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Radioactive Effluents in the Savannah River: Summary Report for 1989 (U)

W. G. Winn

Westinghouse Savannah River Company
Savannah River Site
Aiken, SC 29808



Prepared for the U. S. Department of Energy under contract no. DE-AC09-89SR18035

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Radioactive Effluents in the Savannah River— Summary Report for 1989 (U)

By W. G. Winn

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Abstract

Researchers at the Savannah River Site have low-level radiometric studies of the Savannah River to distinguish between the effluent contributions of the Savannah River Site and Plant Vogtle. Since the startup of Plant Vogtle in 1987, researchers have routinely detected neutron-activated isotopes in controlled releases, but all have been well below the Department of Energy's (DOE) guidelines. The study has found that processing improvement at Plant Vogtle during 1989 have lowered the activities of effluents from Plant Vogtle. These studies will continue on a routine basis because they provide disturbing trends before actual health concerns evolve.

Introduction

During 1989 low-level radiometric studies of the Savannah River continued to distinguish between effluent contributions from Plant Vogtle and the Savannah River Site. Both institutions have interest in the measurements of the radioactive effluents. Analysis of the data helps the institutions address disturbing trends before they become health and legal concerns.

The Environmental Technology Section (ETS) conducted radiometric studies of Plant Vogtle (since late 1986) prior to its startup.¹ The plant has two 1100 MWe pressurized water reactors developed by Westinghouse: Unit 1 started commercial operations in June 1987, and Unit 2 followed in May 1989. During powered operations ETS routinely detected neutron-activated isotopes in controlled releases, but all activities have been several orders of magnitude below the DOE guide values.^{2,3,4} Processing improvements in 1989 lowered the activities of Vogtle effluents in the river. The Plant Vogtle release data and the ETS measurements correlated well from 1987-1989.

Researchers measure effluents by using ETS ultralow-level radiometric measurement techniques. The techniques generally include concentrating river samples on resins, preparing them in the lab, and then counting the effluents in the underground counting facility in 735-A.⁵ Researchers use gamma spectrometry for most of the counting and liquid-scintillation counting for tritium. Gamma-ray analysis provides detection limits that are thousands of times lower than those routinely achievable. (Tritium analysis also uses state-of-the-art detection sensitivity.) An underwater NaI(Tl) at

Highway 301 bridge provides continuous gamma monitoring of the Savannah River.

Discussion

During 1989 observers found noticeably less radioactive effluents in the Savannah River than in 1987-1988. This observation correlates with improvements made at Vogtle for processing liquid effluents. The Vogtle effluents continue to be dominated by ⁵⁸Co, which had a maximum concentration of 3.9 pCi/L at the beginning of 1989, but stayed below 0.6 pCi/L after February. Observers found trace amounts of ⁵¹Cr, ⁵⁴Mn, ⁵⁷Co, ⁵⁹Fe, ⁶⁰Co, ⁹⁵Nb, and ⁹⁵Zr in the effluents; similar observations were made in 1987 and 1988. Monitoring of tritium and ¹³⁷Cs found their levels consistent with known SRS sources.

River Sampling

Researchers collected samples from the Savannah River at Shell Bluff, areas near the Vogtle outfall, and by the Highway 301 Bridge. Effluents from Vogtle and SRS cause the increase in activity from Shell Bluff to the Highway 301 bridge. Measurements upstream and downstream of the Vogtle outfall (0.3 mi above, 0.1 mi below, and at the Vogtle outfall) identify Vogtle contributions.

Three types of samples were collected—resin, water, and sediment. Resin-concentrator samples, which consist of approximately 25 g of resin in a porous nylon bag, stay in the water for two weeks before being returned to the lab. Researchers directly collect water samples. A specially

designed sampler that is pulled along the bottom of the river scoops sediment samples. The Vogtle prestartup study details the sampling methods.¹

Laboratory Measurements

The resin concentrator samples provide the most comprehensive isotopic information. Technicians dried and turned each sample into ash to achieve a smaller volume and better geometry for the high-purity germanium detector (HPGe) detector. Typically, two HPGe detectors, which have ⁶⁰Co standard efficiencies of 20% and 25%, counted the samples. (A new detector, with a ⁶⁰Co standard efficiency of 90%, analyzed special cases.) Analysts processed the samples overnight (or over the weekend) to assure good counting statistics for detecting low-level activities. An IBM/PC-XT used the GRABGAM code to analyze the resulting HPGe gamma-ray spectra to yield the activity (fCi) of each isotope detected in the resin sample.⁵ By using empirical calibration data that relate the resin mass and collection time to the effective water volume (L), analysts obtained the average isotopic concentration (fCi/L) for the collection period.

A TRICARB LL 2050A low-level liquid scintillation analyzer examined samples with 3 mL of water and 20 mL of OPTI-FLUOR™ scintillant for tritium. Duplicate samples with 0.25 mL of tritium standard checked the performance of the automatic quench corrections.

Sediment samples were transferred to 1-L Marinelli beakers and then counted on the HPGe detectors. Analysts divided the sample isotopic activities (fCi) from the gamma-ray analysis 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 the Highway 301 bridge. The detector includes a 9 in. diameter by 4 in. long NaI(Tl) detector, four photo-multiplier tubes, and a high-voltage/preamp unit; everything is contained in a waterproof stainless steel housing. The detector views its surroundings through a thin hemispherical dome.

