

Radioactive Effluents in Savannah River - Summary Report for 1991

by

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Radioactive Effluents in Savannah River
Summary Report for 1991 (U)

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July 16, 1992

**Radioactive Effluents in Savannah River
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Introduction

During 1991, low-level radiometric studies of the Savannah River continued to distinguish between effluent contributions from Plant Vogtle and the Savannah River Site. Measurements of these radioactive effluents are of mutual interest to both institutions, as they can address disturbing trends before they become health and legal concerns.

The Environmental Technology Section (ETS) has conducted radiometric studies of Plant Vogtle since late 1986, prior to its startup [ref 1]. The plant has two 1100 MWe pressurized water reactors developed by Westinghouse. Unit 1 started commercial operations in June 1987, and Unit 2 began in May 1989. During powered operations, ETS has routinely detected neutron-activated isotopes in controlled releases [refs 2-5] but all activities have been several orders of magnitude below the DOE guide values [ref 6]. In 1991, Vogtle effluents continue to have low activities in the river. The Vogtle release data and the ETS measurements have tracked well over the past five years.

ETS ultra low-level radiometric measurement techniques are used in this work. In general, river samples are concentrated on resins, prepared in the lab, and then counted in the Underground Counting Facility at 735-A [ref 7]. The bulk of the counting is by gamma spectrometry, but tritium is examined by liquid scintillation counting. The overall sampling/counting technique for gamma-ray analysis provides detection limits that are thousands of times lower than those routinely achievable. The tritium analysis also utilizes state-of-the-art detection sensitivity. An underwater NaI(Tl) at Hwy 301 Bridge provides continuous gamma monitoring of the Savannah River.

Summary

During 1991, the radioactive effluents in the Savannah River were somewhat less than those observed in 1990. This decreasing trend has followed Vogtle improvements in pre-processing their liquid effluents. These effluents continue to be dominated by ^{58}Co , which had a maximum concentration of only 0.21 pCi/L, which is an order of magnitude lower than the maximum observed in 1990. Many of the other man-made radionuclides observed in 1987-1990 [refs 2-5] have now decreased to where some are not even detected, and no new radionuclides were detected in the 1991 Vogtle effluents. Some ^{54}Mn and ^{60}Co were frequently observed, but traces of ^{95}Nb and ^{95}Zr were seen were observed only during the time of maximum ^{58}Co . No ^{51}Cr , ^{57}Co , or ^{59}Fe were seen in 1991. Tritium and ^{137}Cs were also monitored, but their levels generally remain consistent with known SRS sources. The maximum tritium observed near Vogtle was 10 pCi/mL; however, the maximum downstream tritium was an order of magnitude higher due to a tritium release from K-reactor in December [ref 8], which occurred after the tritium samples of the present study were retrieved.

River Sampling

Samples were collected from the Savannah River at Shell Bluff, near the Vogtle outfall, and at Hwy 301 bridge. The increase in activity from Shell Bluff to Hwy 301 Bridge is due to the effluents from both Vogtle and SRS. Activities measured just upstream and downstream of the Vogtle outfall identify contributions from that plant alone. Specifically, samples were taken 0.3 mi above, 0.1 mi below, and at the Vogtle outfall. A map of the sampling sites is given in the appendix.

Resin, water, and sediment samples were collected using methods detailed in the Vogtle pre-start study [ref 1]. Each resin concentrator sample consists of \approx 25 g of resin in a porous nylon bag. The samplers were in the river for two weeks, after which they were retrieved and returned to the lab. Water samples

were collected directly, and sediment samples were scooped into a specially designed sampler pulled along the bottom of the river.

Laboratory Measurements

The resin concentrator samples provide the most comprehensive isotopic information. Each of these samples was dried and ashed, leaving a smaller volume and thus better geometry for the HPGe detector. Typically, two HPGe detectors, with ^{60}Co standard efficiencies of 20% and 25%, were used in counting these samples; both are located in the Underground Counting Facility. Also, a larger detector with ^{60}Co standard efficiency of 90% and an active/pассив shield has been calibrated to examine these samples. Samples were counted overnight (or over the weekend) to assure good counting statistics for detecting low-level activities. The resulting HPGe gamma-ray spectra are analyzed with an IBM/PC-XT using the GRABGAM code [ref 7] to yield the activity (fCi) of each isotope detected in the resin sample. The average isotopic concentration (fCi/L) for the collection period is obtained by using empirical calibration data that relate the resin mass and collection time to the effective water volume (L).

The water samples were examined for tritium by liquid scintillation counting. A TRICARB™ LL 2050A low-level liquid scintillation analyzer was used. Vials with 3 mL of sample and 20 mL of OPTI-FLUOR™ scintillant were counted. Duplicate samples with 0.25 mL of tritium standard checked that the automatic quench corrections were being performed accurately.

Sediment samples were transferred to 1-L Marinelli beakers and then counted on the HPGe detectors as described above. The sample isotopic activities (fCi) from the gamma-ray analysis are divided by the dry weight (g) of each sample to yield the corresponding concentrations (fCi/g).

Underwater NaI(Tl) Measurements

The underwater NaI(Tl) detector operates from a stationary pontoon boat near Hwy 301 bridge. The detector includes a 9" diam by 4" long NaI(Tl) detector, four photo-multiplier tubes, and a high voltage/preamp unit - all contained in a waterproof stainless steel housing. The detector views its surroundings through a thin hemispherical dome.

The detector receives operating power via a waterproof cable connected to a \approx 25 V DC supply in the cabin of the pontoon boat. A unit within the detector assembly boosts this voltage to about 1000 V for photo-tube operation. The AC-coupled detector signals are transmitted via the same cable to the cabin, where they are

amplified and input to a multichannel analyzer.

Spectral data are collected on the multichannel analyzer, which comprises a COMPAQ 286 computer and an ACE MCA card from EG&G ORTEC. Spectra are stored on floppy disk in sequenced 24-hr periods for 2-3 weeks. The disks are retrieved and analyzed at the laboratory. This detector system provides better time resolution (1 day) than the resin concentrator method (2 weeks), but its sensitivity is lower by an order of magnitude [refs 3-5].

