KERR-MCGEE CHEMICAL CORPORATION

POST OFFICE BOX 55 . HENDERSON, NEVADA 89015

October 26, 1982

RECEIVED OCT 2 7 1982

CERTIFIED MAIL NO. P26 0233713

Mr. Bill Wilson, Chief Technical Assessment Section Toxics and Waste Programs Branch U. S. Environmental Protection Agency Region IX 215 Fremont Street San Francisco, CA 94105

Re: Second Quarter 1982 RCRA Groundwater Monitoring Data, Henderson, Nevada Facility, Kerr-McGee Chemical Corporation, EPA ID No. NVD 008290330

Dear Mr. Wilson:

Please find attached in Tables I (landfill) and II (surface impoundments) completed second quarter 1982 RCRA groundwater monitoring data for the Kerr-McGee Chemical Corporation facility at Henderson, Nevada. These data supplement the partial data submitted on August 17, 1982. This data summary is supplied in accordance with 40 CFR 265.94(2)(i).

The cadmium concentration in all reported wells exceeds the NIPDWS limit of 0.01 mg/L. The turbidity limit of 1 NTU is also exceeded in all wells. Chromium exceeds the NIPDWS limit of .05 mg/L in the surface impoundment wells M-1, M-2, and M-3. Nitrate limits of 10 mg/L as N are exceeded in wells M-1, M-3, and M-4. The NIPDWS silver limit of 0.05 mg/L is exceeded in wells M-2 and M-3. In the landfill wells, the NIPDWS limit of 1.4-2.4 mg/L for fluoride is exceeded in M-5, M-6, and M-7. Coliform NIPDWS limits of 1/100 mL are exceeded in wells M-5, M-6, and M-7.

Sincerely, Machur

Ŕ. B. Chase Plant Manager

RBC:jc Attachment

cc: H. L. Rosse-Carson City

TABLE I

SECOND QUARTER 1982 RCRA GROUNDWATER MONITORING DATA KERR-MCGEE CHEMICAL CORPORATION - HENDERSON FACILITY

		Upgradient Wel	Ll Do	wngradient	Wells ·
Paramete:	r	M-5	M-6	M-7	H-28
Arsenic	(mg/L)	0.02	<0.01	<0.01	0.01
Barium	**	0.10	0.08	0.12	0.09
Cadmium	ŤŤ	0.02	0.02	0.02	0.02
Chromium	**	0.01	0.01	<0.01	0.01
Fluoride	TI	3.6	2.5	2.5	1.0
Lead	11	<0.01	0.02	<0.01	<0.01
Mercury	11	<0.001	<0.001	<0.001	0.012
Nitrate (as N)	11	0.45	0.22	0.22	0.2
Selenium	*1	<0.005	<0.005	<0.005	<0.005
Silver	11	0.02	0.01	0.01	0.02
Endrin	**	<0.0002	<0.0002	<0.0002	<0.000
lindane	**	<0.004	<0.004	<0.004	<0.004
lethoxychlor	11	<0.1	<0.1`	<0.1	<0.1
ſoxaphene	11	<0.005	<0.005	<0.005	<0.005
2,4-D	**	<0.1	<0.1	<0.1	<0.1
2,4,5-TP Silve	x "	<0.01	<0.01	<0.01	<0.01
Gross Alpha	(pCi/L)	<40	<20	<40	< 4 0
Gross Beta	(pCi/L)	40 <u>+</u> 30	40 <u>+</u> 20	60 <u>+</u> 20	50±20
Radium	(pCi/L)	<2	<2	< 2	< 2
Coliform	(MPN/100 mL)	240	93	4	<2.2
Turbidity	(NTII)	2000	2000	2000	35

LANDFILL

Analysis by Truesdail Laboratories, Inc., Los Angeles, CA.

TABLE II

SECOND QUARTER 1982 RCRA GROUNDWATER MONITORING DATA KERR-MCGEE CHEMICAL CORPORATION - HENDERSON FACILITY

·		Upgradient Well	Dov	wngradient	Wells
Paramete	r	M-1	<u>M-2</u>	M-3*	M-4*
Arsenic	(mg/L)	<0.01	0.03	<0.01	<0.01
Barium	23	0.18	0.23	0.27	0.09
Cadmium	11 ·	0.03	0.05	0.06	0.01
Chromium	11	12.9	10.0	44	0.01
Fluoride	11	0.7	1.0	0.7	1.0
Lead	11	<0.01	<0.01	<0.01	<0.01
Mercury	TI	<0.001	<0.001	<0.001	<0.001
Nitrate (as N)	TT	11.2	8.8	44	11.1
Selenium		<0.005	<0.005	<0.005	<0.005
Silver	**	0.04	0.10	0.09	0.03
Endrin	11	<0.0002	<0.0002	<0.0002	<0.000
Lindane	11	<0.004	<0.004	<0.004	<0.004
Methoxychlor	"	<0.1	<0.1	<0.1	<0.1
Toxaphene	*1	<0.005	<0.005	<0.005	<0.005
2,4-D	11	<0.1	·<0.1	<0.1	<0.1
2,4,5-TP Silve:	x "	<0.01	<0.01	<0.01	<0.01
Gross Alpha	(pCi/L)	<100	<100	<200	<40
Gross Beta	(pCi/L)	80±20	720±40	300 ±40	30 ±20
Radium	(pCi/L)	<2	<2	<2	<2
Coliform	(MPN/100 mL)	<2.2	<2.2	<2.2	<2.2
Turbidity	(NTU)	45	57	2.5	85

SURFACE IMPOUNDMENTS

Analysis by Truesdail Laboratories, Inc., Los Angeles, CA.

* Monitor wells M-3 and M-4 were replaced by wells M-8 and M-9 following second quarter analysis.

-							EPA T or Gon	SD Fac. Na. N.	V D 0	0 8 2 9	0 3 3 0
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orameter Units	Cround Watte Eler. Fr. Sampia Occurvice	pH Standard	Conductivity Jamhos mg/l	Total Organic Carbon mg/l	Totel Organic Helogan moli	Chlarida nia/l	lron mg/l	Manganese mg/l	Phanols mg/l	Sodium mg/l	Sulfate mg/l
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il reat true	nitial) Background		•	•							
14/32	1751.63	7.3	11,350	4.3	528	1450	0.27	0.07	0.01	1260	868
1/82	1750.51	7.4	12,250	18.3	51.5	-1625	0.16	0:04	<0.01	1085	1190
./5/82	1750.72	7.3	11,600	21.3	615	2000	0.40	0.09	<0,01	1170	1120
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Units	Arsenic mg/l	Barium mg/l	Cadmium , mg/l	Chromium mg/l	Fluorida mg/l	Lead mg/l	Mercury mg/l	Nitrate mg/l (as N)	Selenium mg/l	Silver mg/l	
14/82	0.01	0.21	0.03	12.2	1.0	<0.01	0,001	0.91	<0.005	0,04	-
/1/82	<0.01	0.18	0.03	12.9	0.7	<0.01	<0.001	11.2	<0,005	0.04	
:/5/82	<0.01	0.20	0.03	12.7.	1 • 0 • 1	<0.01	<0.001	18.7	<0.005	0.02	
Units	Endrin mg/l	Lindane . Mg/I	Methoxychlor .mg/l	Toxaphene mg/l	2,4.D mg/l	24.5-79 Filver MgA	Radlum pCI/I	Gross Atpha pCI/I	Gross Beta pC1/1	Coldorn B science 1/100 ml	
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/14/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<15	<20	<2.2	
1/82	<0.0002	<0.004	<0,1	<0.005	<0.1	<0.01	<2	<100	80±20	<2.2	
0/5/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<50	50±30	<2.2	
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Farameter Units	Ground Water Elev. FL: Sample Occurance	pH Standard	Conductivity Junhos mg/l	Total Organic Carbon mg/l	Youl Organic Halogan Mall	Chloride ma/i	Iron mg/l	Manganese mg/l	Phonois mg/i	Sodium mg/l	Sulfate mg/i
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124,/82	1746.77	7.3	20,000	5	870	3800	0,03	0.04	<0.01	4430	2900
/1/82	1744.40	7.2	34,000	6	23	7210	0.27	0.05	<0.01	6350	4180
0/5/82	1746.35	7.3	22,000	45	29	4750	0.34	0,07	<0.01	4190	1190
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Parameter Units	Arsenic mg/l	Barium mg/l	Cadmium , mg/l	Chromium mg/l	Fluorido mg/l	Lead mg/l	Mercury mg/l	Nitrato mg/l (ag N)	Setenium rng/l	Silver mg/l	
21										•	
14/82	0.02	0.18	0.04	9.0	0.95	<0.01	<0.001	0.45	<0.005	0.05	
/1/82	0.03	0.23	0.05	10.0	1.0	<0.01	<0.001	8.8	<0.005	0.10	
./5/82	0.02	0.13	0.04	9,15	2.0	<0.01	<0.001	14.7	<0.005	0.03	
Units	Endrin mg/l	Lindane 1/0m	Methoxychlor Ng/I	Toxaphane mg/l	2,4-D mg/l	2.4.6-70 Silvy MpA	Radlum pCI/I	Gross Alpha pCI/I	Gross Beta pC1/1	Colulorm Bectria 1/100 ml	
-											
/14/82	<0.0002	<0.004	<0.1	<0.005	<0,1	<0.01	<3	<15	<20	<2.2	
/2/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<100	720±40	<2.2	
:/5/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	-09>	300+100	<2.2	
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0./5/82	0.01	0.23	0.02	5.1	2.0	<0.01	<0.001	14.7	<0.005	0,02	
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5/16/82	1716.49	6.5	10,500	126	35	2750	22.19	8.94	0.17	1010	2350
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Parameter Units	Arscnic mg/l	Barium mg/l	Cadmium , mg/l	Chromium mg/l	Fluorido mg/l	Lead I/gm	Marcury mg/l	Nitrato mg/l (AS N)	Selonium mg/l	Silver mg/l	
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5/16/82	0.02	0.10	0.02	0.01	3.6	<0.01	<0.001	0.45	<0.005	0.02	
-0/5/82	<0.01	0.15	0.02	0.02	2.0	<0.01	<0.001	0.10	<0.005	0.02	
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Parometer Units	Endrin mg/l	Lindane mg/l	Methoxychlor .mg/l	Toxephene mg/l	2,4-D mg/l	2.4:8-7P Silver Aga	Radium pCI/I	Gross Alpha pCUI	Gross Bata pC1/1	Coliform 8 ktiviu 1/100 ml	
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./16/82	1697.82	6.6	9600	47	8	2270	9.73	6.51	0.06	1060	2550
0/5/82	1697.57	6.4	9500	80	12	2370	15.9	2.88	<0.01	1150	1170
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Parameter Units	Arsenic mg/l	Barium mg/l	Cadmium ng/l	Chromium ng/l	Fluorido mg/l	Lead mg/l	Mercury mg/i	Nitrate mg/i (as N)	Selenium mg/l	Silver Ing/l	
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/16/82	<0.01	0.08	0.02	0.01	2.5	0.02	<0.001	0.22	<0.005	0.01	
/5/82	<0.01	0.10	0,02	0.02	1.0	<0.01	0.001	0.10	<0.005	0,03	
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L'units	Endrin mg/l	Lindane mg/l	Methoxychlor mg/l	Toxephene mo/l	2,4-D mg/l	2.4:5-7.0 Silvar 2.4:20	Radium pCi/l	Gross Alpha pCi/l	Gross Beta pC1/1	history 1/100	
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16/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<20	40±20	93	
1/5/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<50	300±100	<2.2	
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Parameter Units	Ground Water Elev. Fl. Sumple Decurence	pH Standard	Conductivity Jumhoi ing/i	Yotel Organic Carbon mg/l	Tatal Drazhie Halogan Nan	Chloride ma/l	lron mg/l	Manganese mg/l	Phenols mg/l	Sadium mg/l	sulfato 1/gin
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c/16/82	1701.83	6.8	10,000	50	6 ·	2300	4.87	4.28	0.08	1180	2500
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Parameter Units	Arsenic mg/l	Barium mg/l	Cadmium mg/l	Chromium mg/l	Fluoride mg/l	Lead mg/l	Mercury mg/l	Nitrate mg/i (as N)	Selenium mg/l	Silver mg/l	
e.									-		
/16/82	<0.01	0.12	0.02	<0.01	2.5	<0.01	<0.001	0.22	<0.005	0.01	
0/5/82	<0.01	0,18	0.02	<0.01	1.0	<0.01	0.001	0.10	<0.005	0.02	
-stameter L'aire	Endrin moli	Lindane 2007	Methoxychlor mo/t	Toxaphene ma/l	2,4.D ma/l	2.4:579 Bilogy	Radium	Gross Alpha aCl/i	Gross Bata	Coldora Bactura	
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¢.							-	· · · ·			
/16/82	-:0,0002	<0.004	<0.1	<0.005	<0,1	<0,01	<2	<40	60±20	4	-
0/5/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<20	<40	16	
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Parameter Duits	Ground Nater Elev. Ft. Sample Occurence	pH Standard	Conductivity Lanhos mg/l	Yatel Organic Carbon mg/l	Yatal Organ's Hulogen mo/l	Chlorida ma/l	lron mg/l	Manganose mg/l	Phenols mg/l	Sodium mg/l	Sulfato mg/I
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First Year lit	itial) Background		:	•		•					
10/97	1699 81	7.0	10.500	3	23	2130	0,21	2.42	<0.01	1680	730
70/6/-	TD.0071	2 4	800	18	7.9	2140	2.28	2.16	<0.01	1195	2110
01=100	1608 33	(·)	000.6	, 4	11	2250	0,61	1.77	<0.01	0111	0611
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15/S2	0.02	0.08	0.02	0.02	2.0	<0.01	<0.001	0.53	<0.005	0.01	
a erameter Units	E narin mg/l	Lindane mg/l	Methoxychlor mg/l	Toxapheno ng/l	2,4-D mg/l	2.4.5.7 P Salvoy Mun	Redium pCi/l	Gross Alpha pCl/l	Gross Beta pC1/1	Coliform Buciria 1/103 ml	
						-					
(5/0/0)	<0.0002	<0.004	<0.1	<0.005	<0.1	<0,01	<2	81	<40	<2.2	
65/ ty	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<40	50+20	<2.2	
15/82	<0.0002	<0.004	<0.1	<0.005	<0.1	<0.01	<2	<40	60+40	<2.2	
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KERR-MCGEE CHEMICAL CORPORATION