The detector receives operating power through a waterproof cable connected to a ≈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 same cable transmits the AC-coupled detector signals to the cabin, where they are amplified and sent to a multichannel analyzer.

A multichannel analyzer, which is comprised of a COMPAQ 286 computer and an ACE MCA card from EG&G ORTEC, collects spectral data. Spectra are stored on a floppy disk in sequenced 24-hr periods for two to three weeks until researchers retrieve the disk and analyze it at the laboratory. This detector system provides better time resolution (one day) than the resin concentrator method (two weeks), but its sensitivity is one order of magnitude less.³

Results

This section summarizes the measurement results and comments briefly on their significance. The next section (Discussion) provides a more detailed account.

The resin sampler analysis continued to indicate ⁵⁸Co, ⁶⁰Co, and ¹³⁷Cs, which were the most significant gamma-emitting radionuclides detected in the earlier studies.^{1,2,3} The 1989 results for these isotopes are given in tables 1-3, along with plots in figures 1-3. The other detected man-made radionuclides correlate with ⁵⁸Co and are included in Table 4. The activities are the lowest observed since the Vogtle startup in 1987.

Water analysis for tritium followed techniques used earlier.^{1,2} Table 5 summarizes the 1989 tritium results. These low-tritium levels have essentially not changed over the past few years.

Sediments were analyzed for ⁵⁸Co, ⁶⁰Co, and ¹³⁷Cs to appraise whether any deposition is occurring during the transport of these isotopes in river water. The corresponding results (see Table 6) show that only very low levels of those man-made radionuclides continue to be observed.

Figure 4 presents an isometric plot of count rate vs gamma energy and date from the data collected by the underwater NaI(Tl) detector. Figures 5 and 6 compare the ⁵⁸Co detected by the underwater NaI(Tl) with that detected with the resin concentrator samples. The two methods continue to correlate well.

Conclusions

In 1989 overall man-made radioactivity in the Savannah River was significantly lower than in the preceding year. This was primarily due to improvements in Plant Vogtle processing of effluent waste. River water samples show that Vogtle effluents continue to contain ⁵¹Cr, ⁵⁴Mn, ⁵⁷Co, ⁵⁸Co, ⁵⁹Fe, ⁹⁵Nb and ⁹⁵Zr. Aside from these Vogtle contributions, the other activities remained consistent with levels of SRS radionuclides ³H, ⁶⁰Co, and ¹³⁷Cs detected prior to Vogtle

Table 1. ⁵⁸Co Concentrations in 1989 (Values in fCi/L)^a

Date	Shell Bluff fCi/L	Above Vogtle fCi/L	Below Vogtle fCi/L	H301 Bridge fCi/L
12/14/88 - 1/5/89	-2.48	-1.96	3897.07	443.40
1/5 - 1/19/89	-2.55	-2.65	2113.70	40.22
1/19 - 2/2/89	-3.38	-2.85	1099.72	69.64
2/2 - 2/16/89	-2.49	-1.97	377.67	33.80
2/16 - 3/2/89	-4.24	-5.77	505.16	51.12
3/2 - 3/16/89	-2.80	-9.46	176.61	30.71
3/16 - 3/30/89	-6.65	-9.49	121.79	20.79
3/30 - 4/13/89	-7.01	-9.91	98.91	22.78
4/13 - 4/27/89	-3.88	-7.59	82.53	5.78
4/27 - 5/11/89	-4.63	-3.58	29.37	-3.53
5/11-5/25/89	-3.87	-5.85	143.15	-5.82
5/25 - 6/08/89	-5.51	na	246.54	8.07
6/08-6/22/89	na	-5.25	31.44	8.88
6/22 - 7/6/89	-6.92	-7.05	24.50	-6.85
7/6-7/20/89	-5.75	-6.20	124.14	8.85
7/20 - 8/3/89	-5.48	-5.02	50.68	-5.41
8/3 - 8/17/89	-3.29	-5.45	171.97	7.12
8/17-8/31/89	-4.92	-4.41	91.99	39.23
8/31 - 9/14/89	-4.74	-3.56	180.23	12.44
9/14 - 9/28/89	-4.09	-5.36	107.69	3.34
9/28 - 10/12/89	-3.95	-4.69	121.87	6.50
10/12 - 10/26/89	-10.28	-18.13	540.51	17.50
10/26 - 11/09/89	-5.19	-5.61	755.61	15.63
11/09 - 11/22/89	na	-5.03	422.52	163.47
11/22 - 12/07/89	-4.11	-4.71	299.10	16.92
12/07 - 12/21/89	-9.91	-7.26	35.59	-9.78
12/21/89 - 01/04/90	-5.26	-6.29	20.40	-9.07