Results

The resin sampler analyses continued to indicate ^{58}Co , ^{60}Co , and ^{137}Cs , the most significant gamma-emitting radionuclides detected in the earlier studies [refs 2-5]. The 1991 results for these isotopes are given in Tables 1-3, along with plots in Figures 1-3. The other detected manmade gamma-emitting radionuclides are given in Table 4, where they are compared with the ^{58}Co levels.

Water analysis for tritium followed techniques used earlier [ref 1,2]. The 1991 tritium results are summarized in Table 5.

Sediments were analyzed for ^{58}Co , ^{60}Co , and ^{137}Cs to appraise whether any deposition is occurring during the transport of these isotopes in river water. The corresponding results are shown in Table 6.

Data for the underwater NaI(Tl) detector are presented in Figures 4 and 5. Figure 4 is an isometric plot of countrate vs gamma energy and date. Figure 5 compares the ^{58}Co detected by the underwater NaI(Tl) with that detected with the resin concentrator samples.

Discussion

In 1991, the decreasing trend of manmade radioactivity in the Savannah River continued. This was primarily due to continuing improvements in Plant Vogtle processing of effluent waste. River water samples show that Vogtle effluents contain fewer detectable nuclides, including ^{54}Mn , ^{58}Co , ^{60}Co , ^{95}Nb , and ^{95}Zr . No evidence of ^{51}Cr , ^{57}Co , or ^{59}Fe was observed in 1991, consistent with their overall decreasing trend from 1987 to 1990. Aside from these Vogtle contributions, the other activities were essentially consistent with levels of SRS radionuclides ^3H , ^{60}Co , and ^{137}Cs detected prior to Vogtle startup in 1987. The sediment samples complement the results observed for the river water. The underwater NaI(Tl) detector again provided continuous tracking of Vogtle effluents. As in the past, all measured radionuclide concentrations are well below DOE guides for drinking water.

The Vogtle effluents continue to be dominated by ^{58}Co , and effluent discharges from Vogtle track well with the temporal levels of ^{58}Co observed in the SRTC samples. However, in contrast to earlier observations, the river concentrations for ^{58}Co [Figure 1] do not track as well with those for ^{60}Co [Figure 2], although the ^{60}Co itself does track well with Vogtle release data for ^{60}Co . This is due to the variance in neutron activation intervals (now ranging up to 18 months) for the effluents released to the river, as activation-buildup of $^{58}\text{Co}(71\text{d})$ levels out much earlier than that for $^{60}\text{Co}(5\text{y})$. The data also show that the bulk of the ^{60}Co is from Plant Vogtle, as illustrated by ^{60}Co levels 0.3mi above and 0.1mi below the outfall [Table 2].

Relative to ^{58}Co , the river concentrations of other Vogtle-produced isotopes [Table 4] are generally lower than in earlier years [refs 2-5]. However, the relative concentrations of longer-lived $^{54}\text{Mn}(313\text{d})$ and $^{60}\text{Co}(5\text{y})$ increased in 1991, which is contrary to their decrease in 1990 [ref 5] that marked a departure from an increasing trend observed during 1987-1989 [ref 4]. This is probably due to (1) activation-buildup differences of the longer-lived ^{54}Mn and ^{60}Co relative to ^{58}Co holdup practices and (2) larger statistical ratio fluctuations due to the generally lower levels of all effluent Vogtle radionuclides. Indeed, the largest relative amounts of Mn-54 and Co-60 occurred for extremely low ^{58}Co . From 1987 to 1989 the relative concentrations of both ^{54}Mn and ^{60}Co had increased by more than an order of magnitude; during 1990 their concentrations dropped by about a factor of ~3; in 1991 they increased relative to ^{58}Co that were considerably lower.

During 1991, SRS still appeared to be the major source for the ^3H and ^{137}Cs concentrations in the river. Excluding the tritium release from K reactor in late December [ref 8], Table 5 indicates that the average ^3H at Hwy 301 Bridge was 4.1 pCi/mL, which is similar to the 3.0 pCi/mL measured prior to Vogtle startup. The maximum ^3H concentrations observed in this work (prior to the December tritium release) were 10 and 8 pCi/mL at the Vogtle outfall on March 3 and March 14, but these somewhat high levels were readily diluted to normal levels by the river. Concentrations ranging up to a maximum of 45 fCi/L ^{137}Cs were observed at Hwy 301 bridge, and these are consistent with an overall decreasing trend observed in recent years. The values near Vogtle are noticeably lower, averaging 5.2 ± 0.5 fCi/L and 5.8 ± 0.7 fCi/L for samples 0.3 mi above and 0.1 mi below the Vogtle outfall. This implies a ^{137}Cs contribution of 0.5 ± 0.9 fCi/L from Vogtle. By contrast, such ^{137}Cs contributions during 1987-1989 increased from 1 to 5 fCi/L, suggesting an increasing trend [ref 4]. The 1990-1991 data do not support such a trend. The future program will continue to monitor for this type of ^{137}Cs contribution.

The sediment samples contained detectable ^{58}Co , ^{60}Co , and ^{137}Cs [Table 6]. All activities observed in 1991 are comparable to the low values observed in 1990, except for the 32 fCi/g ^{58}Co measured 0.1 mile downstream of Vogtle on 2/7. Although this value is larger than its 1990 maximum of 10 fCi/g [ref 5], it is lower than the 1989 maximum of 98 fCi/g [ref 4]; thus, statistical fluctuations in sampling may explain this outlier. In general, the ^{137}Cs values are comparable to those observed in earlier years [refs 2-5].

The underwater NaI(Tl) detector at Hwy 301 Bridge continues to illustrate that Plant Vogtle effluents can be monitored by detection of the ^{58}Co activity. An isometric plot of countrate vs gamma energy and time is given in Figure 4, where the peak near 800 keV is identified with the 811 keV gamma of ^{58}Co . Figure 5 compares a time plot of this peak with a similar one for ^{58}Co concentrations measured with the resin samples. Although 1991 yielded lower ^{58}Co levels at Hwy 301 Bridge, the data are sufficient to show consistency for the two profiles, indicating continuation of the good tracking exhibited earlier [refs 3-5]. The 795 keV and 835 keV gammas of ^{228}Ac and the 835 keV of ^{54}Mn can also contribute to the peak near 800 keV in the spectra, due to resolution limits of the underwater NaI(Tl) detector [ref 4]. No such resolution problem existed for the resin samples because they were counted on HPGe detectors. Thus in Figure 5, the underwater NaI(Tl) plot of Co-58 includes a background component whereas the corresponding plot for HPGe-counted resins does not.

