

New York State Department of Environmental Conservation Division of Hazardous Waste Remediation 50 Wolf Road Albany, New York 12233-4015

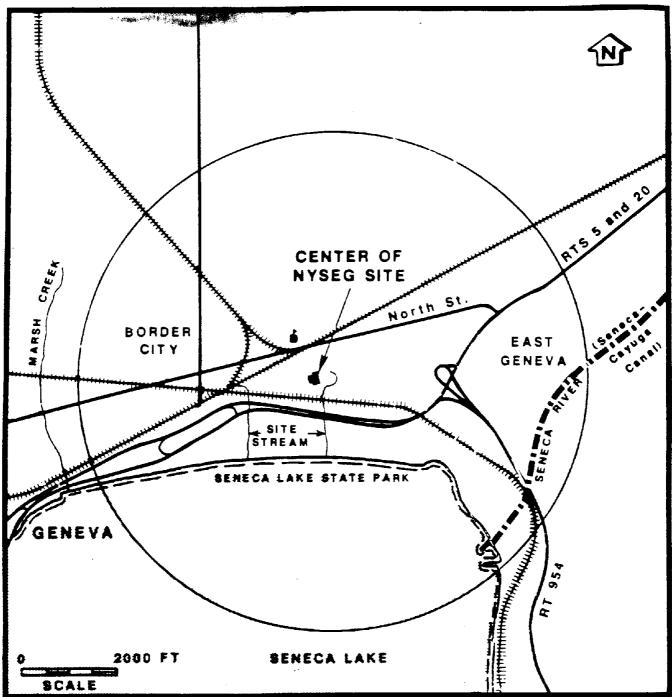
Add	
Modify	
Reclassify	X
Delist	

ADDITIONS/CHANGES TO REGISTRY OF INACTIVE HAZARDOUS WASTE DISPOSAL SITES

RECEIVED

JUN 4 1 1989

	HUREAU DE HAZARDOUS SEE CUREROL DIVISION OF HAZARDOUS
Site Name Former Geneva Coal Gasification Plant	DEC ID Numbers RESIDERION
Site Address North Street, Waterloo	County Seneca
Add New Site: (Potential hazardous waste site, Site Inspection Summa Report, EPA Preliminary Assessment Form and Registry Form must be completed and attached)	ry
Modify Registry data (detail below)	
χ Reclassify from class $2a$ to class 2 . (justify below)	
Delist (justify below)	
Detail/Justification	
The Geneva Coke Plant of New York State Electric & Gasolid wastes which were disposed of on-site. The disposal from the Ammonia Concentrate building and the By-Product & considered as hazardous waste disposal (6NYCRR Part 371 Salk K060, K087). Analysis of groundwater samples has indicated exceeded and standards for PAH's, total phenols, antimony, beryllic cyanide. Similar contaminants were also found at elevated water and stream sediments (refer enclosed letter from Miles).	l of sludge/tar generated building (Task 1 Report) is ection 371.4, EPA Hazardous Waste ences of regulatory guidelines um, iron, lead, sulfate and d levels in soils, surface
	e e e
Prepared by Manmohan Mehta D	late: June 15, 1989
Approvals:	14. 1/2
Reg. Haz. Waste Eng.	Date: 6/4/89
R. Tramontano NYSDOH	Date: 10/185
R. Dana/DEE Kichard / Dana [Date: 7/10/89
W. Demick/J. Swartwout/ Wird June 1	Date: 6/30/89
R. Marino	Date: 10/18/89
E. Barcomb (Date: 10/19/89



Base Map From USGS Topographic Map

(circle indicates 1 mile radius from site center)

Figure 3-1 Site Location

in soils where former gas plant structures (such as gas holders) had been located.

The predominant liquid waste generated at the Geneva site was quench water from the coking operations. Initially this water was discharged to a site stream. In 1923 a concrete-lined sludge basin was constructed and the water was pumped to the basin and allowed to separate. The supernatant was discharged to the site stream while the lower liquid layer was pumped into an 8-inch diameter, 336-foot deep injection well.

The environmental investigation has provided historical, geological, and hydrological information, as well as chemical data for ground water, stream water and sediment, lake sediment, soil, wastes, and air. Samples were analyzed for chemicals commonly found at coal gasification sites including polynuclear aromatic hydrocarbons (PAHs), volatile organic compounds, cyanides, non-chlorinated phenols and metals.