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

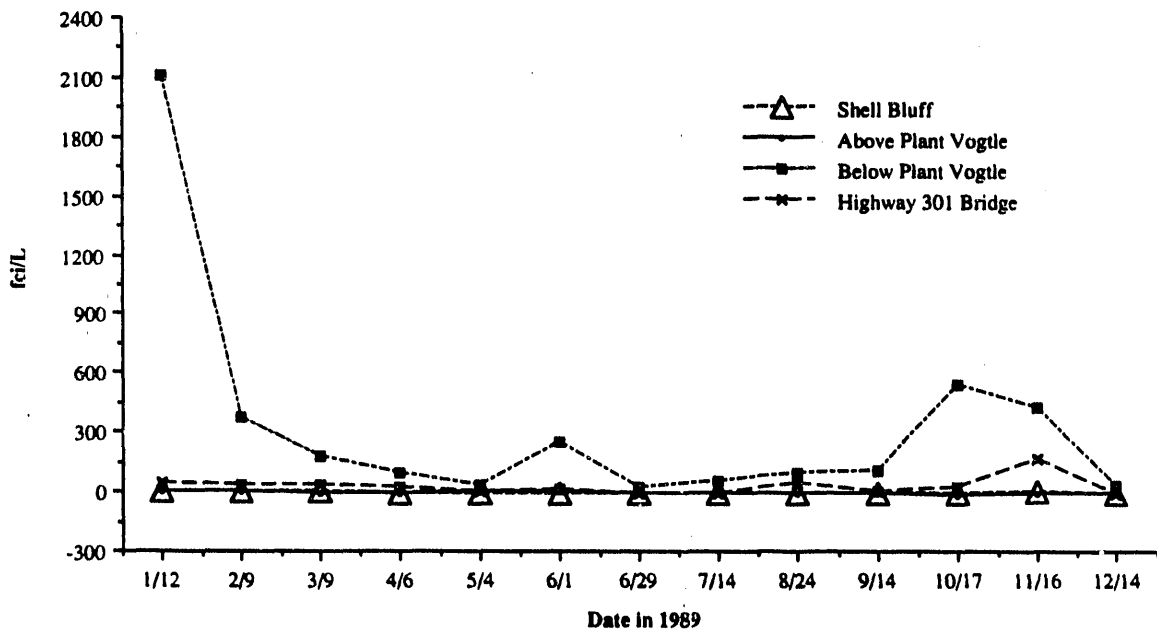


Figure 1. ⁵⁸Co in the Savannah River in 1989 [Negative Values are Minimum Detectable Amounts (MDAs)]

Table 2. ⁶⁰Co Concentrations in 1989 (Values in fCi/L)^a

Date	Shell Bluff fCi/L	Above Vogtle fCi/L	Below Vogtle fCi/L	H301 Bridge fCi/L
12/14 - 1/5/89	-2.00	-1.60	225.50	46.90
1/5 - 1/19	-2.40	-2.40	94.10	5.90
1/19 - 2/2	-3.30	-2.70	93.10	8.60
2/2 - 2/16	-2.40	-1.90	59.10	8.40
2/16 - 3/2	-3.40	-4.70	47.40	9.30
3/2 - 3/16	-2.50	-8.50	49.60	7.40
3/16 - 3/30	-5.00	-8.00	37.50	13.50
3/30 - 4/13	-5.60	-7.40	13.50	-5.70
4/13 - 4/27	-3.50	-6.60	30.00	-3.30
4/27 - 5/11	-3.70	-2.80	13.70	-3.50
5/11-25	-3.70	-4.90	45.30	-4.80
5/25 - 6/08	-6.40	na	63.10	-4.40
6/08-22	na	-4.50	14.90	4.80
6/22 - 7/6	-6.10	-6.20	20.70	-6.40
7/6-20/89	-5.40	-5.20	25.30	8.30
7/20 - 8/3/89	-4.80	-4.80	8.20	-5.10
8/3 - 8/17/89	-3.00	-4.90	44.70	11.20
8/17-31/89	-4.70	-4.40	35.60	7.80
8/31 - 9/14/89	-4.10	-3.00	95.70	8.80
9/14 - 9/28/89	-5.00	-4.90	31.30	-3.00
9/28 - 10/12/89	3.60	-7.50	10.20	-4.50
10/12 - 10/26/89	-5.10	-9.60	88.70	6.20
10/26 - 11/09/89	-3.50	-3.40	44.70	-7.20
11/09 - 11/22/89	na	-3.10	21.30	7.50
11/22 - 12/07/89	-2.60	-3.00	49.60	7.60
12/07 - 12/21/89	-7.30	-5.50	-10.00	-6.50
12/21 - 01/04/90	-3.20	-8.60	-6.20	-5.90