Table 7 compares the maximum-detected river concentrations with the DOE guide limits for drinking water [refs 6,8]. All concentrations are well below the guide limits. Furthermore, the table illustrates that maximum concentrations in 1991 were generally lower than those during 1987-1990.

Continuation of Study

These studies have continuously monitored the Savannah River since their inception in 1986. They will continue on a routine basis, as they provide early detection/correction of disturbing trends before actual health and legal concerns evolve. The program is mutually beneficial to both Vogtle and SRS, and cooperative efforts between the two sites continue to enhance the measurements program.

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Table 1. Co-58 Concentrations in 1991

(Values in fCi/L)^a

Date	Plot X	Shell Bluff fCi/L	Above Vogtle fCi/L	Below Vogtle fCi/L	H301 Bridge fCi/L
01/03-01/17/91		-5.91	-4.62	112.68	5.72
01/17-01/31/91	01/24	-3.31	-4.57	80.79	3.62
01/31-02/14/91		-7.56	-5.06	127.40	16.49
02/14-02/28/91	02/21	-5.50	-10.05	30.23	-3.44
02/28-03/14/91		-13.21	-3.80	32.11	-4.20
03/14-03/27/91	03/21	-2.42	-7.42	154.83	-1.65
03/27-04/10/91		missing	missing	81.33	4.92
04/10-04/24/91	04/17	-5.59	-4.90	21.40	-4.76
04/24-05/08/91		-8.09	-6.63	23.39	-5.24
05/08-05/22/91	05/15	-5.49	-6.57	11.10	-5.53
05/22-06/05/91		-6.35	-20.52	-11.52	-4.63
06/05-06/19/91	06/12	-7.15	-5.53	90.50	-7.48
06/19-07/02/91		-5.94	-4.08	48.03	-7.46
07/02-07/17/91	07/09	-5.84	-8.11	20.12	-6.47
07/17-07/31/91		-5.56	-5.41	-6.51	-8.02
07/31-08/14/91	08/07	missing	missing	missing	-5.44
08/14-08/28/91		-5.53	-8.70	-2.18	-5.95
08/28-09/12/91	09/05	-3.40	-2.26	2.08	-3.81
09/12-09/25/91		-6.17	-8.96	6.12	-9.44
09/25-10/09/91	10/02	-1.99	-4.57	7.75	-6.76
10/09-10/24/91		-2.92	-2.12	18.51	-4.43
10/24-11/06/91	10/31	-4.41	-5.96	23.42	-5.91
11/06-11/19/91		-7.82	-2.58	37.54	-5.39
11/19-12/04/91	11/26	-5.99	-5.57	95.88	-6.07
12/04-12/18/91		-11.94	-6.32	214.59	8.21
12/18-01/15/92	12/31	-6.81	-7.72	32.68	6.88

a) Counting error estimated at $\leq 0.82 \sqrt{V}$ where V is entry value. Values with minus (-) are minimum detectable amounts or MDAs.

Table 2. Co-60 Concentrations in 1991

(Values in fCi/L)^a

Date	Plot X	Shell Bluff fCi/L	Above Vogtle fCi/L	Below Vogtle fCi/L	H301 Bridge fCi/L
01/03-01/17/91		-5.80	-4.20	15.00	-4.30
01/17-01/31/91	01/24	-3.30	-5.00	12.80	-2.00
01/31-02/14/91		-8.30	-12.40	19.30	-9.70
02/14-02/28/91	02/21	-5.50	-9.60	8.10	3.50
02/28-03/14/91		-13.20	-4.10	32.10	-4.20
03/14-03/27/91	03/21	2.90	-7.00	58.70	-6.60
03/27-04/10/91		missing	missing	17.00	8.10
04/10-04/24/91	04/17	-6.60	-5.30	7.00	-4.60
04/24-05/08/91		-8.80	-6.60	-13.00	-5.20
05/08-05/22/91	05/15	-6.10	-6.40	4.80	-5.20
05/22-06/05/91		-6.30	-19.30	10.40	-3.60
06/05-06/19/91	06/12	-8.20	-5.00	36.90	11.40
06/19-07/02/91		-6.10	-4.30	109.50	13.50
07/02-07/17/91	07/09	-3.80	-6.00	37.70	4.90
07/17-07/31/91		-5.20	-4.30	17.70	6.50
07/31-08/14/91	08/07	missing	missing	missing	6.20
08/14-08/28/91		-5.20	-8.40	14.80	-6.00
08/28-09/12/91	09/05	-3.90	-2.60	2.70	-4.00
09/12-09/25/91		-6.50	-7.90	-4.70	-9.10
09/25-10/09/91	10/02	2.50	-4.10	-5.00	-6.60
10/09-10/24/91		-3.40	-2.00	8.60	-4.40
10/24-11/06/91	10/31	-4.50	-5.60	25.40	-6.70
11/06-11/19/91		-8.90	-2.30	7.20	-4.20
11/19-12/04/91	11/26	-5.30	-5.70	21.10	-4.00
12/04-12/18/91		-11.50	-5.80	32.90	-4.70
12/18-01/15/92	12/31	-6.10	-7.30	8.80	-7.20

a) Counting error estimated at $\leq 0.75 \sqrt{V}$ where V is entry value. Values with minus (-) are minimum detectable amounts or MDAs.