Specific "chemicals of interest" were selected for consideration in the risk assessment. Chemicals were included in the analysis if they were found at the site in elevated concentrations, have the potential for exerting acute or chronic health effects, and/or were present at levels exceeding established guidelines or standards. The assessment integrates two bodies of information for these chemicals: 1) site specific exposure analysis, and 2) health/environmental effects data. The latter information is taken from the available literature and is often summarized by regulatory agencies (primarily EPA) in the form of "potency factors" or "Acceptable Intake Chronic or Acute Values".

Various transport models are used to estimate exposure point concentrations from laboratory measurements of field samples. In this analysis nominal values are used where data are reported as less than a

presence of "blue billy" found coating some rocks near the site of the gas holder and purifier building.

The use of a deep injection well to dispose of coke quench water suggests that there may be coal tar constituents at depths greater than 200 feet, i.e., in the bedrock. Within a one mile radius of the site, there are no potable water wells in the bedrock formation.

Air quality impacts from volatile organic compounds appear to be minor and present minimal risk to on-site workers. The measured levels are at or below the benzene threshold limit value (TLV) of 10 ppm (Time Weighted Average)(Lederer, 1985) established by the American Conference of Industrial Hygienists. One high reading was detected in the transmission room of the natural gas compressor building, an area not frequented by employees. It is suspected that this reading was caused by minor natural gas leakage. Natural gas is a simple asphyxiant, dangerous in very high concentrations; however, no TLV has been established for this substance.

These previous investigations identified direct contact, ground water, surface water, stream sediments, and possibly fugitive dust from the former disposal area, as primary potential contaminant pathways. Fugitive dust was later eliminated as a possible pathway due to the moist nature of the site.

At the conclusion of Task 1 the spatial extent of the constituents in both the near surface and deeper levels was not known. In addition, the data collected were insufficient to establish the type(s) and quantity of the material disposed of and the extent of any plume that may exist because of that disposal.

The method by which the suspected tars entered the stream sediments also has not been determined. This information is important because a direct disposal route suggests that the level of contamination is not increasing