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

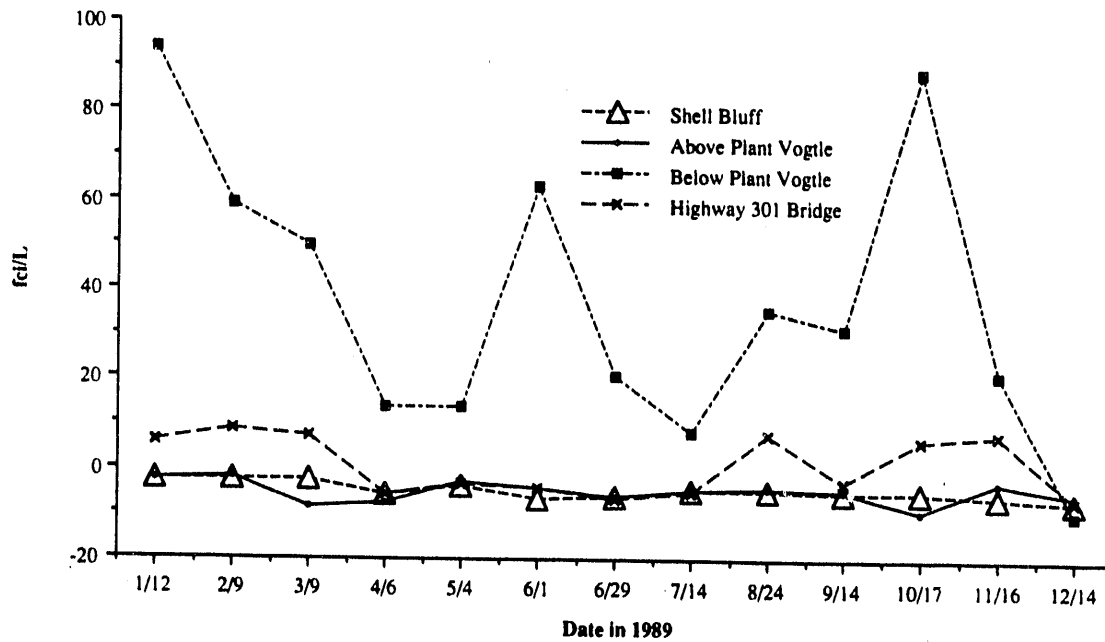


Figure 2. ⁶⁰Co in the Savannah River in 1989 (Negative Values are MDAs)

Table 3. ¹³⁷Co Concentrations in 1989 (Values in fCi/L)^a

Date	Shell Bluff fCi/L	Above Vogtle fCi/L	Below Vogtle fCi/L	H301 Bridge fCi/L
12/14/88 - 1/5/89	-3.10	4.45	11.86	56.44
1/5 - 1/19/89	2.28	2.30	11.62	19.83
1/19 - 2/2/89	5.06	3.38	4.05	34.09
2/2 - 2/16/89	3.60	6.37	3.41	30.61
2/16 - 3/2/89	8.37	11.35	10.99	37.13
3/2 - 3/16/89	5.77	11.36	13.26	77.54
3/16 - 3/30/89	8.45	15.66	18.33	56.15
3/30 - 4/13/89	7.12	11.68	16.26	46.92
4/13 - 4/27/89	7.23	11.63	17.51	26.52
4/27 - 5/11/89	6.45	6.80	13.01	30.35
5/11-5/25/89	5.52	17.13	23.43	30.52
5/25 - 6/08/89	6.59	na	14.54	51.89
6/08-6/22/89	na	11.12	8.45	46.44
6/22 - 7/6/89	14.78	20.18	29.16	66.65
7/6 - 7/20/89	12.65	25.25	40.20	118.79
7/20 - 8/3/89	7.90	15.96	22.53	47.80
8/3 - 8/17/89	9.61	13.32	13.71	46.30
8/17-8/31/89	9.54	6.31	11.30	51.04
8/31 - 9/14/89	4.91	9.31	8.81	34.45
9/14 - 9/28/89	6.82	9.87	19.47	23.84
9/28 - 10/12/89	3.03	14.39	7.94	31.87
10/12 - 10/26/89	8.74	14.00	12.90	27.66
10/26 - 11/09/89	5.25	4.17	7.91	50.61
11/09 - 11/22/89	na	2.31	8.62	33.36
11/22 - 12/07/89	2.82	7.10	13.25	26.06
12/07 - 12/21/89	11.42	-3.71	13.38	23.55
12/21/89 - 01/04/90	5.25	4.39	10.32	32.72