POST OFFICE BOX 55 . HENDERSON, NEVADA 89015

August 6, 1982

RECEIVED

AUG 9 - 1982

ENVIRONMENTAL PROTECTION

CERTIFIED MAIL NO. P26 0233706

Mr. H. Laverne Rosse Program Director - Waste Management State of Nevada Division of Environmental Protection Capitol Complex Carson City, NV 89710

Dear Mr. Rosse:

Enclosed is Kerr-McGee Chemical Corporation's first quarter 1982 RCRA groundwater monitoring data. Through an oversight, this report was not mailed to you when it was issued.

Sincerely,

C. B. Armstrong Plant Manager

CBA:jc Enclosure

KERR-MCGEE CHEMICAL CORPORATION

POST OFFICE BOX 55 . HENDERSON, NEVADA 89015

May 14, 1982

CERTIFIED MAIL - NO. P26 0233534 RETURN RECEIPT REQUESTED

Mr. Bill Wilson, Chief Technical Assessment Section Toxics and Waste Programs Branch U. S. Environmental Protection Agency Region IX 215 Fremont Street San Francisco, CA 94105

Re: First Quarter 1982 RCRA Groundwater Monitoring Data

Dear Mr. Wilson:

Please find attached in Table I, first quarter 1982 RCRA groundwater monitoring data for the Kerr-McGee Chemical Corporation facility in Henderson, Nevada. This data summary is supplied in accordance with 40 CFR 265.94(2)(i).

The arsenic concentration in downgradient well H-28 exceeds the EPA National Interim Primary Drinking Water Standard (NIPDWS) of 0.05 mg/l. The NIPDWS level of 0.01 for cadmium is exceeded in all wells except downgradient well M-4, while the NIPDWS level for chromium of 0.05 mg/l is exceeded in all wells except downgradient well H-28. The silver concentration in downgradient well M-3 also exceeds the NIPDWS level of 0.05 mg/l.

Additionally, the NIPDWS gross alpha level of 15 PCI/l is exceeded in well H-28. At this time we are investigating possible causes for these exceedances. No other values exceed NIPDWS levels.

Sincerely,

C. B. Armstrong Plant Manager

CBA:jc Attachment

cc: H. L. Rosse - Certified Mail No. P26 0233706 Division of Environmental Protection Carson City, NV 89710

TABLE I

FIRST QUARTER 1982 RCRA GROUNDWATER MONITORING DATA DRINKING WATER PARAMETERS

KERR-McGEE CHEMICAL CORPORATION

Henderson, Nevada Facility

	Upgradient		Downgradie	ent Wells	
Parameter	<u>M-1</u>	M-2	<u>M-3</u>	<u>M-4</u>	H-28
Arsenic (mg/1)	0.01	0.02	0.02	<0.01	0.47
Barium	0.21	0.18	0.37	<0.12	0.08
Cadmin	0.03	0.04	0.06	0.01	0.02
Chromium	/ 12,2 [*] /	9.0	31.1	0.18	0.03
Fluoride	1.00	0.95	1.00	1.00	0.55
Logd	<0.01	<0.01	<0.01	<0.01	<0.01
Morgury	0.001	<0.001	0.001	0.001	<0.001
Nitrate (as N)	0.91	0.45	<0.10	0.80	<0.10
Solonium	<0.005	<0.005	<0.005	<0.005	<0.005
Selection	0.04	0.05	0.06	0.02	0.02
Silver	<0.0002	<0.0002	<0.0002	<0.0002	<0.0002
	<0.004	<0.004	<0.004	<0.004	<0.004
Lindane	<0.01	<0.1	<0.1	<0.1	<0.1
Methoxychiol	<0.005	<0.005	<0.005	<0.005	<0.005
Toxaphene	<0.10	<0.10	<0.10	<0.10	<0.10
	<0.01	<0.01	<0.01	<0.01	<0.01
2, 4, 5-1P Silvex	<2	<3	<2	<2	<2
Total Radium (PCI/I)	~15	<15	<15	<15	81
Gross Alpha (PCI/I)	<20	<20	<20	<20	<40
Gross Beta (PCI/1)	<20	<2.2	 <2 2	<2.2	<2.2
Coliform Bacteria (#/100 ml)	<2.2	73	7 3	7.6	7.0
pH	7.3, 7.3		1.5	,	•••

' BR-MCGEE CHEMICAL CORPORATION

Henderson Facility

RCRA Monitor Well No.M-5 Well Construction and Completion Table

Date Started	June 1, 1982
Date Completed	June 3, 1982 (except for surface grout)
Location	Approx. 100' South of landfill
Elevation from Top of Well Cover	1747.83'
Drilling Method	Rotary rig
Drilling Fluid	Revert
Depth to Muddy Creek	31'.
Total Depth of Well	43'
Borchole Diameter	9 5/8"
Well Casing Diameter/Type	5" ID/threaded steel pipe
Well Casing Interval	39' to surface
Perforated Interval	39' - 29'
Perforation Type/ Size /Open Area	Factory slot/ 1/8" x 2" /4 in ² per 10'
Casing Above Ground (Well Cover)	Approx. 15"
Gravel Pack Interval	43' - 28'
Type of Gravel	1/8" - 3/8" pea gravel
Surface Seal Interval	22' to surface (grout)
Completion:	022.0' grout 22.0'-24.0' well cuttings 24.0'-28.0' bentonite pellets 28.0'-43.0' gravel pack
Comments: Open hole with gravel from 43.0' to 39.0'	
Steel 6-inch well cover with cap cemented in place 6-16-82	

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TORR-MCGEE CHEMICAL CORPORATION

Henderson Facility

RCRA Monitor Well No. M-6 Well Construction and Completion Table

Date Started	June 2, 1982
Date Completed	June 3, 1982 (except for surface grout)
Location	NW corner of landfill
Elevation from Top of Well Cover	1729.17'
Drilling Mathod	Rotary rig
Drilling Fluid	Revert
Depth to Muddy Creek	32'
Total Depth of Well	43'
Borehole Diameter	.9 5/8"
Nell Casing Diameter/Type	5" ID/threaded steel pipe
Well Casing Interval	<u>34' to surface</u>
Perforate: Interval	35' - 25'
Perforation Type/ Size /Open Area	Factory slot/ 1/8" x 2" / 4 in ² per 10'
Casing Above Ground (Well Cover)	Approx. 15"
Gravel Pack Interval	43' -22'
Type of Gravel	1/8" - 3/8" pea gravel
Surface Seal Interval	15' to surface
Completion:	0 -15.0' grout 15.0'-18.0' well cuttings 18.0'-22.0' bentonite pellets 22.0'-43.0' gravel pack
Comments: Open hole with gravel from 43.0' to 35.0'.	
Steel well cover with cap cemented in place on 6-16-82.	

