	5.27 Y	4.118E-07	2.79E-07	1.362E 00	5.20E-01
Zn-65	1115.22	243.94 D	1.744E-06	4.09E-07	3.071E 00	1.15E 00

MATERIALS LICENSE

Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 40 and 70, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). This license shall be deemed to contain the conditions specified in Section 183 of the Atomic Energy Act of 1954, as amended, and is subject to all applicable rules, regulations and orders of the Nuclear Regulatory Commission now or hereafter in effect and to any conditions specified below.

Licensee		3. License number	04-14395-02E
1.	General Atomics P. O. Box 85608 San Diego, California 92138	4. Expiration date	October 31, 1993
2.		5. Docket or Reference No.	030-29179
6. Byproduct, source, and/or special nuclear material	7. Chemical and/or physical form	8. Maximum amount that licensee may possess at any one time under this license	
A. Any byproduct material with atomic numbers between 1-83	A. Irradiated processed topaz	A. Not applicable	
9. Authorized use			
A. In accordance with Section 32.11, 10 CFR Part 32, distribution of processed topaz containing byproduct material to persons exempt from licensing pursuant to Section 30.14, 10 CFR Part 30 or equivalent regulations of any Agreement State.			

CONDITIONS

10. Licensed material shall be distributed only from the licensee's facilities located at 10955 John Jay Hopkins Drive, San Diego, California.
11. Licensed material shall be distributed by, or under the supervision of Dr. Junaid Razvi or Dr. William Whittemore.
12. This license does not authorize possession or use of licensed material.
13. A. The licensee shall file periodic reports as specified in Section 32.12, 10 CFR Part 32.
B. The licensee shall file a report by November 1, 1989 and a new report at intervals not to exceed 14 months, listing the names and license numbers of all specific licensees to which the licensee has transferred irradiated gems pursuant to Section 30.41 of 10 CFR Part 30.

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