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

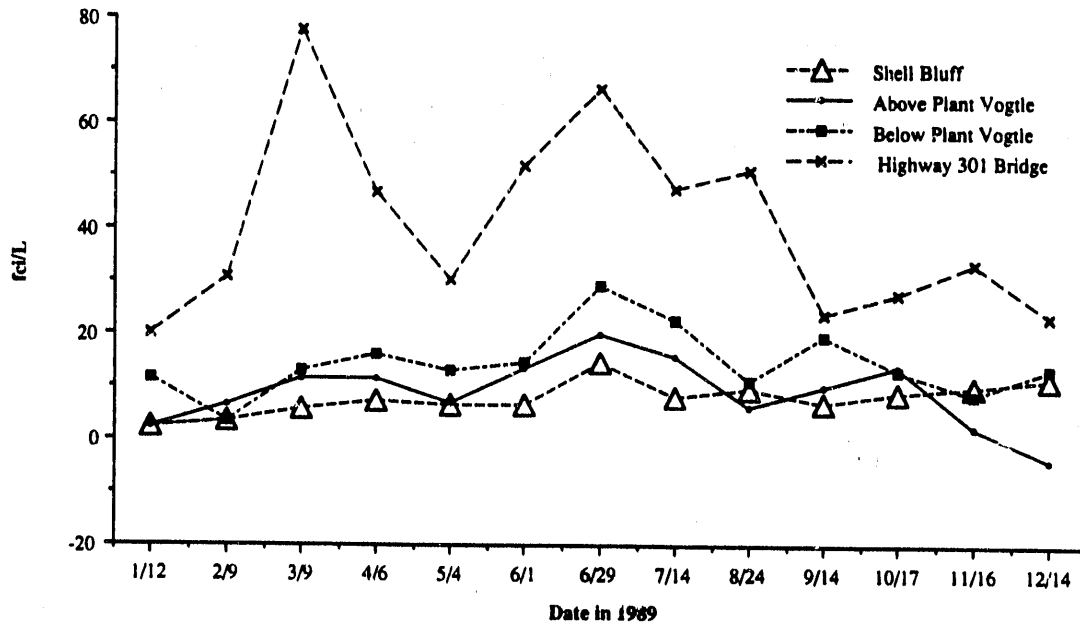


Figure 3. ¹³⁷Cs in the Savannah River in 1989 (Negative Values are MDAs)

Table 4. Relative Isotopics Below Vogtle During 1989^a

Date	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
12/14/88 - 1/5/89	3897.07	7.94	15.73	—	100.00	1.29	5.79	2.63	1.15	0.30
1/5 - 1/19/89	2113.70	—	13.96	0.28	100.00	0.42	4.45	1.09	—	0.55
1/19 - 2/2/89	1099.72	5.00	12.25	—	100.00	1.95	8.47	2.91	1.49	0.37
2/2 - 2/16/89	377.67	11.54	15.64	—	100.00	5.81	20.18	4.94	2.57	0.90
2/16 - 3/2/89	505.16	—	18.02	—	100.00	—	9.38	—	2.61	2.18
3/2 - 3/16/89	176.61	—	24.41	—	100.00	—	28.09	9.25	6.24	7.51
3/16 - 3/30/89	121.79	—	23.70	—	100.00	—	30.82	10.89	—	15.05
3/30 - 4/13/89	98.91	—	18.67	—	100.00	—	13.67	—	—	16.44
4/13 - 4/27/89	82.53	—	43.03	—	100.00	—	36.30	6.67	—	21.22
4/27 - 5/11/89	29.37	—	42.33	—	100.00	—	46.67	—	—	44.30
5/11 - 5/25/89	143.15	0.00	22.58	—	100.00	—	31.64	—	12.93	16.37
5/25 - 6/08/89	246.54	—	32.14	—	100.00	—	25.60	8.35	—	5.90
6/08 - 6/22/89	31.44	—	40.30	—	100.00	—	47.47	21.73	—	26.87
6/22 - 7/6/89	24.50	—	—	—	100.00	—	84.59	—	—	119.04
7/6 - 7/20/89	124.14	—	8.32	—	100.00	—	20.35	—	—	32.38
7/20 - 8/3/89	50.68	—	9.84	—	100.00	—	16.12	—	—	44.46
8/3 - 8/17/89	171.97	—	20.25	—	100.00	—	43.92	—	—	7.97
8/17 - 8/31/89	91.99	—	17.28	—	100.00	—	38.67	8.80	—	12.28
8/31 - 9/14/89	180.23	—	20.47	—	100.00	—	53.08	10.91	—	4.89
9/14 - 9/28/89	107.69	—	13.03	—	100.00	—	29.03	—	—	18.08
9/28 - 10/12/89	121.87	—	3.67	—	100.00	—	6.14	—	—	6.52
10/12 - 10/26/89	540.51	—	6.69	—	100.00	—	16.41	—	—	2.39
10/26 - 11/09/89	755.61	—	0.88	—	100.00	—	5.92	—	—	1.05
11/09 - 11/22/89	422.52	—	1.40	—	100.00	—	5.04	—	—	2.04
11/22 - 12/07/89	299.10	—	6.47	—	100.00	—	16.57	—	—	4.43
12/07 - 12/21/89	35.59	—	—	—	100.00	—	-28.19	—	—	37.60
12/21/89 - 01/04/90	20.40	—	—	—	100.00	—	-30.46	—	—	50.59

a) All values are relative activities on resin sample, except for ¹³⁷Cs, which is corrected for collection efficiency. (Note: This is also true for the 1987 data of Reference 2; however, the 1988 data of Reference 3 inadvertently used the relative activity of ¹³⁷Cs.)

startup in 1987. The sediment samples reflect many of the results observed for the river water. The underwater NaI(Tl) detector 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 ⁵⁸Co. Furthermore, the river concentrations for ⁵⁸Co (see Figure 1) continued to coincide with those for ⁶⁰Co (see Figure 2). This correlation results because Plant Vogtle supplies the bulk of the ⁶⁰Co, which is illustrated by ⁶⁰Co levels 0.3 mi above and 0.1 mi below the outfall (see Table 2).

In 1989, the ⁵⁸Co decreased noticeably, but the decrease in ⁶⁰Co was less pronounced. Neutron activations in the coolant piping creates both of those isotopes, but ⁵⁸Co (71 days) builds up toward saturation and decays more rapidly than ⁶⁰Co (five years). Therefore, ⁵⁸Co reduction is more pronounced than that for ⁶⁰Co during processing that retains the effluents longer before discharge to the river. Both isotopes had their

maximum concentrations just below Vogtle at the beginning of the year when concentrations of 3900 fCi/L ⁵⁸Co and 226 fCi/L ⁶⁰Co were measured.