 $\sum_{i=1}^{n}$

KERR-ECGEE CHEMICAL CORPORATION

Henderson Facility

RCRA Monitor Well No. M-7 Well Construction and Completion Table

Date Started	June 3, 1982
Date Completed	June 3, 1982 (except for surface grout)
Location	Approx. 120' east of M-6
Elevation from Top of Well Cover	1729.83'
Drilling Method	Rotary rig
Drilling Fluid	Revert
Depth to Muddy Creek	29.5'
Total Dapth of Well	37'
Borchole Diameter	9 5/8"
Well Casing Diameter/Type	<u>5" ID/threaded steel pipe</u>
Well Casing Interval	34' to surface
Ferforated Interval	351 - 251
Perforation Type/ Size /Open Area	Factory slot / 1/8" x 2" /4 in ² per 10'
Casing Above Ground (Well Cover)	Approx. 15"
Gravel Pack Interval	37' - 22'
Type of Gravel	1/8" - 3/8" pea gravel
Surface Seal Interval	15' to surface
Completion	0 -15.0' grout 15.0'-18.0' well cuttings 18.0'-22.0' bentonite pellets 22.0'-37.0' gravel pack
Comments: Open hole with gravel from 35.0' to 37.0'	
Steel well cover with cap cemented in place 6-16-82.	

KERR-MCGLE CHEMICAL CORPORATION

Henderson Facility

RCRA Monitor Well No. M-8 Well Construction and Completion Table

Date Started	June 14, 1982
Date Completed	June 15, 1982 (except for surface grout)
Location	NE corner of S-1 pond
Elevation from Top of Well Cover	1780.00'
Drilling Method	Rotary rig
Drilling Fluid	Revert
Donth to Muddy Creek	42.5'
Total Depth of Well	45'
Borehole Diameter	9 5/8"
Well Casing Diameter/Type	5" ID/threaded steel
Well Casing Interval	40' to surface
Perforated Interval	40' - 30'
Perforation Type/ Size /Open Area	Factory slot/ 1/8" x 2" /4 in ² per 10'
Casing Above Ground (Well Cover)	Approx. 15"
Gravel Pack Interval	45' - 27.5'
Type of Gravel	1/8" - 3/8" pea gravel
Surface Seal Interval	22' to surface
Completion:	0 -22.0' grout 22.0'-24.0' well cuttings 24.0'-27.5' bentonite pellets 27.5'-45.0' gravel pack
Consdents: Open hole with gravel from 40.0' to 45.0'.	
Steel well cover with cap cemented in place on 6-16-82.	

KTRR-MCGEE CHEMICAL CORPORATION

Henderson Facility

RCRA Monitor Well No. M-9 Well Construction and Completion Table

•--• • •

Date Started	June 15, 1982
Date Completed	June 15, 1982(except for surface grout)
Location	NW corner of S-1 pond
Elevation from Top of Well Cover	1778.92'
Drilling Method	Rotary Rig
Drilling Fluid	Revert
Depth to Muddy Creek	42'
Total Depth of Well	45'
Boreholc Diameter	9 5/8"
Well Casing Diameter/Type	5" ID/threaded steel pipe
Well Casing Interval	40' to surface
Perforated Interval	40' - 30'
Perforation Type/ Size /Open Area	Factory slot/ 1/8" x 2" /4 in ² per 10'
Casing Above Ground (Well Cover)	Approx. 15"
Gravel Pack Interval	45' - 28'
Type of Gravel	1/8" - 3/8" pea gravel
Surface Seal Interval	22' to surface
Completion:	0 - 22.0' grout 22.0'- 24.0' well cuttings 24.0'- 28.0' bentonite pellets 28.0'- 45.0' gravel pack
Comments: Open hole with gravel from 45.0' to 40.0'.	
Steel well cover with can	

WELL CONSTRUCTION DETAILS

WELL H-28

Depth: 51 feet

Borehole Diameter: 10 inches

Casing Diameter and Type: 6" steel I.D.

Casing Length: 51.7 feet

Top of Casing Eelvation: 1730.33

Screened Interval: 37.4 to 50.5 feet, 6" factory slotted steel well screen

Gravel Pack Interval: 28 to 51.7 feet

Seal Interval: 0-28 feet: cement

Date Completed: 12-18-80

Data from Geraghty and Miller, Inc., 1980.

KERR-McGEE CORPORATION

TO Distribution

November 18, 1981

FROM T. L. Bentley/S. M. Logan SUBJECT RCRA Groundwater Quality Assessment Outline

A written outline for a groundwater quality assessment program is required by RCRA for hazardous waste surface impoundments, landfills and landfarms by November 19, 1981 (45 FR 33241). This outline would be used if a groundwater quality assessment is ever required.

DATE

Attached is an outline to be kept on file at each of your facilities.

T. L. Bentley Environmental Affairs

S. M. Logan Engineering Services

CORPORATE

TLB/SML/dp

414.1.4

Distribution

B. J. Montgomery/S. Livingston cc: G. E. Adam R. A. Clark C. B. Armstrong/R. F. Wohletz P. C. Gaskin (6) E. A. Anglada R. A. Napier W. J. Broussard M. W. Merrill/K. J. Cox S. W. Foster/H. A. Perkins J. E. Warn/W. L. Hoelscher W. B. Hayes S. Pia B. Hoffman T. L. Hurst J. R. Kelley D. F. Schiesz M. Moore D. R. Oakes D. G. Bond J. H. Stallings K. Brothers F. W. McCann L. H. Albright F. D. Lyons W. J. Ganus

GROUNDWATER QUALITY ASSESSMENT PROGRAM OUTLINE

- I. Evaluate Actual and Potential Migration of Hazardous Waste (HW) from the Existing HW Management Area.
 - A. Evaluate physical and chemical characteristics of groundwater from monitoring well data.
 - B. Identify and characterize alternate sources of HW that may influence the site.
 - 1. Existing and Historical on-site
 - 2. Existing and Historical off-site
 - C. Evaluate construction details, operating procedures and operating history of the HW facility.
 - D. Perform a water balance (if possible) around the HW management area.

- E. Examine water levels adjacent to the HW facility.
- II. Define Hydrogeologic Environment
 - A. Potentially impacted aquifer(s):
 - 1. Depth
 - 2. Thickness
 - 3. Areal Extent
 - a. local recharge and discharge points
 - b. determine groundwater usage of aquifer(s)
 - B. Estimate aquifer parameters:
 - 1. Transmissivity
 - 2. Storage coefficient
 - 3. Hydraulic conductivity
 - 4. Effective porosity
 - C. Estimate local and regional directions and average velocities of groundwater flow:
 - 1. Develop potentiometric maps of aquifer systems using water
 - levels from current monitoring points.
 - 2. Develop depth-to-water maps.
- III. Define an Expanded Monitoring Well System
 - A. Review historical groundwater data (if any).
 - B. Establish locations and depths for new monitoring wells to delineate boundaries of impacted areas.
 - C. Sample and analyze monitoring wells for the HW constituents at the facility.

- 1. Determine parameters
- 2. Determine sampling and analysis procedures
- 3. Determine frequency of sampling
- IV. Evaluate Impact from Existing HW Management Area
 - A. Evaluate physical and chemical characteristics of groundwater and determine concentrations of HW constituents.
 - B. Estimate rate of HW migration.
 - C. Estimate extent of HW migration.
 - D. Interpret analytical results in terms of water quality criteria and other applicable regulations.
 - E. Assess significance of analytical data in terms of compliance requirements.
 - V. Prepare Schedule of Implementation
 - A. Submit assessment plan of EPA or to the State if it has obtained RCRA interim authorization.
 - B. Install additional monitoring wells.
 - C. Initiate sampling and analysis.
 - D. Evaluate analytical and physical data.
 - E. Install additional wells, if necessary, and monitor.
 - F. Estimate area impacted from existing HW Management Area.
 - G. Submit assessment report to EPA or to the State if it has obtained RCRA interim authorization.

C DESERT RESEARCH INSTITUTE I UNIVERSITY OF NEVADA SYSTEM



Kerr-McGee Pond Sampling Program

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May, 1982

· WATER RESOURCES CENTER

Summary of Laboratory Analyses Table

Sample Sitc	Sample Type	AS Mg/R	Se mg/&	Ba mg∕&	Cđ mg/&	Total Cr mg/g	Сr+6 mg/g	y by Mg	da Mg∕&	g H g M
AP1	Liquor Sludge	0.05< 0.05<	0.04 0.04<	0.8 0.5<	0.05< 0.05<	0.30	0.2< 0.2<	0.32 0.05<	0.5< 0.5<	0.005< 0.005<
AP2	Liquor Sludge	0.05< 0.05<	0.06 0.04<	0.8 0.5<	0.08 0.05<	0.48 0.60	0.2<	0.63 0.05<	0.5< 0.5<	0.005< 0.005<
AP4	Liguor Sludĝe	0.05< 0.05<	0.04< 0.04<	0.6 0.6	0.05< 0.05<	.0.20 0.20	0.2<	0.15 0.05<	0.5< 0.5<	0.005< 0.005<
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IMPLEMENTATION PROCEDURES

TO COMPLY WITH

RCRA INTERIM STATUS STANDARDS

FOR

GROUNDWATER IMPACT ASSESSMENT AND MONITORING

Prepared By:

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Terry L. Bentley

Sr. Staff Environmental Engineer

Kerr-McGee Corporation

April 1, 1981

SUMMARY

Owners and operators of hazardous waste (HW),¹ surface impoundments, landfills, and landfarms are required by RCRA² to implement one of three alternate groundwater monitoring programs no later than November 19, 1981 (45 FR 33239). These alternate programs are:

- Demonstrate a low potential for migration of HW to water supply wells or surface water; or
- Implement a groundwater monitoring program to determine impact on groundwater quality; or
- 3. Implement an expanded groundwater monitoring program if it is assumed or known that HW has significantly affected groundwater quality.

Dr. W. J. Ganus' Hydrology Department will inspect each Kerr-McGee site to determine the proper location of monitoring wells, help coordinate well drilling, and supervise well installation. Your groundwater monitoring programs should then be implemented in accordance with Figure 1. A description of each item of Figure 1 and designation of responsibilities is provided in the text which follows.

l Hazardous waste as defined in 40 CFR Part 261.

²Resource Conservation and Recovery Act, PL 94-580 dated October 21, 1976.

IMPLEMENTATION PROCEDURES

FOR COMPLIANCE WITH

RCRA GROUNDWATER IMPACT ASSESSMENT AND MONITORING PROGRAMS

Figure 1 is a schematic flow sheet for guidance in establishing and conducting groundwater .impact assessment and monitoring programs required by RCRA.

Item 1: Submit Part A RCRA Permit Application.

These applications were submitted to EPA prior to November 19, 1980, for existing operating facilities.

Responsibility: Submitted by Operating Facility. Approved by Environmental Affairs.

Item 2: Perform First Site Visit.

A reconnaissance visit by Hydrology and Environmental Affairs people will be made to learn of past and present practices, compile hydrologic and geologic data and make a preliminary judgement whether groundwater degradation has occurred.

Responsibility: Hydrology and Environmental Affairs.