Table 3. Cs-137 Concentrations in 1991

(Values in fCi/L)^a

Date	Plot X	Shell Bluff	Above Vogtle	Below Vogtle	H301 Bridge
		fCi/L	fCi/L	fCi/L	fCi/L
01/03-01/17/91		-3.54	-2.15	-1.61	6.78
01/17-01/31/91	01/24	2.22	4.00	4.09	9.16
01/31-02/14/91		11.92	11.08	11.43	44.99
02/14-02/28/91	02/21	-3.20	-5.65	8.04	35.50
02/28-03/14/91		18.63	9.44	11.99	35.25
03/14-03/27/91	03/21	2.86	5.99	10.45	15.43
03/27-04/10/91		missing	missing	5.88	18.62
04/10-04/24/91	04/17	8.73	3.85	3.87	10.96
04/24-05/08/91		8.93	5.03	8.63	30.59
05/08-05/22/91	05/15	4.19	-3.28	2.62	10.33
05/22-06/05/91		4.99	-9.87	12.18	26.68
06/05-06/19/91	06/12	5.30	-3.32	8.38	36.16
06/19-07/02/91		-4.02	5.10	10.47	27.49
07/02-07/17/91	07/09	-2.67	5.77	6.28	15.74
07/17-07/31/91		-3.72	-2.82	-3.39	-2.84
07/31-08/14/91	08/07	missing	missing	missing	15.88
08/14-08/28/91		6.49	-4.23	5.33	15.51
08/28-09/12/91	09/05	-1.44	-1.28	0.94	7.02
09/12-09/25/91		6.52	-5.43	5.96	34.11
09/25-10/09/91	10/02	1.69	-2.57	-2.89	29.52
10/09-10/24/91		2.32	2.81	3.40	12.68
10/24-11/06/91	10/31	-3.14	6.39	4.49	18.41
11/06-11/19/91		8.40	5.40	3.87	9.35
11/19-12/04/91	11/26	6.97	5.81	3.40	7.05
12/04-12/18/91		11.99	8.42	4.16	9.58
12/18-01/15/92	12/31	-3.66	-4.01	-2.14	14.77

a) Counting error estimated at $\leq 0.58 \sqrt{V}$ where V is entry value. Values with minus(-) are minimum detectable amounts or MDAs.

Table 4. Relative Isotopics Below Vogtle During 1991^{a,b}
 (Sample site 0.1 mi downstream of Vogtle)

Date	Plot X	Co-58 fCi/L	Isotopics % of Co-58 Activity below Vogtle								
			Cr-51	Mn-54	Co-57	Co-58	Fe-59	Co-60	Nb-95	Zr-95	Cs-137
01/03-01/17/91		112.68	0.00	7.60	0.00	100.00	0.00	13.33	0.00	0.00	-1.43
01/17-01/31/91	01/24	80.79	0.00	0.00	0.00	100.00	0.00	15.88	0.00	0.00	5.06
01/31-02/14/91		127.40	0.00	0.00	0.00	100.00	0.00	15.11	0.00	0.00	8.97
02/14-02/28/91	02/21	30.23	0.00	23.08	0.00	100.00	0.00	26.70	0.00	0.00	26.59
02/28-03/14/91		32.11	0.00	12.57	0.00	100.00	0.00	33.86	0.00	0.00	37.34
03/14-03/27/91	03/21	154.83	0.00	11.95	0.00	100.00	0.00	37.92	0.00	0.00	6.75
03/27-04/10/91		81.33	0.00	0.00	0.00	100.00	0.00	20.85	0.00	0.00	7.23
04/10-04/24/91	04/17	21.40	0.00	0.00	0.00	100.00	0.00	32.74	0.00	0.00	18.09
04/24-05/08/91		23.39	0.00	61.11	0.00	100.00	0.00	-55.56	0.00	0.00	36.89
05/08-05/22/91	05/15	11.10	0.00	35.00	0.00	100.00	0.00	43.12	0.00	0.00	23.59
05/22-06/05/91		-11.52	0.00	0.00	0.00	*100.00	0.00	+90.00	0.00	0.00	+105.70
06/05-06/19/91	06/12	90.50	0.00	17.43	0.00	100.00	0.00	40.79	0.00	0.00	9.26
06/19-07/02/91		48.03	0.00	34.25	0.00	100.00	0.00	228.02	0.00	0.00	21.80
07/02-07/17/91	07/09	20.12	0.00	0.00	0.00	100.00	0.00	84.98	0.00	0.00	31.21
07/17-07/31/91		-6.51	0.00	0.00	0.00	*100.00	0.00	+271.43	0.00	0.00	*52.06
07/31-08/14/91	08/07	missing	missing	missing	missing	missing	missing	missing	missing	missing	missing
08/14-08/28/91		-2.18	0.00	+330.43	0.00	*100.00	0.00	+678.26	0.00	0.00	+244.33
08/28-09/12/91	09/05	2.08	0.00	0.00	0.00	100.00	0.00	129.63	0.00	0.00	45.20
09/12-09/25/91		6.12	0.00	0.00	0.00	100.00	0.00	-76.19	0.00	0.00	97.32
09/25-10/09/91	10/02	7.75	0.00	0.00	0.00	100.00	0.00	-63.92	0.00	0.00	-37.30
10/09-10/24/91		18.51	0.00	0.00	0.00	100.00	0.00	46.21	0.00	0.00	18.37
10/24-11/06/91	10/31	23.42	0.00	26.25	0.00	100.00	0.00	108.33	0.00	0.00	19.17
11/06-11/19/91		37.54	0.00	0.00	0.00	100.00	0.00	19.14	0.00	0.00	10.31
11/19-12/04/91	11/26	95.88	0.00	0.00	0.00	100.00	0.00	21.96	0.00	0.00	3.55
12/04-12/18/91		214.59	0.00	4.93	0.00	100.00	0.00	15.31	56.40	16.02	1.94
12/18-01/15/92	12/31	32.68	0.00	0.00	0.00	100.00	0.00	26.79	18.87	0.00	-6.55

a) All values are relative activities on resin sample, except for ¹³⁷Cs, which is corrected for collection efficiency.

b) MDA values of ⁵⁸Co are denoted with a minus(-) sign. Isotopic % is denoted with an astrick(*) for MDA cases, and with a plus(+) for detected cases.

Table 5. Tritium Concentrations in 1991
(Values in pCi/mL)^a

Date	Shell Bluff	Vogtle Vicinity ^b			Hwy 301
		+0.3 mi	0.0 mi	-0.1 mi	
1/03/91	<2.0	<3.0	10.0±1.0	2.7±0.8	4.2±0.8
2/14/91	2.4±0.8	2.2±0.8	4.3±0.9	5.2±0.9	3.4±0.9
3/14/91	1.9±0.8	5.6±0.9	8.0±1.0	2.5±0.8	3.0±0.9
4/10/91	<1.0	<1.0	<1.0	<1.0	2.8±0.7
5/08/91	<1.0	<1.2	<1.0	<1.0	<1.6
6/05/91	1.7±0.8	2.1±0.9	<1.9	3.0±0.9	3.9±0.9
7/02/91	3.7±0.9	2.1±0.9	2.9±0.9	2.9±0.9	5.8±1.0
8/14/91	<0.9	<0.8	<0.6	<0.8	2.2±0.6
9/12/91	<0.7	<0.6	<0.9	<1.0	3.8±0.6
10/09/91	<1.1	<1.0	<0.5	<0.8	5.0±0.5
11/06/91	<0.5	<0.6	4.3±0.5	2.3±0.4	2.0±0.4
12/04/91	<0.4	<0.8	<1.2	<0.9	1.1±0.4 ^c

(a) Errors are 2σ values (other errors in report are 1σ).