Relative to ⁵⁸Co the river concentrations (Table 4) of most Vogtle-produced isotopes have not dramatically changed since these studies began in 1987. However, like ⁶⁰Co (five years), ⁵⁴Mn (313 days) has a long half-life relative to ⁵⁸Co (71 days); thus, its buildup/decay can result in increasing its relative activity. In fact from 1987 to 1989, the relative concentrations of both ⁵⁴Mn and ⁶⁰Co have increased by more than an order of magnitude. By contrast, such large variations are not seen in the relative activities of ⁵¹Cr, ⁵⁹Fe, ⁹⁵Nb, and ⁹⁵Zr, which have shorter half-lives and compare better to ⁵⁸Co. The ⁵⁷Co also has a relatively long half-life, but its detection was too rare for comparison.

SRS still appears to be the major source for the ³H and ¹³⁷Cs concentrations in the river. In 1989 the average ³H was 3.5 pCi/mL at Highway 301 bridge (see Table 5), which agrees

Table 5. Tritium Concentrations in 1989 (Values in fCi/mL)^a

Date	Shell Bluff	Vogtle Vicinity ^b			Hwy 301
		+0.3 mi	0.0 mi	-0.1 mi	
1/26/89	<1.3	4.2 ±0.9	4.7 ± 0.9	4.2 ±0.9	5.8 ± 0.9
2/16/89	<1.5	2.8 ±1.4	2.3 ± 1.1	<1.4	3.9 ± 1.1
3/16/89	<1.2	<1.7	<1.0	<1.1	3.9 ± 1.0
4/13/89	<1.5	<1.7	<1.8	<1.5	3.0 ± 1.0
5/16/89	<1.4	<1.2	<1.6	<2.0	3.2 ± 0.9
6/08/89	<1.0	<2.3	<1.3	<1.5	4.6 ±1.0
7/20/89	<1.4	<1.2	1.4 ± 1.4	4.2 ± 1.2	2.1 ± 0.6
8/17/89	<1.4	1.8 ±0.9	<1.7	<1.5	2.7 ± 0.9
9/28/89	<1.5	<1.1	<1.1	<1.0	2.5 ± 0.8
10/12/89	<1.8	3.3 ± 1.0	1.9 ± 0.9	2.6 ± 1.1	3.4 ± 1.0
11/22/89	<2.2	1.9 ± 0.9	1.5 ± 1.1	3.0 ± 1.0	5.0 ± 1.0
12/21/89	<1.5	<1.1	<1.7	<1.5	1.3 ± 0.6

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

b) Miles are measured upstream of Vogtle outfall.

well with the 3.0 pCi/mL measured prior to Vogtle startup; ³H measurements at other river locations remained at approximately 2 pCi/mL, which is also comparable to pre-startup values. A maximum concentration of 119 fCi/L ¹³⁷Cs was observed at Highway 301 bridge; this is comparable to the approximate 100 fCi/L average that is typically observed annually.

The ¹³⁷Cs concentrations near Vogtle are noticeably lower, averaging 10.1 ±1.2 fCi/L and 14.3 ±1.6 fCi/L for samples 0.3 mi above and 0.1 mi below the Vogtle outfall. This implies a ¹³⁷Cs contribution of 4.2 ±1.9 fCi/L from Vogtle and may signal that very small leaks in the reactor fuel are releasing trace fission products. Compared to the 1987 and 1988 yields of 1.1 ±1.3 fCi/L and 5.8 ±2.1 fCi/L respec-

Table 6. Sediment Concentrations in 1989^a

Isotope	Date	Vogtle Vicinity ^a			Hwy 301
		+0.3 mi	0.0 mi	-0.1 mi	
⁵⁴ Mn ^b	2/9/89	<4.2	<4.3	25.1 ±1.3	<4.7
	7/13/89	<4.0	<3.4	9.9 ±1.2	<6.6
⁵⁸ Co	2/9/89	<4.3	16.4 ±2.4	98.2 ±1.3	9.0 ±1.2
	7/13/89	<13	18.6 ±5.4	17.9 ±2.9	<13
⁶⁰ Co	2/9/89	<1.3	<1.5	6.9 ±0.7	2.3 ±0.7
	7/13/89	<1.3	<1.9	4.6 ±0.7	<1.6
¹³⁷ Cs	2/9/89	11.5 ±1.5	7.5 ±2.0	11.4 ±1.0	131 ±2
	7/13/89	11.4 ±1.5	7.7 ±1.6	19.5 ±1.1	148 ±3

a) Miles are measured upstream of Vogtle outfall.

b) ⁵⁴Mn values corrected for interference of 835.8 keV gamma of ²²⁸Ac.