Item 3: Prepare Preliminary Site Evaluation Report.

Existing information will be summarized and recommendations made for obtaining any additional data which may be required.

Responsibility: Hydrology and Environmental Affairs.

<u>Item 4:</u> Perform Site Soil/Water Testing (If Existing Data Insufficient).

Soil borings, permeability tests, groundwater quality tests, etc., may be required to further establish a data base.

Responsibility: Operating Facility. Drilling by outside contractor supervised by Hydrology. Concurrence by Environmental Affairs.

(2)

Item 5: Prepare Final Site Evaluation Report and Recommendations.

Preliminary report (Item 3) will be revised to include results of Item 4. One of three groundwater programs will be recommended. A waiver demonstration is preferable, but may not always be possible.

Responsibility: Hydrology, Environmental Affairs, and Operating Facility.

Item 6: Select a Course of Action.

A groundwater program will be selected.

Responsibility: Hydrology. Concurrence by Environmental Affairs and Operating Facility.

Item 7: Prepare Waiver Demonstration Report.

All or part of the monitoring requirements may be waived if low potential for migration of HW to water supply wells or surface water can be demonstrated. A written report, certified by a qualified geologist or geotechnical engineer, must be prepared by November 19, 1981 and kept at the facility.

Responsibility: Hydrology. Concurrence by Environmental Affairs and Operating Facility.

Item 8: Design Groundwater Monitoring System.

A minimum of four groundwater monitoring wells are required. At least one must be upgradient (in the direction of increasing static head) and at least three downgradient (in the direction of decreasing static head).

Separate monitoring systems for each surface impoundment, landfill, or landfarm are not required as long as HW migration can be detected.

The well casing must be screened or perforated and packed with gravel or sand to allow sample collection at proper depths. The casing must also be sealed with cement grout or bentonite clay to prevent contamination of sampling zone. Responsibility: Hydrology. Concurrence by Environmental Affairs and Operating Facility.

Item 9: Install Monitoring Wells.

Wells installed per Item 8. Dr. W. J. Ganus of the Hydrology Department is scheduling installation. A Computer target date of September 1, 1981, has been set for 260ct51 installing all wells.

Responsibility: Operating Facility. Installation by outside contractor supervised by Hydrology. Concurrence by Environmental Affairs.

Item 10: Review Annually.

Groundwater surface elevations must be reviewed annually to insure that the wells are located properly. If not, the number, location, or depth of wells must be modified.

Responsibility: Hydrology and Environmental Affairs.

Item 11: Prepare Groundwater Sampling and Analysis Plan.

A written plan must be prepared by November 19, 1981, and kept at the facility. This plan must include procedures and techniques for:

- 1. Sample Collection;
- 2. Sample Preservation and Shipment;
- 3. Analytical Procedures; and
- 4. Chain of Custody Control.

The plan must require:

- Monitoring wells to be sampled and analyzed quarterly for the parameters in Table I for the first year to establish a baseline. After the first year, groundwater quality parameters must be analyzed annually, indicator parameters semi-annually, and analysis of drinking water parameters can be discontinued. See Appendix I for instructions concerning sampling and analysis techniques.
- 2. Four replicate measurements of each of the indicator parameters to be obtained for upgradient wells the first year. After the

first year, four replicate measurements of each of the indicator parameters must be obtained for all wells.

- 3. Groundwater surface elevations to be determined each time a sample is obtained.
- 4. Sample analysis to be continued during the active life of HW facilities and during the post-closure care period of HW disposal facilitities (normally 30 years).

Responsibility: Operating Facility and Environmental Affairs.

TABLE I

REQUIRED ANALYSES

GROUNDWATER MONITORING WELLS

Drinking Water Parameters:

Arsenic	Methoxychlor
Barium	Toxaphene
Cadmium	2,4-D
Chromium ,	2,4,5-TPSilvex
Fluoride	Radium
Lead	Gross Alpha
Mercury	Gross Beta
Nitrate (as N)	Turbidity
Selenium	Coliform Bacteria
Silver	
Endrin	
Lindane	

Groundwater Quality Parameters:

Chloride	Phenols
Iron	Sodium
Manganese	Sulfate

Contamination Indicator Parameters:

pH Specific Conductance Total Organic Carbon Total Organic Halogen¹

¹There is no standard accepted procedure for the determination of Total Organic Halogen. Environmental Affairs has requested clarification from EPA and will advise.

Item 12: Prepare Outline of Groundwater Quality Assessment Plan.

An outline of a more comprehensive groundwater program than the one implemented the first year (Refer to Item 8) must be prepared by November 19, 1981, and kept at the facility: The program must be capable of determining:

- 1. Whether HW or HW constituents have entered the groundwater.
- 2. Rate and extent of migration of HW or constituents.
- 3. Concentration of HW or constituents in the groundwater.

Responsibility: Hydrology and Environmental Affairs.

Item 13: Implement Sampling and Analysis Program.

Implement program described in Item 11.

Responsibility: Operating Facility. Approved by Environmental Affairs.

Item 14: Establish Initial Background Values During First Year.

Quarterly analyses performed the first year must be recorded and summarized to establish a baseline.

Responsibility: Hydrology and Environmental Affairs.

Item 15: Report Results to EPA.

Results obtained the first year must be sent to Environmental Affairs and Hydrology immediately after they are available to allow time for review prior to submitting to EPA. Results of the <u>drinking</u> water analyses for each well must be submitted to the EPA Regional Administrator within 15 days after receipt, or to the State if it has obtained RCRA interim authorization. Concentrations or values which exceed the Interim Primary Drinking Water standards must be identified.

Reporting of groundwater quality parameters is not

(8)

required. Indicator parameters must be reported in the Annual Report (Refer to Item 17).

Responsibility: Reported by Operating Facility after approval by Environmental Affairs.

Item 16: Continue Sampling and Analysis During Active Life of the Facility.

Continue the program identified in Item 11.

Responsibility: Operating Facility. Approved by Environmental Affairs.

Item 17: Submit Results in Annual Report to EPA.

An Annual Report must be submitted to the EPA Regional Administrator, or to the State, if it has obtained RCRA interim authorization, by March 1 of each year. This report must include:

- 1. Concentrations or values of indicator parameters for each well.
- 2. Statistical evaluations of indicator parameters (Refer to Item 18).
- 3. Results of the evaluation of groundwater surface elevations performed in Item10 and modifications required to the monitoring system.

Responsibility: Submitted by Operating Facility. Approved by Environmental Affairs.

Item 18: Perform Statistical Analyses on Indicator Parameters.

First year indicator parameter measurements must be pooled for each upgradient well and the initial background arithmetic mean and variance of each parameter calculated. After the first year, the arithmetic mean and variance of each indicator parameter must be calculated semi-annually for each upgradient and downgradient well using the four replicate measurements taken on each sample.

Responsibility: Hydrology and Environmental Affairs.

Item 19: Compare Semi-Annual Indicator Results to Initial Background Values.

Each semi-annual arithmetic mean calculated for a specific parameter must be compared with its background arithmetic mean using the student's t-test (statistical test) at the 0.01 level of significance to establish whether a "statistically significant" change has occurred. The calculation for the student's t-test is as follows:

$$t = \frac{\bar{x} - \bar{y}}{s} - \sqrt{\frac{nm}{n+m}}$$

- where, x = average of four replicate measurements obtained during semiannual testing.
 - y = average of sixteen (four replicate measurements per quarter) measurements establishing initial background.
 - s = overall standard deviation for all measurements under examination.

n = 16

therefore, $t = \frac{\overline{x} - \overline{y}}{s}$ (1.79)

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A "t" value exceeding 2.84 represents a "statistically significant" change.

Responsibility: Hydrology and Environmental Affairs.

Item 20: Has Degradation Occurred? (Upgradient Wells)

Student's t-test results must be evaluated for upgradient wells. Groundwater sampling and analysis must continue.

Responsibility: Hydrology and Environmental Affairs.

Item 21: Submit Results in Annual Report to EPA.

Significant increases (or pH decreases) in upgradient wells must be submitted in the Annual Report due March 1 of each year. Groundwater sampling and analysis must continue. Responsibility:

Submitted by Operating Facility. Concurrence by Environmental Affairs and Hydrology.

Item 22: Has Degradation Occurred? (Downgradient Wells)

Student's t-test results must be evaluated for downgradient wells: If degradation has not occurred, the original sampling and analysis program must be continued.

Responsibility: Hydrology and Environmental Affairs.

Item 23: Prepare Assessment Plan from Outline (See Item 12).

If comparisons for the downgradient wells show significant increases (or pH decreases), a specific plan for a groundwater quality assessment program must be prepared. It must be certified by a qualified geologist or geotechnical engineer and specify.

- 1. The number, location, and depths of wells.
- Sampling and analytical methods for those HW constituents in the facility.
- 3. Evaluation procedures, including any use of previously gathered information.
- 4. An implementation schedule.

Responsibility: Hydrology. Concurrence by Environmental Affairs.

Item 24: Obtain Additional Samples and Analyze.

Additional samples must be obtained immediately from downgradient wells where significant increases (or pH decreases) were detected. Samples must be split in two and four replicate measurements of each split sample taken for indicator(s) showing significant differences.

Responsibility: Operating Facility. Approved by Environmental Affairs.

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Item 25: Were Results Due to Lab Error?

Measurements must be compared to background data to determine if results were due to laboratory error. If they were, the original sampling and analysis program must be continued.

Responsibility: Environmental Affairs and Operating Facility.

Item 26: Notify EPA.

The EPA Regional Administrator, or the State if it has obtained RCRA interim authorization, must be notified in writing within seven days if degradation has occurred.

Responsibility: Notification by Operating Facility. Approved by Environmental Affairs.

Item 27: Submit Assessment Plan to EPA.

The assessment plan must be submitted within 15 days after the notification of Item 26.

Responsibility: Submitted by Operating Facility. Approved by Environmental Affairs.

Item 28: Implement Assessment Plan.

The assessment plan must be implemented as soon as technically feasible and determine:

- 1. Rate and extent of HW migration in the groundwater.
- 2. Concentrations of HW constituents in the groundwater.

Responsibility: Operating Facility, Hydrology, and Environmental Affairs.

Item 29: Submit Assessment Report to EPA.

A written report assessing the groundwater quality must be submitted within 15 days after the first determination of Item 28. It must be submitted to the EPA Regional Administrator, or State if it has obtained RCRA interim authorization. Responsibility:

Submitted by Operating Facility. Prepared by Hydrology. Approved by Environmental Affairs.

Item 30: Has Degradation Occurred?

If the first determination shows that HW has <u>not</u> entered the groundwater, the original sampling and analysis program must be continued. If HW has entered the groundwater, proceed to Item 31.

Responsibility: Hydrology and Environmental Affairs.

Item 31: Determine Closure Status.