(b) Miles are measured upstream of Vogtle outfall.

(c) Level peaked at 63 pCi/mL on 12/28 following tritium release from K-Reactor (details per ref 8).

Table 6. Sediment Concentrations in 1991
 (Values in fCi/g = pCi/Kg)

Isotope	Date	Vogtle Vicinity ^a			Hwy 301
		+0.3 mi	0.0 mi	-0.1 mi	
⁵⁸ Co	02/07/91	<2.6	<4.3	32.0 ± 1.7	<5.0
	05/29/91	<2.1	<2.6	<2.6	<3.5
	09/04/91	<4.8	<10.1	<4.4	<5.9
	12/11/91	<2.0	<4.8	2.3 ± 0.8	<3.7
⁶⁰ Co	02/07/91	<3.0	<3.2	<2.9	<3.7
	05/29/91	<1.7	<1.9	<2.1	<2.6
	09/04/91	<3.1	<5.0	<3.0	<5.9
	12/11/91	<3.1	<3.4	<2.1	<2.4
¹³⁷ Cs	02/07/91	8.6 ± 1.3	<3.9	4.7 ± 1.2	85 ± 3
	05/29/91	5.6 ± 0.8	3.6 ± 0.9	4.8 ± 1.0	69 ± 2
	09/04/91	4.6 ± 1.5	<8.8	<3.5	124 ± 3
	12/11/91	4.6 ± 1.4	<4.0	2.6 ± 0.8	105 ± 1

(a) Miles are measured upstream of Vogtle outfall.

Table 7. Comparison of Maximum 1987-1991 Levels with DOE Guides for Drinking Water

All measured levels are from samples 0.1 mi downstream of Vogtle outfall unless specified otherwise.

Isotope	Maximum Concentration (pCi/L)					
	1987	1988	1989	1990	1991	DOE Guide ^a
³ H	47,300 ^b	4200 ^c	5800 ^c	42,600 ^b	10,000 ^d	2,000,000
⁵¹ Cr	3.7	2.8	0.31	0.08	---	1,000,000
⁵⁴ Mn	0.06	0.40	0.61	0.03	0.02	50,000
⁵⁷ Co	0.02	0.02	0.01	0.005	---	100,000
⁵⁸ Co	16.8	15.5	3.90	2.37	0.21	40,000
⁵⁹ Fe	0.22	0.49	0.05	0.02	---	20,000
⁶⁰ Co	0.14	0.49	0.23	0.08	0.11	5,000
⁹⁵ Nb	0.10	0.50	0.10	---	0.12	40,000
⁹⁵ Zr	0.17	0.23	0.04	---	0.03	60,000
¹³⁷ Cs	0.15 ^c	0.39 ^c	0.12 ^c	0.09 ^c	0.05 ^c	3,000

a) DOE 5400.5 (details per reference 9).

b) Value at Vogtle outfall.

c) Value at Highway 301 Bridge.

d) Value at Vogtle outfall, which is less than 63,000 pCi/L observed at Hwy 301 Bridge on 12/28 per K-Reactor release (details per reference 8).

e) Not detected at any location.

FIGURE 1. Co-58 in Savannah River in 1991

Locations are defined as:
SB = Shell Bluff
VA = 0.3 mi Upstream of Vogtle
VB = 0.1 mi Downstream of Vogtle
BR = Highway 301 Bridge