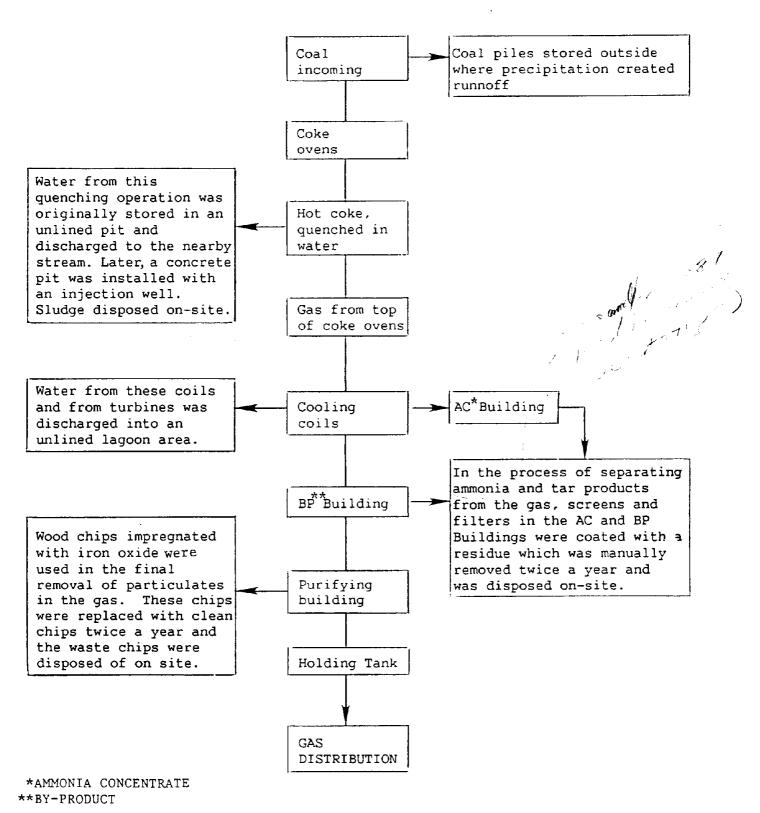


Figure 3-3 Waste Generation Flow Diagram Geneva Coal Gasification Plant

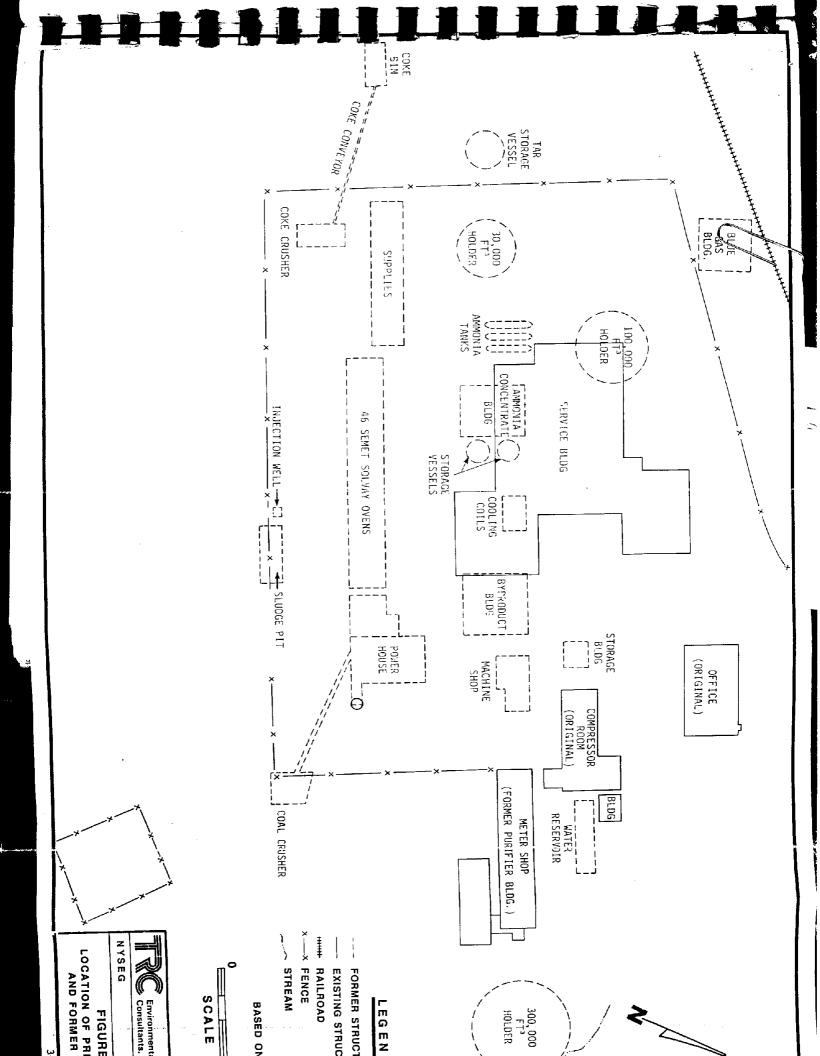


TABLE E-29 (Cont.)

GENEVA GROUND WATER SAMPLES

RESULTS FROM COMPUCHEM - SUMMARY OF RESULTS - ROUND 1

CONSTITUENT CONSTITUENT CONSTITUENT CONSTITUENT CONSTITUENT CONSTITUENT COMPOUNDS ACID EXTRACTABLES PHENOL 0.00 2,4-DIMETHYLPHENOL VOLATILE ORGANICS METHYLENE CHLORIDE C.05 CONORGANIC COMPOUNDS	DETECTION LIMIT MG/L	MG/L	MG/L
PHENOL 0.001 2,4-DIMETHYLPHENOL VOLATILE ORGANICS METHYLENE CHLORIDE 0.05			
PHENOL 0.001 2,4-DIMETHYLPHENOL VOLATILE ORGANICS METHYLENE CHLORIDE 0.05			
2,4-DIMETHYLPHENOL VOLATILE ORGANICS METHYLENE CHLORIDE 0.05			
	0.010 0.010	0.036 0.011	N D N D
NORGANIC COMPOUNDS	0.010	ND	0.010
METALS			
MERCURY, TOTAL 0.002 ZINC, TOTAL 5.0	0.00020 0.020	0.00032 0.080	0.0070 0.16
CYANIDES			
total cyanide 0.2	0.010	ND	1.8

TABLE E-30 (Cont.)

GENEVA GROUND WATER SAMPLES

RESULTS FROM COMPUCHEM - ROUND 2

			AMPLE ID DATE MPLE TYPE	MW-3S 5/1/86 GRAB	MW-3D 5/1/86 GRAB
CONSTITUT		DI STAN- - DARO	ETECTION LIMIT MG/L	MG/L	MG/L
INORGANIC COMPOUNDS					
METALS	5				
	ANTIMONY, TOTAL ARSENIC, TOTAL BERYLLIUM, TOTAL CADMIUM, TOTAL CHROMIUM, TOTAL COPPER, TOTAL LEAD, TOTAL MERCURY, TOTAL NICKEL, TOTAL SELENIUM, TOTAL SILVER, TOTAL THALLIUM, TOTAL ZINC, TOTAL	0.025 AL 0.025 0.004	0.050 0.020 0.010 0.050 0.10 0.050	ND 0.075 ND ND ND 0.16 0.00038 ND ND ND ND	ND N
PHENOLS	5				
	TOTAL PHENOI	10000	0.010	0.034	N D
CYANIDI	ES .				
	TOTAL CYANIDE	E 0.2	0.010	0.075	ND

TABLE E-31 (Cont.)