Was the assessment made before or after final closure of the facility?

Responsibility: Operating Facility and Environmental Affairs.

Item 32: Make Quarterly Assessments Until Final Closure.

Determinations of Item 28 must be continued quarterly until final closure. Results must be submitted in the Annual Report due March 1 of each year.

Responsibility: Submitted by Operating Facility. Prepared by Hydrology. Approved by Environmental Affairs.

Item 33: Cease Determination.

Assessments may be discontinued if the assessment plan was implemented during the post-closure care period.

Responsibility: Operating Facility and Environmental Affairs.

Item 34: Design Expanded Groundwater Monitoring System.

An expanded groundwater monitoring program may be implemented if it is assumed or known that groundwater quality has been significantly affected. Responsibility: Hydrology. Concurrence by Environmental Affairs and Operating Facility.

Item 35: Install Monitoring Wells.

Wells installed per Item 34. Dr. W. J. Ganus of the Hydrology Department is scheduling installation. A target date of September 1, 1981 has been set for installing all wells.

Responsibility: Operating Facility. Installation by outside contractor supervised by Hydrology. Concurrence by Environmental Affairs.

Item 36: Submit Groundwater Quality Assessment Plan to EPA.

A specific plan for a groundwater quality assessment program must be submitted by November 19, 1981 to the EPA Regional Administrator, or to the State if it has obtained RCRA interim authorization. The plan must specify:

- 1. The number, location, and depth of wells.
- Sampling and analytical methods for those HW constituents in the facility.
- 3. Evaluation procedures, including any use of previously gathered information.
- 4. An implementation schedule.

The first groundwater samples must be analyzed prior to November 19, 1981. Implementation must be in accordance with Items 28-33 with the exception of Item 30, which does not apply.

Responsibility: Submitted by Operating Facility. Prepared by Hydrology and Environmental Affairs.

Prepared By: Concurred By Approved By: L. Hurst Т م الم الم المست Dated:

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APPENDIX I

SAMPLING AND ANALYSES TECHNIQUES

GROUNDWATER MONITORING PROGRAMS

APPENDIX I

SAMPLING AND ANALYSES TECHNIQUES GROUNDWATER MONITORING PROGRAMS

The following standard sampling and preservation techniques must be employed when monitor wells are sampled and analyzed.

Preparation for Sampling

- 1. Measure depth to water from the top of the well casing within 1/10 of an inch.
- 2. Clear standing water from the casing as specified below:
 - (a) <u>Pumping</u>: A pump should be used for sampling whenever possible. Three casing-volumes of water should be pumped from the well before sampling. For wells that have a slow recovery rate, the well should be evacuated and allowed to recover at least once prior to sampling.
 - (b) <u>Bailing</u>: When pumping is not possible (i.e. a 2" casing) the well casing must be bailed to dryness at least once then allowed to recover before sampling. If this is not possible due to a rapid recovery rate, one casing-volume should be bailed from the well before sampling.

Sampling Volumes, Containers, and Preservation Requirements

The attached Table I lists EPA recommended sample volumes, preservatives, containers, and holding times. For example, the following samples, preservatives, and containers would be required for RCRA groundwater monitoring.

Container Preservative		Parameter		
Plastic/Glass (polyethylene w/ polypropylene top recommended) 1 gallon	Filter on site using a 0.45 v membrane filter and pressure apparatus. Add HNO ₃ to pH <2	As, Ba, Cd, Cr, Pb, Hg, Se, Ag, Fe, Mn, Na, Ra Gross α, Gross β.		
Glass only l quart	Add H_3PO_4 to pH <4, 1.0 g CuSO ₄ /1.	Phenols		

Container

Preservative

Glass only l quart

Cool to 4°C.

Plastic/Glass l gallon

Cool to 4°C.

Plastic/Glass l quart

Cool to 4°C. H_2SO_4 or HCl to pH <2.

ph^{*}must be determined on site.

*For establishing baseline data, four replicate determinations must be obtained for each sample of the upgradient well for these parameters.

All glass containers should be cleaned thoroughly and rinsed with distilled water before sampling. Preferably new sterile plastic containers should be used when possible. Each container should be rinsed at least once with the sample before filling one sample

The RCRA regulations require total organic halogen analysis for which there is no standard procedure. Environmental Affairs is awaiting clarification from EPA on this requirement.

Analyses

All analyses must be conducted by using procedures found in "Standard Methods for the Examination of Water and Wastewater", 14th edition; "Methods for Chemical Analysis of Water and Wastes" (EPA 600/4-79-020, March, 1979); or the Annual Book of ASTM Standards, Part 31, "Water" (1976).

Environmental Affairs has prepared a list of certified laboratories. Except for those facilities that have complete in-house capability and routinely perform NPDES or equivalent analysis, it is imperative a competent State/EPA certified laboratory do the sampling and analyses at each facility.

Chain-of-Custody

Proper chain-of-custody records are essential to avoid any questions regarding sample integrity. A Kerr-McGee standard chain-of-custody form has been prepared which must accompany the sample from the time it is taken to the time it is analyzed. The complete chain-of-custody record must be kept by the K-M

Parameter

Pesticides--must be extracted 7 days after sampling (Proposed in the 12/18/78 FR)

F, C1, SO₄, NO₃(N) turbidity, coliform bacteria, specific conductance*

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site manager and a copy sent to Environmental Affairs with the analytical results.

References

- "Methods for Chemical Analysis of Water and Wastes", (EPA 600/4-79-020) March, 1979.
- "Handbook for Analytical Quality Control in Water and Wastewater Laboratories", (EPA 600/4-79-019) March, 1979.
- "Procedures Manual for Groundwater Monitoring at Solid Waste Disposal Facilities", (EPA 530/SW-611) August, 1977.

TABLE 1

RECOMMENDATION FOR SAMPLING AND PRESERVATION OF SAMPLES ACCORDING TO MEASUREMENT⁽¹⁾

	Vol.			
Measurement	Req. <u>(ml)</u>	Container ⁽²⁾	Preservative	Holding <u>Time⁽³⁾</u>
100 Physical Properties				• •
Color	50	P,G	Cool, 4°C	24 Hrs.
Conductance	100	P,G	Cool, 4°C	24 Hrs."
Hardness	100	P,G	Cool, 4° C HNO ₃ to pH < 2	6 Mos. ⁽⁵⁾
Odor	200	G only	Cool, 4°C	24 Hrs.
pH	25	P,G	Det. on site	6 Hrs.
Residue -				
Filterable	100	P,G	Cool, 4°C	7 Days
Non- Filterable	100	P,G	Cool, 4°C	7 Days
Total	100	P,G	Cool, 4°C	7 Days
Volatile	100	P,G	Cool, 4°C	7 Days
Settleable Matter	1000	P,G	None Req.	24 Hrs.
Temperature	1000	P,G	Det. on site	No Holding
Turbidity	100	P,G	Cool, 4°C	7 Days
200 Metals				
Dissolved	200	P,G	Filter on site HNO_3 to $pH < 2$	6 Mos. ⁽³⁾
Suspended	200		Filter on site	6 Mos.
Total	100	P,G	HNO_3 to $pH < 2$	6 Mos. ⁽³⁾

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TABLE 1 (CONT)

	Vol. Roc			TT - 1 2
Measurement	(ml)	Container ⁽²⁾	Preservative	Time ⁽³⁾
Mercury Dissolved	100	P,G	Filter on site HNO ₃ to pH < 2	38 Days (Glass) 13 Days (Hard Plastic)
Total	100	P,G	HNO ₃ to pH < 2	38 Days (Glass) 13 Days (Hard Plastic)
300 Inorganics, Non-Meta	llics			
Acidity	100	P,G	None Req	24 Hrs.
Alkalinity	100	P,G	Cool, 4°C	24 Hrs.
Bromide	100	P,G	Cool, 4°C	24 Hrs.
Chloride	50	P,G	None Req.	7 Days
Chlorine	200	P,G	Det. on site	No Holding
Cyanides	500 -	P,G	Cool, 4°C NaOH to pH 12	24 Hrs.
Fluoride	300	P,G	None Req.	7 Days
Iodide	100	P,G	Cool, 4°C	24 Hrs.
Nitrogen				
Аттоліа	400	P,G	Cool,4°C H_2SO_4 to $pH < 2$	24 Hrs.
Kjeldahl, Total	500	P,G	Cool, 4° C H ₂ SO ₄ to pH < 2	24 Hrs. ⁽⁶⁾
Nitrate plus Nitrite	100	P,G	Cool, 4℃ H₂SO₄ to pH<2	24 Hrs. ⁽⁶⁾
Nitrate	100	P,G	Cool, 4°C	24 Hrs.
Nitrite	50	P,G	Cool, 4°C	48 Hrs.

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TABLE 1 (CONT)

Measurement	Vol. Req. <u>(ml)</u>	Container ⁽²⁾	Preservative	Holding Time ⁽³⁾
Dissolved Oxygen Probe	300	G only	Det. on site	No Holding
Winkler	300	G only	Fix on site	4–8 Hours
Phosphorus Ortho- phosphate, Dissolved	50	P,G	Filter on site Cool, 4°C	24 Hrs.
Hydrolyzable	50	P,G	Cool, 4°C H₂SO₄ to pH<2	24 Hrs. ⁽⁶⁾
Total	50	P,G	Cool, 4°C H_2SO_4 to $pH < 2$	24 Hrs. ⁽⁶⁾
Total, Dissolved	50	P,G ¹	Filter on site Cool, 4°C H_2SO_* to $pH < 2$	24 Hrs. ⁽⁶⁾
Silica	50	P only	Cool, 4°C	7 Days
Sulfate	50	P,G	Cool, 4°C	7 Days
Sulfide	500	P,G	2 ml zinc acetate	24 Hrs.
Sulfite	50	P,G	Det. on site	No Holding
400 Organics				
BOD	1000	P,G	Cool, 4°C	24 Hrs.
COD	50	P,G	H_2SO_4 to $pH < 2$	7 Days ⁽⁶⁾
Oil & Grease	1000	G only	Cool, 4°C H₂SO₄ or HCl to pH<2	24 Hrs.
Organic carbon	25	P,G	Cool, 4°C H₂SO₄ or HCl to pH<2	24 Hrs.
Phenolics	500	G only	Cool, 4°C H ₃ PO ₄ to $pH < 4$ 1.0 g CuSO ₄ /1	24 Hrs.
MBAS	250	P,G	Cool, 4°C	24 Hrs.

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TABLE 1 (CONT)

	Vol.			
Measurement	Req. <u>(ml)</u>	Container ⁽²⁾	Preservative	Holding <u>Time</u> ⁽³⁾
NTA	50	P,G	Cool, 4°C	24 Hrs.