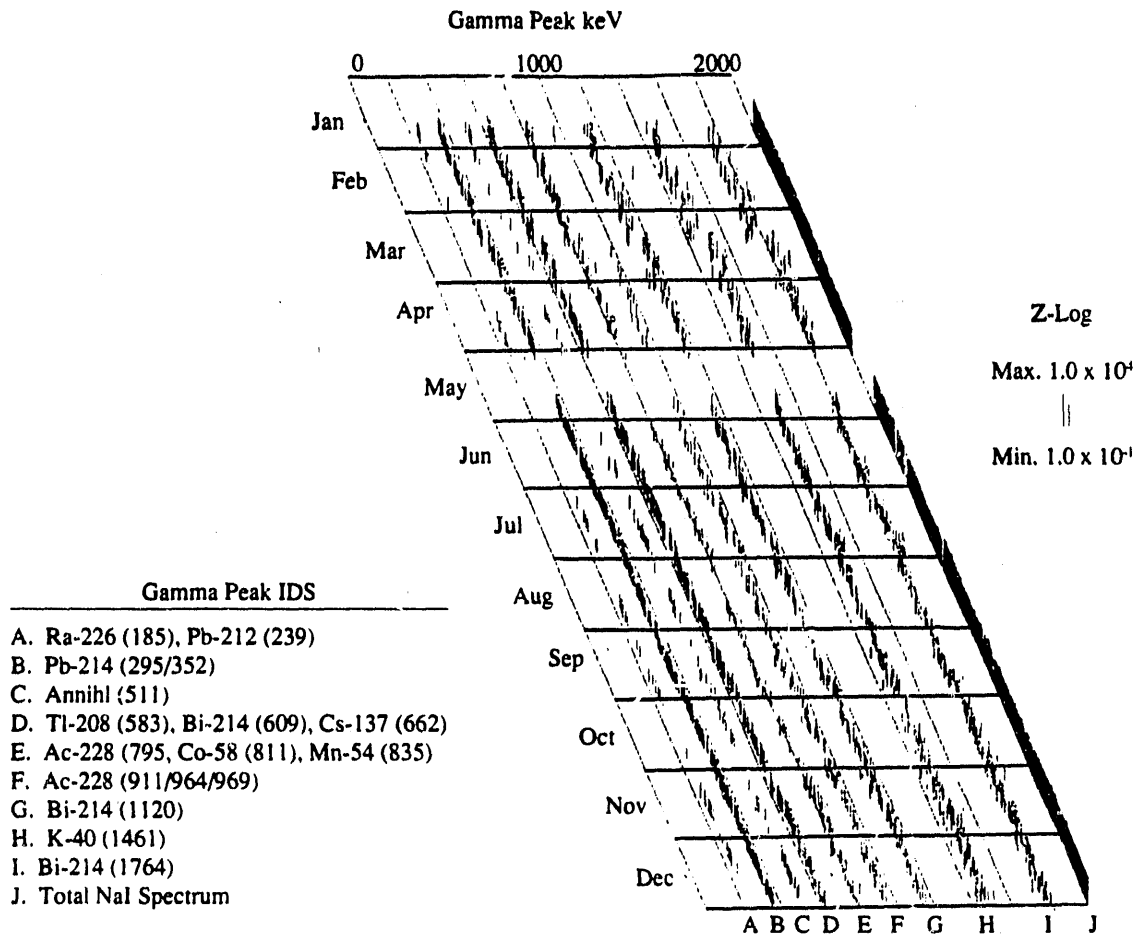


Figure 4. Results for Underwater NaI Detector at the Highway 301 Bridge

tively, the concentrations have noticeably increased. The continuation of these studies will further evaluate this trend.

The sediment samples contained detectable ^{54}Mn , ^{58}Co , ^{60}Co , and ^{137}Cs (see Table 6). This marks the first year that ^{54}Mn was reported, because the relatively long-lived isotope has accumulated significantly in the sediment. The HPGc analysis for ^{54}Mn requires that its 834.8 keV gamma peak be corrected for a 835.6 keV component from background ^{228}Ac . At 0.1 mi below the Vogtle outfall, the activities of ^{54}Mn , ^{58}Co , and ^{60}Co all decreased from the beginning of the year, reflecting the data of the river water samples; their respective maximum concentrations of 25, 98, and 6.9 fCi/g are the largest observed since Vogtle startup. The ^{137}Cs does not display these trends, and its values are comparable to those observed in earlier years.^{2,3}

The underwater NaI(Tl) detector at the Highway 301 bridge continues to illustrate that Plant Vogtle effluents can be monitored per detection of the ^{58}Co activity. An isometric

plot of count rate 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 . Figures 5 and 6 compare a time plot of this peak with a similar one for ^{58}Co concentrations measured with the resin samples. Both plots exhibit similar profiles, indicating good tracking; however, in the plot for 1987-1988³, the ratio (resin/underwater NaI) of the plots was approximately two times larger than that shown in figures 5 and 6. This difference is probably due to interference effects at the lower count rates, particularly the 795 keV and 835 keV gammas of ^{228}Ac and the 835 keV of ^{54}Mn that contribute to the peak near 800 keV. The ^{228}Ac is a natural background detected in sediment, and the long half-life of ^{54}Mn caused the build up relative to ^{58}Co . For the underwater NaI(Tl) detector, the 795, 811, and 835 keV peaks are not resolved because no resolution problem existed for the resin samples because they were counted on HPGc detectors. The underwater NaI(Tl) peak near 800 keV in 1989 is comprised of ^{54}Mn , ^{58}Co , and ^{228}Ac ; however, ^{58}Co dominated this generally larger peak in 1987-1988.