GENEVA GROUND WATER SAMPLES

RESULTS FROM COMPUCHEM - ROUND 3

	SAMPLE ID DATE SAMPLE TYPE	MW-2S 8/7/86 GRAB	MW-2D 8/7/86 GRAB
CONSTITUENT STAND		MG/L	MG/L
VOLATILE ORGANICS			
CHLOROMETHANE	0.313	ND	ND
VINYL CHLORIDE	0.010	ND	ND
CHLORCETHANE	0.010	ND	ND
BROMOMETHANE	0.810	ND	ND
ACROLEIN	0.100	ND	ND
ACRYLONITRILE	0.100	ND	ND
METHYLENE CHLORIDE $D\cdot l$) 5 0.010	0.011*	0.014
1,1-DICHLOROETHYLENE	0.010	ND	ND
1,1-DICHLORGETHANE	0.313	ND	ND
TRANS-1,2-DICHLOROETHYLENE	0.010	ND	ND
CHLOROFORM	0.010	ND	ND
1,2-DICHLOROETHANE	0.010	ND	ND
1,1,1-TRICHLOROETHANE	0.010	ND	ND
CARBON TETRACHLORIDE	0.010	ND	ND
BROMODICHLOROMETHANE	0.010	DИ	ИĎ
1,2-DICHLOROPROPANE	0.010	ND	ND
TRANS-1,3-DICHLOROPROPENE	0.010	ND	ND
TRICHLOROETHYLENE O. 6	0.010	0.021	ND
BENZENE	0.010	ND	ND
CIS-1,3-DICHLOROPROPENE	0.010	ND	ND
1,1,2-TRICHLOROETHANE	0.010	ND	ND
DIBROMOCHLOROMETHANE	0.010	ND	ND
BROMOFORM	0.010	ND	ND
1,1,2,2-TETRACHLOROETHYLENE	0.010	ND	ND
1,1,2,2-TETRACHLOROTHANE	0.010	ND	ND
TOLUENE	0.010	ND	ND
CHLOROBENZENE	0.010	ND	ND
ETHYLBENZENE	0.010	ND	ND
2-CHLOROETHYL VINYL ETHER	0.010	ND	ND

^{*} Detected in method blank in a concentration greater than 1/2 the detection limit and greater than 1/2 the sample concentration.

GENEVA GROUND WATER SAMPLES

RESULTS FROM COMPUCHEM - ROUND 3

	** · 8	SAMPLE ID DATE SAMPLE TYPE	MW-2S 8/7/86 GRAB	MW-2D 8/7/86 GRAB	MW-2S 8/7/86 GRAB	MW-2D 8/7/86 GRAB
CONSTITUENT	STAN PARA	DETECTION LIMIT STAN - MG/L DAR-D	(FILTERED MG/L	SAMPLES) MG/L	(UNFILTERED MG/L	SAMPLES) MG/L
INORGANIC COMPOUNDS METALS			DISSOLVED	LVED	TOTAL	Ţ.
	ANTIMONY	0.050	Q.	QN Q	QN	Q
	ARSENIC 0.02	5 0.050	Q	QN ON	0.052	Q.
	BERYLLIUM	0.020	QN	ND QN	QN	QN
	CADMIUM 0.01	0.010	0.027	0.013	0.035	0.014
	CHROMIUM	0.050	ND	QV QV	0.069	Q
	COPPER 1.0	0.10	QN QN	CN CN	QN	Q
	LEAD 6.02	5 0.050	QN Q	Ð	ON	Q
	MERCURY C. DO	0.00020	S.	QX	0.0047	QN O
	NICKEL	0.10	0.110	QN	0.19	QN
	SELENIUM	0.010	Ø	QN	Ð	QN
	SILVER	0.050	QV	QN	Q	ND
	THALLIUM	0.050	Q	Q.	<u>R</u>	QN
	ZINC 0.020	0.020	0.050	0.065	0.24	0.048

M

TABLE E-31 (Cont.)