- 1. More specific instructions for preservation and sampling are found with each procedure as detailed in this manual. A general discussion on sampling water and industrial wastewater may be found in ASTM, Part 31, p. 72–82 (1976) Method D-3370.
- 2. Plastic (P) or Glass (G). For metals, polyethylene with a polypropylene cap (no liner) is preferred.
- 3. It should be pointed out that holding times listed above are recommended for properly preserved samples based on currently available data. It is recognized that for some sample types, extension of these times may be possible while for other types, these times may be too long. Where shipping regulations prevent the use of the proper preservation technique or the holding time is exceeded, such as the case of a 24-hour composite, the final reported data for these samples should indicate the specific variance.
- 4. If the sample is stabilized by cooling, it should be warmed to 25°C for reading, or temperature correction made and results reported at 25°C.
- 5. Where HNO₃ cannot be used because of shipping restrictions, the sample may be initially preserved by icing and immediately shipped to the laboratory. Upon receipt in the laboratory, the sample must be acidified to a pH <2 with HNO₃ (normally 3 ml 1:1 HNO₃/liter is sufficient). At the time of analysis, the sample container should be thoroughly rinsed with 1:1 HNO₃ and the washings added to the sample (volume correction may be required).
- 6. Data obtained from National Enforcement Investigations Center-Denver, Colorado, support a four-week holding time for this parameter in Sewerage Systems. (SIC 4952).



GROUNDWATER ASSESSMENT OUTLINE

The objective of the following summary is to provide the operator of this facility with guidance should groundwater analysis prescribed by 40 CFR 265.92 - 93 indicate groundwater contamination is occurring or has occurred. This summary is based on the attached corporate outline prepared by T. Bentley November 18, 1981.

After the first year, groundwater monitoring sampling will continue on a semiannual basis for groundwater quality or groundwater contamination parameters, as is appropriate. Each well sampled shall have the indicator parameters, pH, specific conductance, total organic carbon, and total organic halogen, analyzed in quadruplicate and the mean and variance compared with the initial background mean and variance. If there is a statistically significant increase (or decrease in case of pH), the operator shall do the following:

- 1. Resample and analyze to determine if variation was the result of laboratory error.
- 2. Should resampling confirm difference
 - a. On upgradient wells report the findings in the annual report to the administrator.
 - b. On downgradient wells report the results within seven days of receipt of complete laboratory report to administrator that facility may be affecting groundwater.
- 3. Within 15 days of initial notification, develop and submit to administrator a groundwater assessment program. The details to be included in the assessment plan are spelled out in the corporate outline attached to this summary.

It is understood that suggestions and recommendations of the administrator may change the initial plan outline and implementation schedule but every effort will be made to fulfill the regulatory requirements in a timely manner.

ENVIRONMENTAL PROTECTION AGENCY

REGION IX

SURVEILLANCE & ANALYSIS DIVÍSION

NPDES Compliance Monitoring Report

Permittee: Kerr-McGee Chemical Corporation Oklahoma City, Oklahoma

Facility: Kerr-McGee Chemical Corporation Henderson NV

Permit No.: NV0000078

Date of Inspection: June 19, 1980

Inspection Participants:

EPA: Kenneth D. Greenberg Environmental Engineer

Facility: Charles B. Armstrong Plant Manager

> Richard F. Wohletz Superintendent, Plant Technical Services

Report Prepared by: Kenneth D. Greenberg

FINDINGS

Introduction

The Kerr-McGee Chemical Corporation operates an inorganic chemical production plant at the Basic Management, Inc. (BMI) industrial complex in Henderson, Nevada. The Kerr-McGee facility is subject to NPDES Permit No. NV0000078 which became effective on February 24, 1977 and expires on September 30, 1981. Under the permit, Kerr-McGee is authorized to discharge a daily maximum of 4.0 mgd of non-contact cooling water during the period of June 1 through September 30. The non-contact cooling water is discharged through an open ditch to Las Vegas Wash. Other process streams are either recycled or discharged to lined evaporation ponds located at the plant site. Pond parameters are summarized in Table-1 and their locations are shown in Figure 1.

The BMI industrial complex was originally owned by the U.S. Government which produced magnesium metal at the facility. In 1945, the portion of the industrial complex which is currently owned by Kerr-McGee was taken over by the Western Electrochemical Company. Western Electrochemical merged into American Potash and Chemical Corp. which took over operations at the facility in 1955. Finally, American Potash and Chemical Corp. merged into Kerr-McGee which gained control of the plant operations in 1967. Except for expansion to the production of boron compounds in the early 70's, the list of inorganic products at the facility (see details below) is basically unchanged since 1945.

Prior to 1976, liquid waste streams and slurried solid wastes from the facility were discharged to the unlined BMI ponds located across Boulder Highway to the northeast of the production area. In the mid-70's lined ponds were constructed on the Kerr-McGee plant property to accomodate liquid waste and recycle streams. Solid wastes have been and continue to be disposed on the Kerr-McGee plant property. Solid wastes were also disposed at the BMI dump, located northwest of the facility, until the dump closed in early 1980.

Production Processes and Wastewater Streams

Production at the Kerr-McGee facility is divided into four major processes: 1) Sodium chlorate, 2) perchlorates, 3) manganese dioxide, and 4) boron chemicals.

In the first process, sodium chlorate (NaClO₃) is produced in an electrolytic process from raw materials of sodium chloride and water. Sodium chlorate is sold for use in paper pulp bleaches and is also used as an intermediate in the production of perchlorates at the Henderson facility. Waste from the production of sodium chlorate consists of a filter cake containing impurities from the raw materials and filter aid. In the past, the filter cake (containing calcium sulfate, calcium carbonate, graphite, and diatomaceous earth) has been slurried to the BMI ponds or disposed at the BMI dump. The filter cake, which contains 50% moisture, is currently dumped on the ground surface in the northwest corner of the plant property (see Figure 1).

Spills, cooling tower leaks, and excess storm runoff from the sodium chlorate process are discharged to the lined ponds, P-2, and P-3. Water from these ponds is recycled back to the process.

During the summer, non-contact cooling water, used in the sodium chlorate process is discharged to Las Vegas Wash via the BMI storm ditch and the Alpha ditch. Additional details on this discharge are provided in the subsection below on plant effluent.

The second major process at Kerr-McGee involves the production of ammonium perchlorate (NH4ClO4) and potassium perchlorate (KClO4) which are used in the manufacture of rocket fuels. In this process, a solution of sodium chlorate is first electrolytically converted to sodium perchlorate (NaClO4). The sodium perchlorate is then combined with salts of either ammonia or potassium to form the respective perchlorates.

Wastes from the ammonium perchlorate process include a filter cake and chromic hydroxide which is derived from the use of chromium as a filter aid. In the past, the filter cake, containing calcium sulfate and calcium carbonate, was slurried to the BMI ponds. Now the filter cake and chromic hydroxide are discharged in slurry form to the lined ponds AP-1 or AP-2. At the time of the inspection pond AP-2 was not in use and was empty. Liquid from these ponds is recycled back to the process through the pump basin AP-3. Emergency overflows from the ammonium perchlorate cooling tower are discharged to the lined pond AP-4. A minor stream from a caustic scrubber in the ammonium perchlorate process is discharged to pond P-2 along with wastes from the sodium chlorate process (described above). A waste stream from the potassium perchlorate process containing NaCl, KCl, and KClO4 is discharged to the lined pond S-1.

The third major process at Kerr-McGee is the production of manganese dioxide which is sold for use in high performance dry cells. Low grade manganese ore is crushed, roasted, and then combined with sulfuric acid. The resulting manganous sulfate is then converted to manganese dioxide (MnO₂) by electrolysis. Wastes from this process include a solid waste containing silica, alumina, iron, and heavy metals which is filtered from the roasted ore after it has been combined with sulfuric acid. This waste, which amounts to 50% by weight of the raw ore, is currently disposed in piles at the Kerr-McGee plant site (see Figure 1). A minor waste stream of sodium phosphate solution is discharged to pond C-1. The solution, which is used for cleaning the electrolytic cell electrodes, is discharged in batches of approximately 5,000 gallons once or twice per week. All other water used in the production of manganese dioxide is recycled.

The fourth major process at Kerr-McGee is the production of elemental boron (B), boron trichloride (BCl₃), and boron tribromide (BBr₃). Boron trichloride is used in the manufacture of boron filament for aircraft structures.' Boron tribromide is used in semiconductor doping. Elemental boron is used in pyrotechnics. Waste streams from the production of boron chemicals include a leachate stream containing magnesium sulfate (500 gal./day) and a wet scrubber stream (7000 gal./day). These wastes were being discharged to pond S-1 at the time of the inspection.

Pond C-1 receives a waste stream from the plant's main boiler and cooling tower blowdown. The company reported that the discharge to pond C-1 contains 22,450 ppm total dissolved solids. Liquid in pond C-1 is not recycled back to the plant.

Ponds and Pond Leakage Monitoring

The Kerr-McGee discharge permit requires that:

- "1. If any waste waters... are placed in ponds, such ponds shall be located and constructed so as to:
 - a. contain with no discharge the once-in-one-hundred years storm at said location;
 - b. Withstand with no discharge the once-in-one-hundred years flood of said location; and
 - c. prevent escape of waste water by leakage.
- 2. The permittee shall submit to the Director and the Regional Administrator a summary of the results obtained from monitoring for seepage and leakage at the frequency specified in Part 1.C.2."

Plant personnel conduct a program of monitoring for pond leakage which involves 1) checking the level of liquid in each pond once or twice per week and 2) analyzing the concentration of certain salts in each pond every two or three weeks. With this data, large leaks can be detected by looking for unusual changes in the level of a pond or the load of dissolved salts in a pond. The levels of liquid in adjacent ponds is also compared as a means for detecting losses of liquid in excess of the evaporation rate. A spot check of recorded data from this monitoring program revealed no unusual drops in pond level. Kerr-McGee officials stated that the monitoring program had revealed leaks in the liners of ponds P-1 and AP-2 which have now been repaired.

However, the current leakage monitoring program is not capable of detecting small leaks. The following techniques would make leak detection more exact but would still be subject to inaccuracies due to inherent errors in measurements. Δ continuous level recorder at each pond would provide a more complete picture of liner integrity and make comparison of levels in different ponds easier. However, it would still be difficult to separate liquid losses due to evaporation and small leaks. Continuous level recorders would also provide estimates of the volume of inflow to ponds which currently is not measured. TIMET, one of the other companies at the BMI complex, uses a lithium tracer for detection of leaks in their lined ponds. A known quantity of lithium carbonate is placed in each pond. Periodically the lithium concentration and the pond volume is determined. From this data the load of lithium in each pond can be calculated. drop in the amount of lithium in a given pond would be due to loss through leakage since the lithium load is not affected by evaporation. It is not necessary to measure pond inflow with the lithium tracer technique.