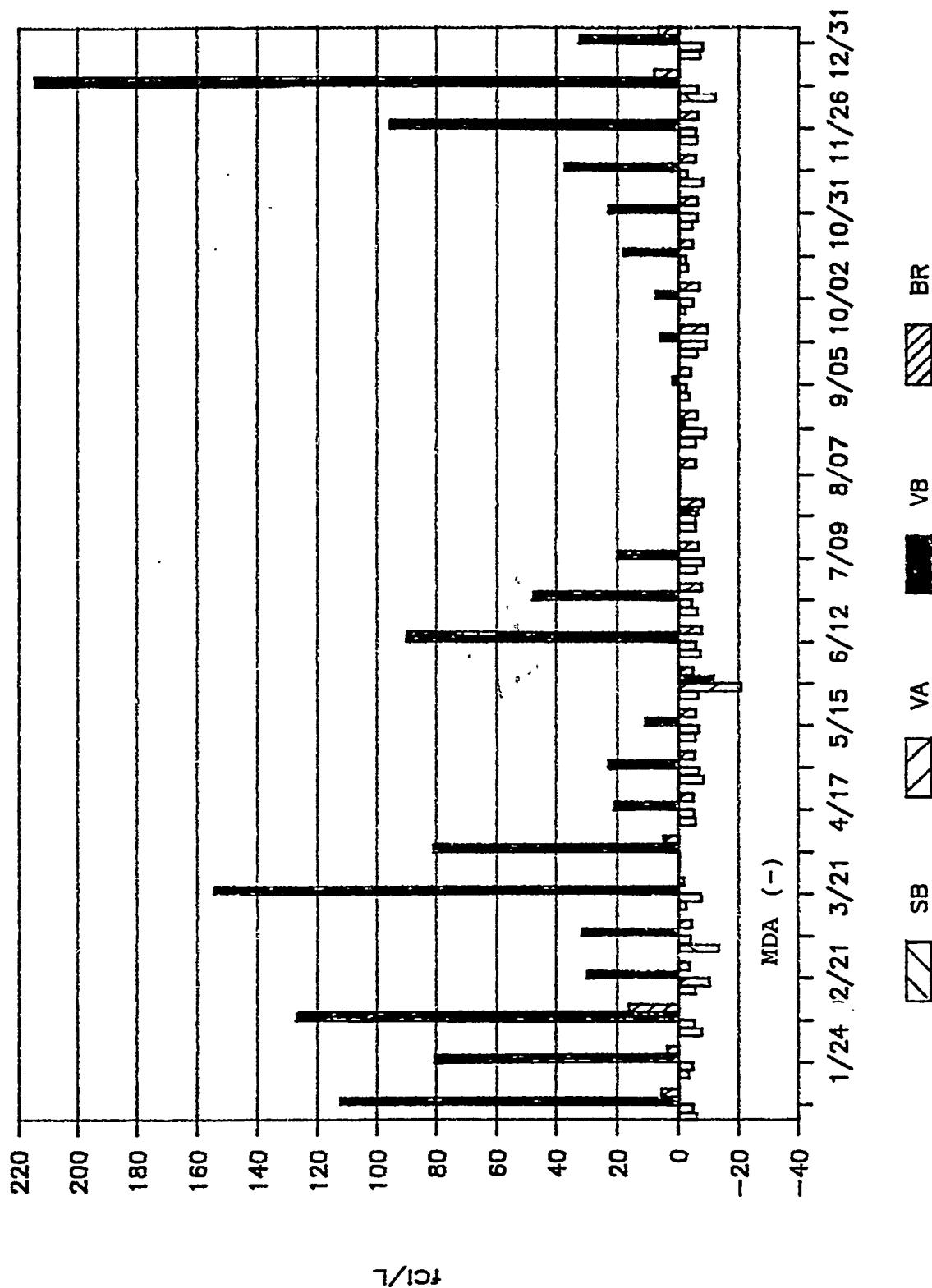


FIGURE 2. Co-60 in Savannah River in 1991

Locations are defined as:
SB = Shell Bluff
VA = 0.3 mi Upstream of Vogtle