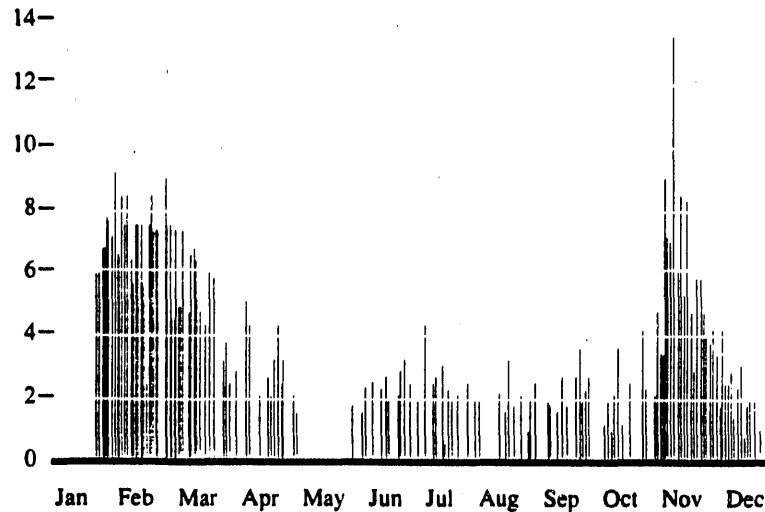


Figure 5. Results of Underwater Nal ⁵⁸Co (1989)

Table 7 compares the maximum-detected river concentrations with the DOE guide limits for drinking water.^{4,6} All concentrations are well below the guide limits. Furthermore, the table illustrates that maximum concentrations in 1989 are generally lower than those of 1987 and 1988. In particular, the short-lived Vogtle-produced isotopes ⁵¹Cr, ⁵⁸Co, ⁵⁹Fe, ⁹⁵Nb, and ⁹⁵Zr noticeably lowered. The longer lived isotopes of ³H, ⁵⁴Mn, ⁵⁷Co, and ¹³⁷Cs concentrations

correlated to those of earlier years; however, the ⁵⁴Mn concentration did exhibit a monotonic increase each year.

Summary

These studies have continuously monitored the Savannah River since their inception in 1986. They will continue on a routine basis, because they provide early detection of

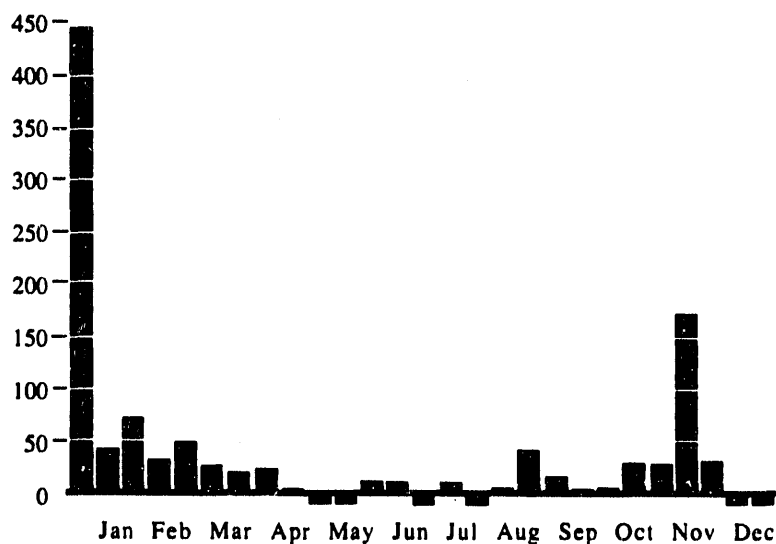


Figure 6. Results of Resin Samples for ⁵⁸Co at Highway 301 Bridge (1989)

Table 7. Comparison of Maximum 1989 Levels with DOE Guides for Drinking Water
(All Measured Levels are from Samples 0.1 mi. Downstream of Vogtle Outfall Unless Specified Otherwise.)

Isotope	Concentration (pCi/L)			
	1987 max	1988 max	1989 max	DOE Guide ^a
³ H	47,300 ^b	4200 ^c	5800 ^c	2,000,000
⁵¹ Cr	3.70	2.80	0.30	11,000,000
⁵⁴ Mn	0.06	0.40	0.61	50,000
⁵⁷ Co	0.02	0.02	0.01	100,000
⁵⁸ Co	16.8	15.5	3.90	40,000
⁵⁹ Fe	0.22	0.49	0.05	20,000
⁶⁰ Co	0.14	0.49	0.23	5,000
⁹⁵ Nb	0.10	0.50	0.10	40,000
⁹⁵ Zr	0.17	0.23	0.04	60,000
¹³⁷ Cs	0.10	0.39 ^c	0.12 ^c	3,000

- a) DOE 5484.1 (details per reference 6).
 b) Value at Vogtle outfall.
 c) Value at Highway 301 Bridge.

disturbing trends before actual health 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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Donald Hallman has continued to provide useful information on Plant Vogtle releases. C.D. Ouzts and H.T. Wilson coordinated the river sampling and analyzed the bulk of the samples.

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