As noted above, Kerr-McGee has been recording data from their leakage monitoring program. However, they have not been reporting this data to the Nevada DEP or EPA as required by the permit. The plant superintendent said that he was not aware of the requirement to report this data.

Originally all of the lined ponds at the Kerr-McGee facility were lined with a single layer of polyvinyl chloride (PVC) on the bottom joined to chlorinated polyethylene (CPE) on the side walls. CPE was used on the side walls of the pond because of its greater resistence to solar radiation. Kerr-McGee officials explained that the PVC/CPE pond liners have been deteriorating over the years because the two membrane materials are incompatable when in contact with each other. In four of the Kerr-McGee ponds, the original PVC/CPE liner developed leaks and have been replaced with a liner made of nylon reinforced rubber. During the inspection the plant superintendent stated that the company planned to take pond S-1 out of service in the near future in order to replace its PVC/CPE lining. The potassium perchlorate waste stream would be rerouted to pond P-1 which has a nylon reinforced rubber liner but was not in use at the time of the inspection. Ponds AP-1 and AP-4 still have the original PVC/CPE liners which the superintendent claimed were in satisfactory condition. In future inspections, the condition of the AP-1 and AP-4 pond linings should be checked. The new nylon reinforced hypolon liners appear to be holding without excessive deterioration.

Plant Effluent and Monitoring Requirements

During the summer months, an average of 3.2 mgd of non-contact cooling water is used in the sodium chlorate process. As authorized by the NPDES permit, this non-contact cooling water is discharged to an unlined, open ditch. The discharge flows approximately 200 ft. to the north at which point it flows into the BMI storm ditch (another unlined open ditch) and continues to the east (see Figure 1). In accordance with their permit, the Kerr-McGee discharge passes under Boulder Highway in the BMI siphon, and through the crossover pipe to the Alpha ditch and on to Las Vegas Wash (see Figure 2).

The BMI storm ditch is also used by Stauffer Chemical Company for their permitted stormwater discharges. Stauffer is located immediately west of Kerr-McGee in the BMI complex. However, under Stauffer's permit, their stormwater discharges are required to flow through the BMI siphon and down the acid ditch which discharges to the upper BMI ponds. If Stauffer discharged stormwater during the summer months it would combine with the Kerr-McGee discharge of non-contact cooling water in the BMI storm ditch. Under their discharge permit, Stauffer would be required to close the crossover pipe in order to route their stormwater discharge to the upper BMI ponds. However, closing the crossover pipe would also cause the Kerr-McGee effluent to flow to the upper BMI ponds. Kerr-McGee is not authorized to discharge to the upper BMI ponds. On the other hand, if the crossover pipe were left open, then the Kerr-McGee effluent and the Stauffer stormwater would flow into the Alpha ditch. The flow of Stauffer stormwater to the Alpha ditch is not allowed under the Stauffer discharge permit. The conflict described above can be resolved either by 1) a permit modification or, 2) a rearrangement of the discharge ditches so the Kerr-McGee and Stauffer discharges do not use a common discharge route.

Under their discharge permit Kerr-McGee is required to monitor their effluent for flow, temperature, pH and oil and grease. The company is also required to measure the change in total dissolved solids and suspended solids of the non-contact cooling water which occurs in the process. Temperature is measured and recorded on a continuous basis while composites for TDS, suspended solids, and oil and grease are collectd manually at the head of the open discharge ditch which carries the non-contact cooling water (see Figure 1). Composites are made once a week by filling 8 glass jars on an hourly basis from discharge water grabbed in a plastic bucket. The plant superintendent explained that the composites are not flow proportioned because the effluent flow is constant. A spot check of effluent flow charts revealed that this is generally true. However, the flow does fluctuate significantly on some days. Therefore, composite samples should be flow proportioned.

An orifice meter located in the plant production area is used to measure the flow which is reported in the discharge monitoring reports . This meter is only capable of measuring the discharge of non-contact cooling water. Other Kerr-McGee discharges which may occur would not be measured by the orifice meter. Other potential discharges to the BMI storm ditch by Kerr-McGee are storm water entering the unbermed ditch, pond overflows, or process spills. Many of the floor drains in the production area have been plugged to prevent such possibilities. However, during an earlier plant visit in August 1979, water leaking from a supply line in the plant was observed to be discharging through the open ditch which joins the BMI storm ditch near pond C-1. Due to the slope of the land and the lack of berms along the open ditches, storm runoff from Kerr-McGee plant property could easily enter the ditches and flow off plant property. Under their NPDES permit, Kerr-McGee is allowed to discharge noncontact cooling water. The discharge of any other liquids is not permitted.

Kerr-McGee has installed a weir and flow meter on the BMI storm ditch at the point it passes to TIMET property (see Figure 1). This meter would be capable of measuring all Kerr-McGee discharges in the BMI storm ditch. However, this meter would also measure any flow which may be discharged in the BMI storm ditch by Stauffer Chemical Company. Furthermore, the weir is not properly installed since it is not perpendicular to the axis of flow in the ditch. It is also possible for Kerr-McGee to discharge through an open ditch which enters TIMET property at a point south of the BMI storm ditch (see Figure 1). There is no flow measuring device on this ditch. In summary, with the flow measuring devices in place at the time of the inspection, it is not possible to measure all potential discharges from the Kerr-McGee plant.

A review of the plant's discharge monitoring reports for the summer months of 1979 revealed that the discharge was within the permitted limits with the exception of some exceedances of the pH limit. In 1979, the maximum limit on pH of 8.5 was exceeded in July (8.9), August (8.8), and October (8.6). The plant water supply (used for cooling water) has an average pH of 8.0 which contributes to the high pH of the discharge. The State of Nevada Division of Environmental Protection granted Kerr-McGee permission to continue their discharge of non-contact cooling water in October 1979. Due to unusually warm weather, the company found it necessary to continue the discharge until October 25, 1979.

- 6 -

SELF-MONITORING DEFICIENCIES

All self-monitoring procedures were in accordance with EPA requirements, EPA recommendations, and NPDES permit specifications with the exception of the following:

- The permittee has failed to report the results of monitoring for leakage from holding ponds as required by the permit. (see detail in Findings Section above).
- 2. Composite samples of the plant effluent are not flow proportioned during collection as required by the permit. Plant personnel claim that, due to the uniform nature of the effluent, analysis results would not change significantly if the sample were flow proportioned. Kerr-McGee should show that this is true by comparing results obtained under both compositing techniques.

TABLE 1: KERR-MCGEE CHEMICAL CORPORATION PONDS

CINOA	Process Waste	Evaporation or Recycle	Liner*	Surface Area (acres)	Capacity (gallons)
	Steam Plant (Boiler), Cooling Tower	Evaporation	PVC w/reinforced rubber walls	1.4	2,750,000
p-1**	KCl04 (future plans)	Evaporation	Reinforced rubber	0.7	000'006<
P-2	NaClO3 spills and NH4ClO3 scrubber	Recycle	Reinforced rubber	0.25	350,000
P-3	NaClO ₃ spills	Recycle	Reinforced rubber	0.25	350,000
AP-1	NH4C104	Recycle	PVC w/CPE walls	0.3	425,000
AP-2**	NH4CIO4	Recycle	Reinforced rubber	0.3	425,000
AP-3	Pump basin for AP-1 and AP-2	Recycle	unknown	0.1	50,000
AP-4	NH4Cl04 Cooling Tower	Evaporation	PVC w/CPE walls	0.4	650,000
S-1	KClO4 and Boron Compounds	Evaporation	PVC w/CPE walls	1.0	2,000,000
1					

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All ponds have single layer linings. Ponds which were not in use at time of inspection (June, 1980).