VB = 0.1 mi Downstream of Vogtle
BR = Highway 301 Bridge

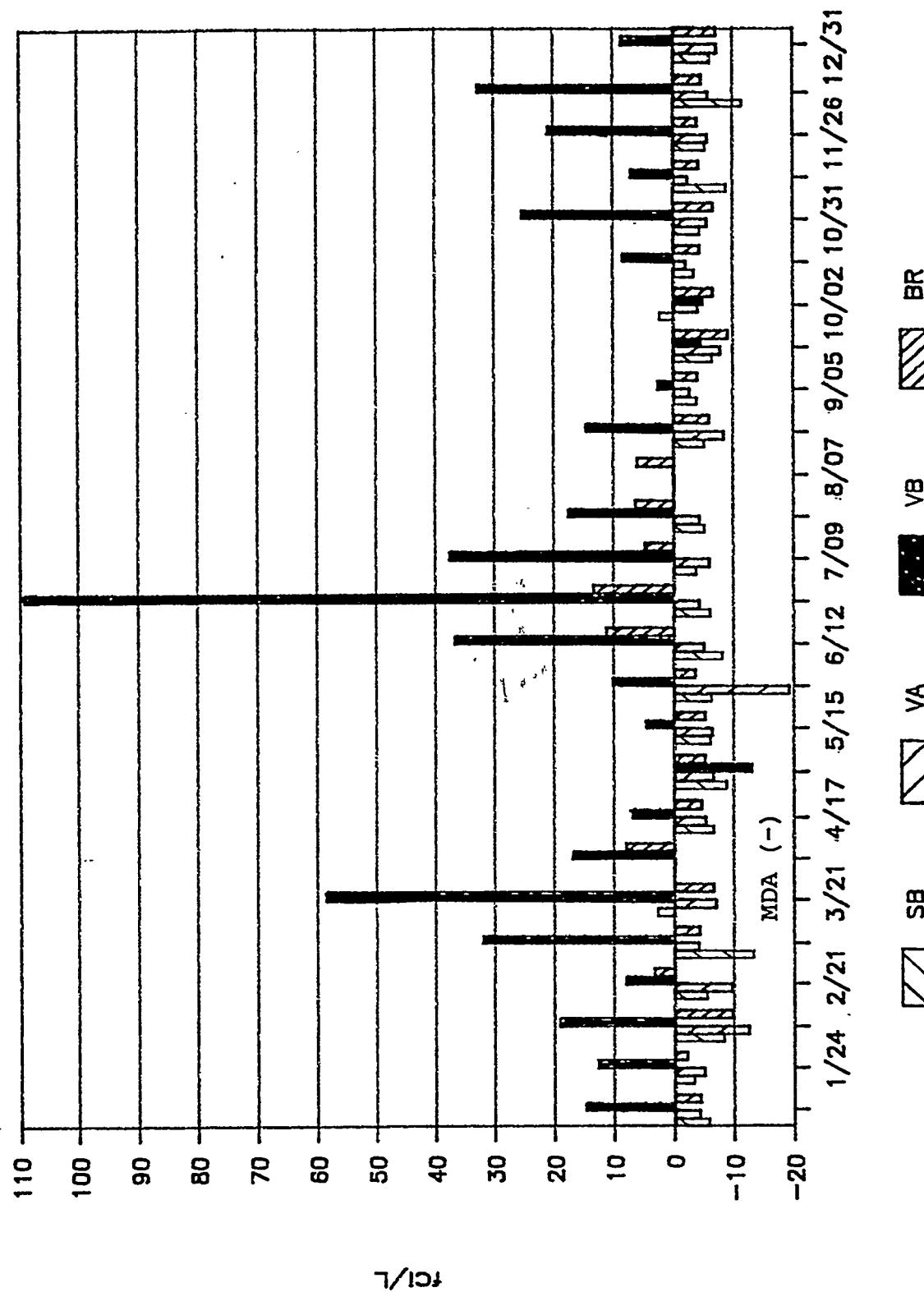
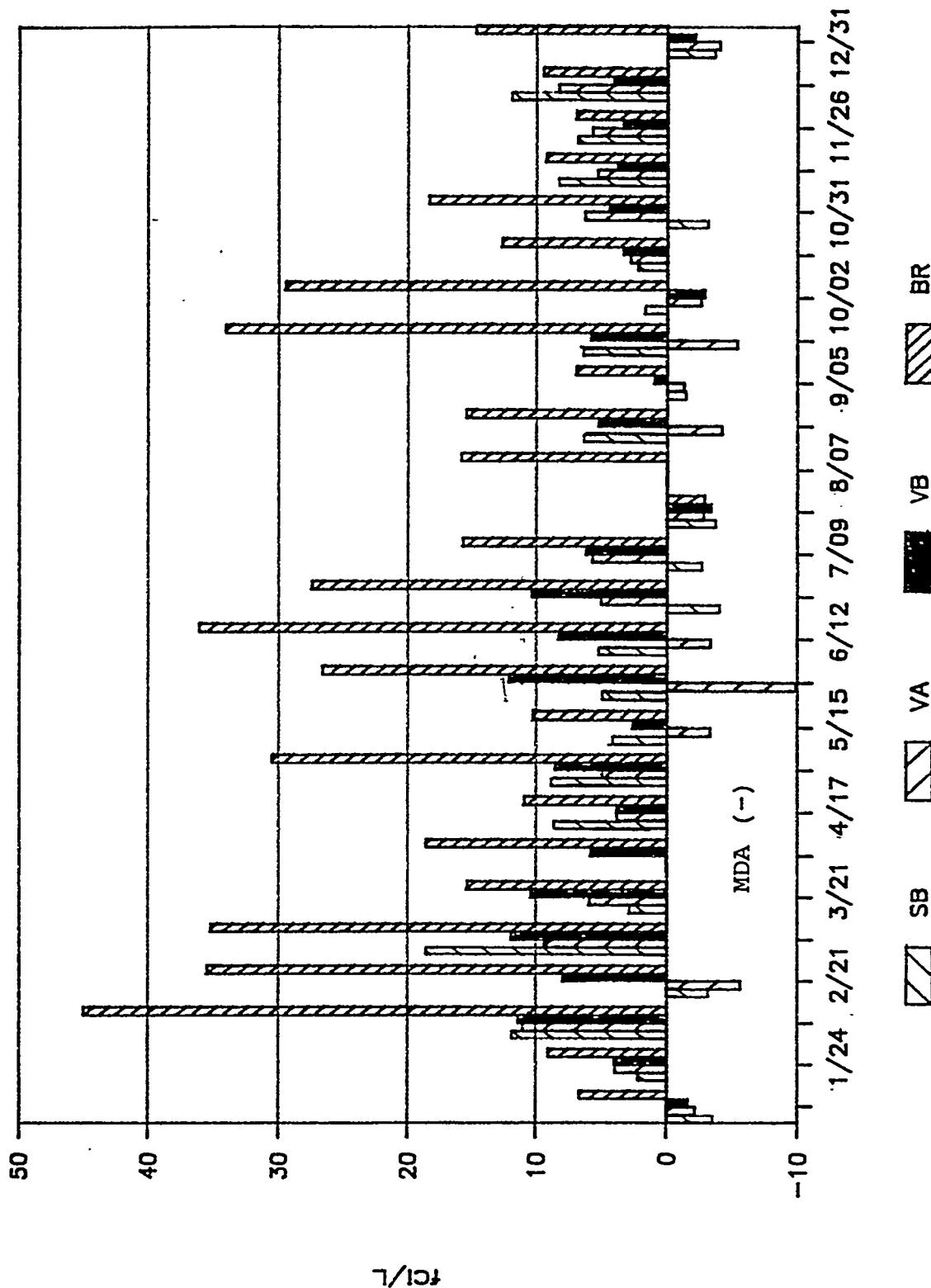