GENEVA GROUND WATER SAMPLES

RESULTS FROM COMPUCHEM - ROUND 3

	S	AMPLE ID DATE	MW-2S 8/7/86	MW-2D 8/7/86
	SA	MPLE TYPE	GRAB	GRAB
CONSTITUENT	_	ETECTION LIMIT		
	STAN DAG	² ø ^{MG/L}	MG/L	MG/L
PHENOLS				
	TOTAL PHENOL 0.001	0.010	0.028	ND
CYANIDES				
	TOTAL CYANIDE 0 12	0.310	0.016	ND



New York State Department of Environmental Conservation

MEMORANDUM

JAM CENTRY

TO: FROM: SUBJECT: Raymond Lupe, Acting Supervisor, Western Investigation Sec Sion!D V:A: E Michael Ryan // / D.E.C. REG. ##

NYSEG - Former Geneva Coal Gasification Plant, Senecace)

DATE:

December 22, 1987

The above-referenced site was investigated as part of an ongoing site study by TRC Environmental Consultants. TRC is investigating the presence of coal gasification process residues at this New York State Electric and Gas Corporation (NYSEG) Facility.

This site is the former location of the Empire Coke Company. Though the controlling interest changed hands in later years, coal gas production at this facility continued until 1934. During its years of operation (1903-1934) the facility consisted of as many as 46 coke ovens, two large gas receivers and a blue gas operation with a holder. The property is currently maintained by NYSEG as a utility substation.

The existing problems are a result of the waste generated by the former operations at this site, specifically the disposal of that waste.

It is reported that both solid and liquid wastes were disposed of onsite. The solid wastes included iron oxide impregnated wood shavings from the purification process and tars. The predominant liquid waste generated was waste water from the coke quenching operation. In the early years of operation the liquid waste was discharged directly into the site stream. This process was altered in 1923 with the construction of a sludge basin. The sludge basin allowed separation of the waste water supernatent and sludge. The supernatent was discharged to the stream while the lower liquid layer was disposed via an 8 inch diameter, 336 foot deep injection well (Task II Report, pg. 3, pg. 24). Other liquid wastes disposed of on site were from random tar spillage, drip boxes or gas lines to collect condensed tar within the system.

As a result of the on-site waste disposal, laboratory analysis has showed contamination of soil, stream sediments, groundwater, and surface water at this site.

Analysis of the soil and stream sediments has shown elevated concentrations of PAH's, purgeable aromatics, phenols, iron, zinc, sulfate, organic compounds (nitrogen) and cyanides. Analysis of groundwater samples demonstrated levels exceeding regulatory guidelines and standards for the following: PAH's, total phenols, antimony, beryllium, iron, lead, sulfate and cyanide (see table). Site surface waters showed elevated levels of PAH's, iron, zinc, copper, lead, phenols and organic nitrogen.

The above noted problems illustrate the need for remedial activity at this site. A site of a class 2 site and should be upgraded accordingly.

Other recommendations and comments:

Prior to any remedial action one or more monitoring wells should be installed in the <u>vicinity of the former injection well</u>. This monitoring well(s) should be at a depth which would allow an accurate assessment of any contamination resulting from the injection well.

Risk assessment is a topic which requires the cooperation of the Health Department. It is suggested that before any risk assessment is completed, the Health Department be consulted and therefore an accurate assessment may be made.

bcc: F. Shattuck

M. Mehta

R. Tramontano

K. Gupta M. Ryan file

MR:kr

SUBSTANCE	STANDARD (ug/l)	MONITORING WELL CONTAMINANT RANGE (ug/1)					
	3	MH-1S MW-1D		MW-2S	MH-5D	MW-35	MW-3D
		.	ROUNDS 1 - 3	*##			
Benzo (a) pyrene	ND	0-0.5	0-1.2	B.E-0	0-0.4		
Iron	300			1,620-2,580		460-1,940	
Sulfate	250,000			535,000-1,180,000	244,000-274,000	1,050,000-1,340,000	0-436,000
Cyanide	200	1		3,200-5,700		0-970	
Phenols (total)	1				0-5.2		
		***	- ROUND 4	***			
Benzo (a) pyrene	CM		1,021,0	8-28			
Antimony	11.5			65- 8 1		64	3 5
Beryllium				3.5	1.4	3.5	00
Iron	300			6,230-6,600	•••	3,160	48-9,800
_ead	25			B,E30 B,000		0,100	0-50
Sulfate	250,000			1,100,000-1,200,000	280,000	1,100,000	320,000
Cyanide	200			1,900-2,100	200,000	1,100,000	520,000
		10	10	12-21	12	10	10
Phenols (total)	1	10	10	15-51	16	10	10

.

-