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	NPDES CO	MPL. CE INS	PECTION	REPC	RT (Coding) Inst		on back of	last	parje)			
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SECTION A - P	ermit Summary	•									·	
Kerr-M PiO Box	DRESS OF FACILI CGee Chemica 55	ry Include Count 1 Corporati	y, State and c A	ZIP con	de)	•		•	EXPIR Sep	ATION	DATE	1
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RESPONSIBLE	OFFICIAL			TITL	E		<u></u>		PHON			·
Charles B. Armstrong					ant Manage	ec	· <u> </u>		(702)	565	<u>- 89</u>	101
Richard F. Wohlerz					echnical Services (702) 565- 8901						701	
SECTION B - E	fluent Characteristic	s (Additional sheet	s attached		-) Data fr	rom DA	NR.for	Sept	ember	1979	1	
PARAMETER/ OUTFALL		MINIMUM	AVERA Daily	GE	MAXIMUM Daily		A	DIT	ONAL		•	
Flow/	SAMPLE MEASUREMENT	- · · ·	3.8,	ngd	4.0 mg.d	ŀ	-	•				
001	PERMIT REQUIREMENT	-			4.0 mgd							
Total Dissolved	SAMPLE MEASUREMENT				•		0 mg/1	net	r cha	nge		
Solids/ 001	PERMIT REQUIREMENT				-	Net effice	change + nt net te	from cx	eed ;	er suj 15 mg,	r pig /1 mei	rc Thy
Temp.	SAMPLE MEASUREMENT		20°0	5	2.2°C							<u> </u>
001	PERMIT		279	c	<u>.</u> 31°C							
Oil and Grease/	SAMPLE MEASUREMENT	÷	1.7	mg/1	2.6 mg/1					•		
001	PERMIT REQUIREMENT		10 mg,	1	15 mg/1				• *		<u> </u>	
рН/	SAMPLE MEASUREMENT	8.1			8.3		· · ·					
001	PERMIT	6.5			8.5						-	
SECTION C - Fa	Citity Evaluation (S =	Satistactory, L' =	Unsetistection	r. N/4	= .Not applicable;		······					
S EFFLUENT WITHIN PERMIT REQUIREMENTS N4:OPERATION					AND MAINTENANCE U SAMPLING PROCEDUPES							
S PERMIT VE	RIFICATION		LOW MEASL	JREMENTS OTHER:				HY PRACTICES				
SECTION D - Co	mments	•		•	·							
SECTION E . Ins	SIGNATUF	1ES			AGENCY DATE DIVIS			ENEN ICN	T			
INSPECTED BY Kén	neth D. H.	eenbers		E	P.A TX	Feb	. 19, 1981	1	COMP	LIANC	ESTAT	rus
INSPECTED BY	•	0	۰.		•					OMPLI	ANCE	N.C =
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Sections F thru L: Complete on all insp	pections, as appropriate.	N/A = Not Applicable	PERM A/	IT NO.	-18
SECTION F - Facility and Permit Backgroun	nd	•	/_	VULLU	10
ADDRESS OF PERMITTEE IF DIFFEREN	IT FROM FACILITY	DATE OF LAST PREVIOUS IN	VESTIGATIO	N BY EPA	TATE
Including City, County and ZIP code)		EPA - Feb.	15,1979		
Rest-McGee Chemical	Corporation	FINDINGS	21.1		
McGee lower					
Oklahoma City, Oklaho	ma	•••			-
SECTION G - Records and Reports					
RECORDS AND REPORTS MAINTAINED	AS REQUIRED BY PERMI		Further explan	ation attache	d X
DETAILS: Leakage meniteri	ing data not report	ed			*
ADEQUATE RECORDS MAINTAINED	OF:				
(I) SAMPLING DATE, TIME, EXAC	TLOCATION		X YES		
(II) ANALYSES DATES, TIMES			1 YES		<u> </u>
(III) INDIVIDUAL PERFORMING AN	ALYSIS		V YES		
(W) ANALYTICAL METHODS/TECH	INIQUES USED		X YES		<u> </u>
(V) ANALYTICAL RESULTS /e.z co	onsistent with self-monitorin	g report data)	S YES		EN.
b) MONITORING RECORDS (e.g., flow, pH	I, D.O., etc.) MAINTAINED	FOR A MINIMUM OF THREE YEARS	5		
INCLUDING ALL ORIGINAL STRIP.CH	HART RECORDINGS (e.g. c	ontinuous monitoring instrumentation	ı .		
canbration and maintenance records).		! 	YES		<u> </u>
ABEQUIPMENT CALIBRATION AND	MAINTENANCE RECORD	DS KEPT.	K YES		Ω N.
S) FACILITY OPERATING RECORDS KE	PT INCLUDING OPERATIO	NG LOGS FOR EACH TREATMENT UNI	T. C YES	O NO	X N.,
DUALITY ASSURANCE RECORDS KEI	PT.		YES		<u> </u>
BECORDS MAINTAINED OF MAJOR C	ONTRIBUTING INDUSTRI	ES (and their compliance status) USING	3		
PUBLICLY OWNED TREATMENT WOR	KS				×N/A
ECTION H - Permit Verification	•	•			
NSPECTION OBSERVATIONS VERIFY TH	HE PERMIT. DYES	ONO ON/A (Further explanatio	on attached	;	
DETAILS:	· · · ·				
B CORRECT NAME AND MAILING ADDR	RESS OF PERMITTEE.		X YES		ŪN/2
) FACILITY IS AS DESCRIBED IN PERM	ит.		K YES		
PRINCIPAL PRODUCTISI AND PRODU	CTION RATES CONFORM	WITH THOSE SET FORTH IN PERMI	T		
APPLICATION.			VES YES		⊡ N ′2
S) TREATMENT PROCESSES ARE AS DE	SCRIBED IN PERMIT APPL	ICATION.			<u> </u>
) NOTIFICATION CIVEN TO EPA/STATE	E OF NEW, DIFFERENT OF	INCREASED DISCHARGES.	VES		<u> </u>
ACCUPATE RECORDS OF RAW WATE	R VOLUME MAINTAINED.		V YES		X.
NUMBER AND LOCATION OF DISCHA	RGE POINTS ARE AS DES	CRIBED IN PERMIT.	S YES		<u> </u>
CORRECT NAME AND LOCATION OF	RECEIVING WATERS.		K YES		<u> </u>
ALL DISCHARGES ARE PERMITTED.			YES		
ECTION I - Operation and Maintenance		·	•		
REATMENT FACILITY PROPERLY OPER	RATED AND MAINTAINED	. I YES I NO X N/A (FI	urther explana	tion attached	t,
STANDBY POWER OR OTHER FOLINA					
ADEQUATE ALARY SYSTEM FOR POL	WER OR FOUNDATENT CAN				<u></u>
REPORTS ON ALTERNATE SCHOOL OF	E POWER SENT TO SAL	TATE AS BEOLUGED AN ATTENT			<u> </u>
SLUDGES AND SOLIDS ADEQUATELY	DISPOSED	ATE AS REQUIRED BY PERMIT.		<u> NO</u>	<u></u>
ALL TREATMENT UNITE IN CERMINE					<u> </u>
CONSULTING ENGINEED DETAILED					<u> </u>
MAINTENANCE PROBLEMS.	DA AVAILABLE FOR CON	SULTATION ON OPERATION AND	[]	D	57
0114115150 005345140 00545					<u> </u>
QUALIFIED OPERATING STAFF FROV	NDED.		II VER		<u> </u>
ESTABLISHED PROCEDURES AVAILA	HDED.	OPERATORS			
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						PERMI	T NO. 1000007	8
SECTION J · Compliance	Schedules	•	•					
PERMITTEE IS MEETIN	G COMPLIANCE SCHEDULE.	1 YES		XN/A	(Further c.	planation a	tieched	
CHECK APPROPRIA	TE PHASE(S):							•
()) THE PERMITT	EE HAS OBTAINED THE NECESSARY TO BEGIN CONSTRUCTION.	YAPPROVAL	SFROM	THE APPR	OPRIATE	•		
(b) PROPER ARR.	ANGEMENT HAS BEEN MADE FOR F	INANCING (n	nortgage c	omnitmen	ts, grants, etc	.). .		
C (c) CONTRACTS	FOR ENGINEERING SERVICES HAVE	BEEN EXEC	UTED.		ن			•
(d) DESIGN PLAN	S AND SPECIFICATIONS HAVE BEEN	N COMPLETE	>.		_			
C (e) CONSTRUCTIO	DN HAS COMMENCED.	• •		•	•			
	ON AND/OR EQUIPMENT ACQUISITIC	ON IS ON SCH	EDULE.					
	ON HAS BEEN COMPLETED.			ţ		•	·	·
(h) START-UP HA	S COMMENCED.			. '	ļ			
(i) THE PERMITT	EE HAS REQUESTED AN EXTENSION	N OF TIME.	•				•	
SECTION K - Self-Monito	ring Program			· · · · · · · · · · · · · · · · · · ·		· · · · ·		
Part I - Flow measure	ment (Further explanation attached	,			•	•		
PERMITTEE FLOW MEA DETAILS:	SUREMENT MEETS THE REQUIREM	ENTS AND IN	TENT O	F THE PEF	RMIT.	YES	D NO	□ Ni
(.) PRIMARY MEASURI	NG DEVICE PROPERLY INSTALLED.					VES YES		
TYPE OF DEVICE	: OWER OPARSHALL FLUME			VENTURI	METER	OTHER IS	pecity Conf	ice
(b) CALIBRATION FREC	IVENCY ADEQUATE. (Date of last calif	bration			/	S YES	I NO	Ξ.
C PRIMARY FLOW ME	ASURING DEVICE PROPERLY OPERA	ATED AND M	AINTAIN	ED.		VES		Ξ.
INSECONDARY INSTRU	IMENTS (totalizers, recorders, etc.) PRG	PERLY OPER	ATED A	ND MAINT	AINED.	X YES		<u> </u>
IN FLOW MEASUREMEN	TEQUIPMENT ADEQUATE TO HAND	DLE EXPECTI	D RANG	ES OF FL	OW RATES.	YES		<u> </u>
Part 2 – Sampling (Fur	ther explanation attached _ X	•			•			
PERMITTEE SAMPLING DETAILS:	MEETS THE REQUIREMENTS AND IN	NTENT OF TH	E PERMI	т.	•			⊡n.,
a) LOCATIONS ADEQUA	ATE FOR REPRESENTATIVE SAMPLE	ES.						<u> </u>
b) PARAMETERS AND S	AMPLING FREQUENCY AGREE WITH	H PERMIT.		•••••••••••••••••••••••••••••••••••••••		S YES		
C) PERMITTEE IS USING	METHOD OF SAMPLE COLLECTION	I REQUIRED	BY PERM	IT.	NCY	VES YES	5 NO	En.
d) SAMPLE COLLECTIO	N PROCEDURES ARE ADEOUATE.					VYES	X NO	
(i) SAMPLES REFI	RIGERATED DURING COMPOSITING					VES .	<u> 2 </u> \c	<u> </u>
(ii) PROPER PRESE	RVATION TECHNIQUES USED	·				VES		Ξ、.
(iii) FLOW PROPOR	TIONED SAMPLES OBTAINED WHER	E REQUIRED	BY PER	літ <u> </u>		O YES	X NO	2
(IV) SAMPLE HOLD	ING TIMES PRIOR TO ANALYSES IN	CONFORMAN	CE WITH	40 CFR 1	36.3	X YES		Ξ× 2
PERMIT.	NALYSES BEING PERFORMED MORE	E FREQUENT	LY THAN	REQUIR	ED BY	—	N	<u> </u>
I) IF (e) IS YES, RESULT	SARE REPORTED IN PERMITTEE'S	SELE-MONIT	BING B	EPORT				
				L: U////.	· ••			- 10 N
ERMITTEE LABORATO	Priner explanation attached					- ·	_	_
DETAILS:	AT PROCEDURES MEET THE REQUI	REMENTS AN	ID INTEN	T OF THE	PERMIT.	DA YES		ŪN/J
) EPA APPROVED ANA	LYTICAL TESTING PROCEDURES US	SED. 1-10 CFR	136.31	•••••	• • •	VES		
b) IF ALTERNATE ANA	LYTICAL PROCEDURES ARE USED, I	PROPER APPI	ROVAL H	AS BEEN	OBTAINED.	VES	I NO	X
PARAMETERS OTHER	R THAN THOSE REQUIRED BY THE P	PERMIT ARE	ANALYZ	ED.		T YES	B 10	Ξ\.
SATISFACTORY CAL	IBRATION AND MAINTENANCE OF I	INSTRUMENT	S AND E	QUIPMEN	т.	🛛 YES	<u> </u>	Ξ
UALITY CONTROL	PROCEDURES USED.					VES YES	C NO	Ξ.
U DUPLICATE SAMPLES	ARE ANALYZED & OF TH	IME.	 	• • • • • • • • • •		YES	C NO	
ONNERCIAL LAR	E USED % OF TIME.					YES		<u> </u>
COMMERCIAL LABOR	TATORY STATE CERTIFICE					U YES	<u> </u>	<u> </u>
	TOTT STALE CEMPIFIED:					U YES		<u> </u>
LAB NAME				• • • •	, 			
	•							

EPA FORM 3560-3 (9-77)

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				<u> </u>		PERMIT NO	0078
OUTFALL NO.	OIL SHEEN	er Observations (GREASE	TURBIDITY	VISIBLE FOAM	VISIBLE FLOAT SOL	COLOR	OTHER
001	None	None	None	None	.None	None	
				·			
				•			·.
	N.L. Effluent/Receiving Water Observations (Further explanation attached						
	e.						
SAMPLE OB MPOSITING FF MPLE REFRIGI MPLE REPRESI	TAINED FROM F REQUENCY ERATED DURING ENTATIVE OF VG	ACILITY SAMP	LING DEVICE	PRE:]NO RGE			
CTION N - Anal	ytical Results (Att	ach report if nec	essary)			•	······································
			•	•	•	•	
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