FIGURE 3. Cs-137 in Savannah River in 1991

Locations are defined as:
SB = Shell Bluff
VA = 0.3 mi Upstream of Vogtle
VB = 0.1 mi Downstream of Vogtle
BR = Highway 301 Bridge



Gamma Peak KeV

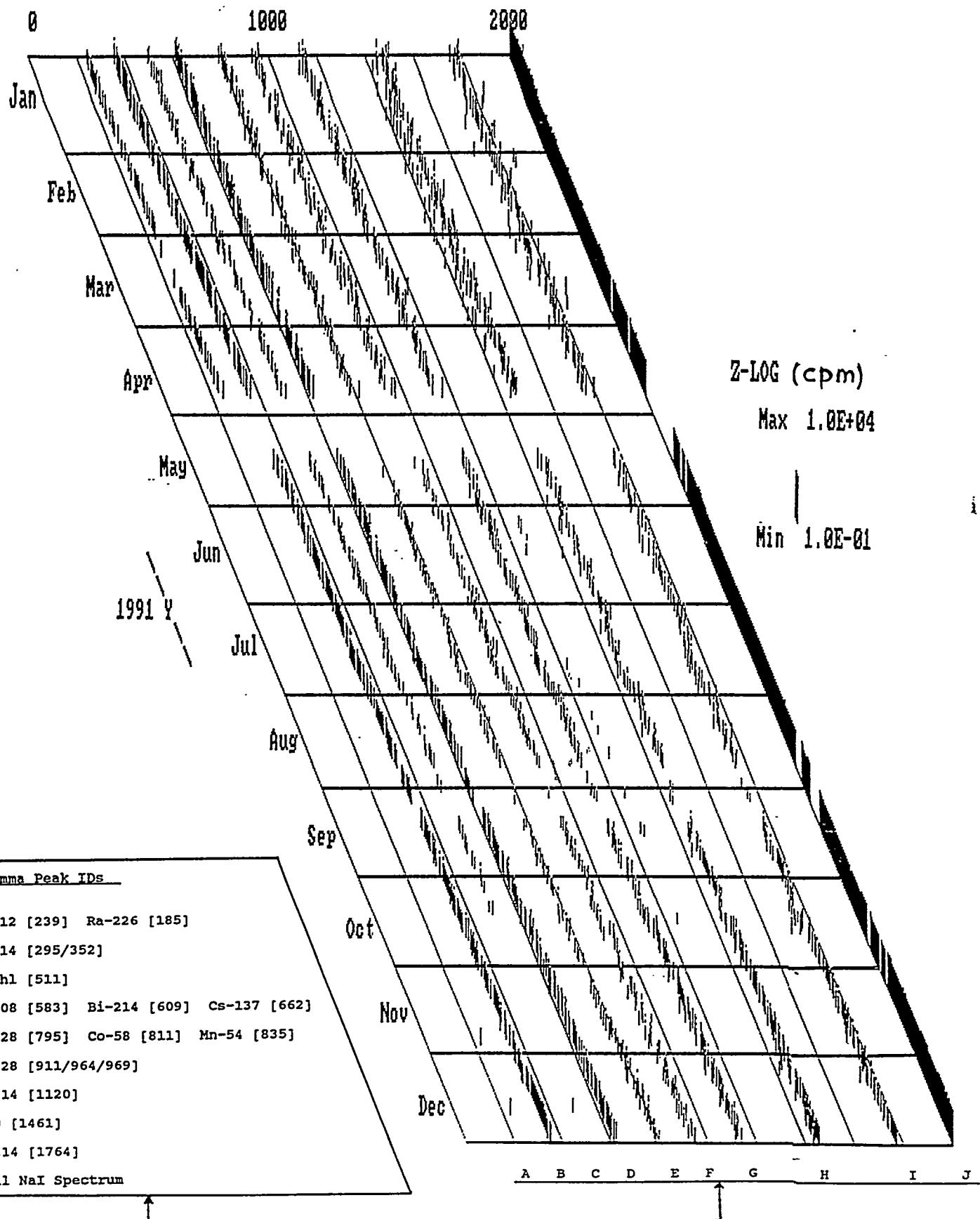
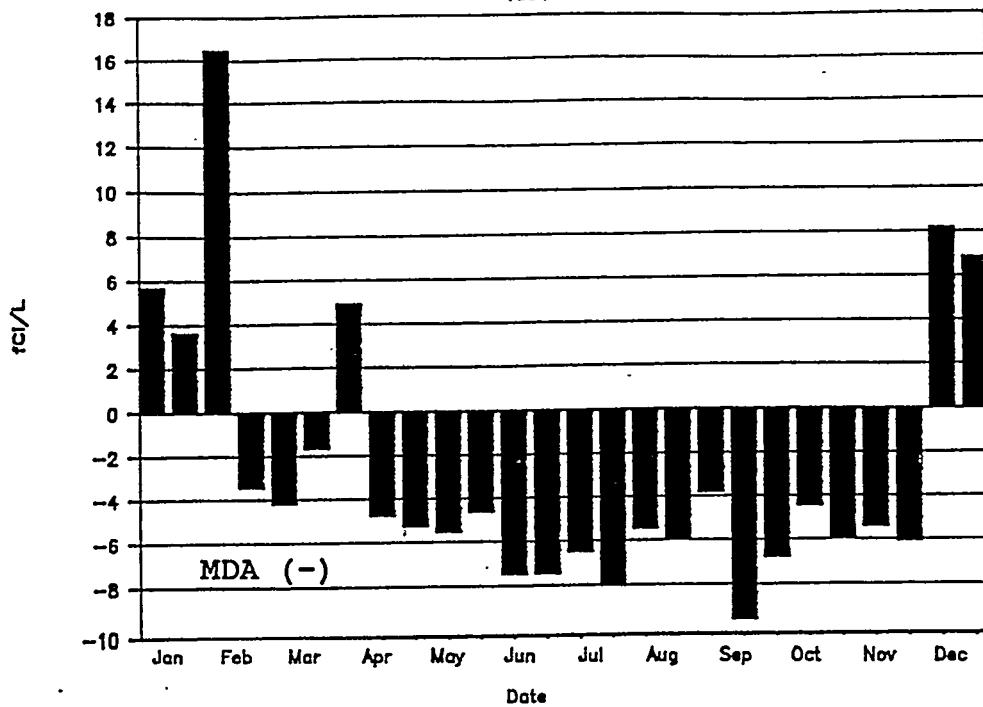


FIGURE 4. Results for Underwater NaI Detector at Hwy 301 Bridge

Resin Co-58 / Hwy301 Bridge

1991



Underwater NaI Co-58

1991

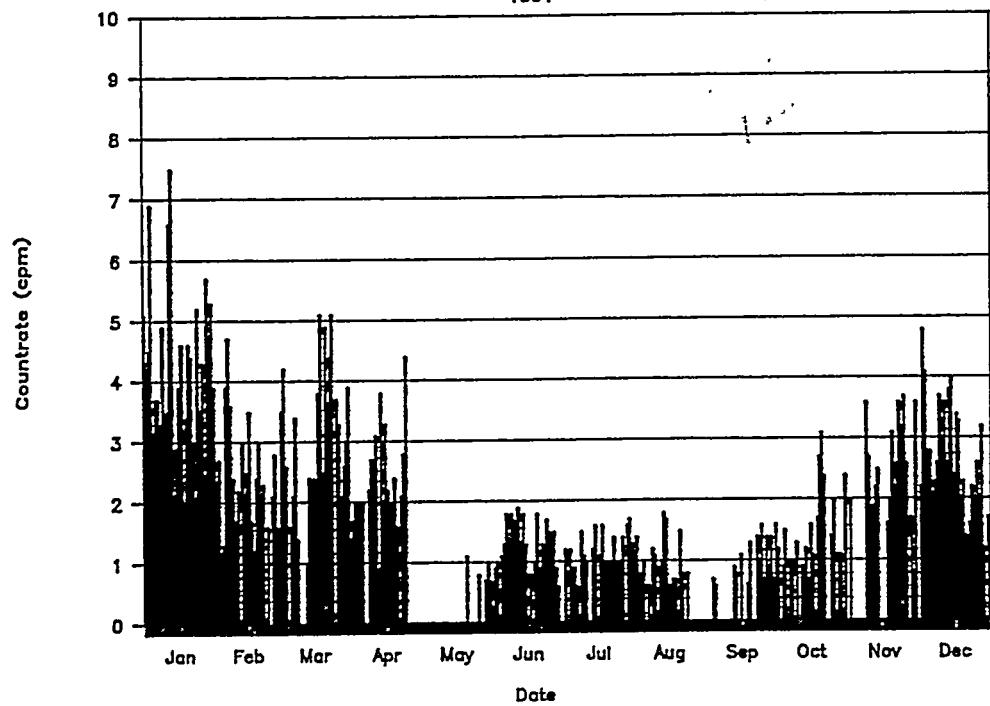


FIGURE 5. Comparison of Underwater NaI and Resin Sample Results for Co-58 detection at Hwy 301 Bridge

Appendix - Map of Sample Locations

