SOIL CLEAN-UP REPORT FOR 1829 CLEMENT AVENUE ALAMEDA, CALIFORNIA



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RE: SOIL CLEAN-UP REPORT

July 18, 1990

KE1179-1, 16314

1829 CLEMENT AVENUE ALAMEDA, CALIFORNIA

Dear Mr. Smith:

Kaldveer Associates is pleased to submit our soil clean-up report for the property at 1829 Clement Avenue, in Alameda, California. The enclosed report contains a description of our investigation, results of soil sample analyses, and our conclusions regarding clean-up of contaminated soils at the site.

We appreciate the opportunity to provide services to you on this project and trust this report meets your needs at this time. If you have any questions or require additional information, please don't hesitate to call.

Very truly yours,

KALDVEER ASSOCIATES, INC.

Dennis Laduzińsky, C.E.G.

Senior Engineering Geologist

David F. Hoexter, C.E.G./R.E.A. Environmental/Geological Services

Associate

DL/DFH:pv

Copies: Addressee (4)

Alameda County Department of Environmental Health (2)

Attention: Mr. Ariu Levi

Alameda Marina (1)

Attention: Mr. Wayne Milani

SOIL CLEAN-UP REPORT

FOR 1829 CLEMENT AVENUE ALAMEDA, CALIFORNIA

To Mr. Loren Smith 3527 Magnolia Drive Alameda, California

By

KALDVEER ASSOCIATES, INC.

Dennis Laduzinsky, C.E.S. Senior Engineering Geologist

David F. Hoexter, C.E.G./R.E.A. Environmental/Geological Services Associate

July, 1990



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SOIL CLEAN-UP REPORT FOR 1829 CLEMENT AVENUE ALAMEDA, CALIFORNIA

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I. INTRODUCTION

This report presents the results of a contaminated soil clean-up program performed at 1829 Clement Avenue in Alameda, California. The site location is shown on the Site Vicinity Map, Figure 1, and the layout of the site is presented on the Site Plan, Figure 2. purpose of this clean-up program has been to contaminated soil, sediment, and other debris from beneath the existing building at the site. The scope of the clean-up program is based on the results of Kaldveer Associates' previous investigations at the site, as outlined in our Soil Testing Report er 1829 Clement Avenue, dated April 4, 1990. The results of that investigation indicated that significant quantities of cyanide or metals were restricted to the top 3 to 6 inches of soil beneath the former etch-process areas at the western portion of the building, and to a dried sediment on top of the asphalt paving located beneath the eastern portion of the building. In accordance with clean-up recommendations outlined in the report, Kaldveer submitted to the Alameda County Department of Environmental Health (ACDEH) a Work Plan for Partial Site Clean-Up, dated March 12, 199 Although this work plan detailed a clean-up program only for the asphalt area beneath the eastern portion of the building, Kaldveer met with representatives of ACDEH on April 16, 1990 to discuss clean-up of the entire site. Subsequent to that meeting, Mr. Ariu Levi of ACDEH issued a Site Remediation letter dated April 17, 1990 that outlined clean-up requirements at the site based on the information contained in Kaldveer's April 4, 1990 report.

The clean-up program described in this report included highefficiency vacuuming of the asphalt pavement surface beneath the
eastern portion of the building and hand-removal of contaminated
surface soils at the western portion of the building. The pavement
surface and concrete footings at the eastern end of the building
were sealed with three inches of cement slurry and stucco.
Concrete footings at the western end of the building were sealed
with latex enamel paint. Contaminated materials were removed to
a licensed disposal facility by a licensed hazardous waste hauler.
An investigation of building interior environmental quality was
performed by a Certified Industrial Hygenist.

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2) LIAE CALPLES ATTEN PAINTING T-00 mD . 671 PAL / cm

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ATTER VACUULIUS

Ps .88 mg/ 100 cm

7.9 us/ 100 cm CW PA, ists).

> AFTER PRINTING TOURS EN . 1 m3/100 cm

3" CONCRETE CAT INSTALLED ATTEN CONCRETE FOUND CN . 007 ms/ 100 c2

CONCLUSIONS II.

Based on the information collected during this soil clean-up program, and the results of the laboratory analyses, it is our opinion that contaminated soil clean-up has been performed in accordance with Alameda County Department of Environmental Health Details of the site clean-up are presented by specifications. specific area of concern in the following sections.

In addition to clean-up of contaminated soil at the site, investigation of building interior environmental quality performed by a Certified Industrial Hygenist indicates that the building is safe for human occupancy. A copy of the report is included herein as Appendix II.

Former Etch Process Room Areas - West End of Building

Surficial soils containing significant levels of cyanide and metals have been removed from beneath the former etch process room areas at the west end of the building. Laboratory analysis of nine surficial soil samples collected following removal of the contaminated soil indicates residual cyanide concentrations of from non-detect to 8.3 mg/kg. Analysis of three of these samples for soluble cyanide using the Waste Extraction Test (Title 22, CCR) indicates soluble cyanide concentrations of from 0.18 to 0.78 ppm (mq/1).

Analysis for selected metals did not indicate the presence of detectable quantities of hexavalent chromium or molybdenum. Laboratory analyses indicate the presence of copper in surficial soils at the west end of the building at concentrations of 5.6 to $100 \, \text{mg/kg}$. These concentrations are well below levels of significant environmental concern.

Following removal of contaminated soils from beneath the west end of the building, concrete footing surfaces were sprayed with latex enamel paint to provide a surface seal. Three of four wipe samples from the painted surface did not contain detectable quantities of cyanide. The fourth sample contained cyanide at 0.071 mg/100 cm2 In our opinion, the latex paint provides an of footing surface. effective seal on the concrete surfaces.

Former Treatment and Storage Room Area - East End of Building

Clean-up beneath the former treatment and storage room area at the east end of the building consisted of high-efficiency vacuuming of the concrete footings and asphalt paving surface underlying this portion of the building. Wipe samples collected from the footing and asphalt paving surfaces following vacuuming indicated the presence of lead at from 0.0088 to 0.88 mg/100 cm², molybdenum from non-detect to 0.23 mg/100 cm², copper at 0.0051 to 0.16 mg/100 cm²,

air

arsenic from 0.019 to 1.6 mg/100 cm², and cyanide at from 0.079 to 7.9 mg/100 cm². Following receipt of the chemical data, two coats of latex enamel paint were applied to the footings and asphalt paving surface to act as a sealant. Analysis of six subsequent wipe samples for cyanide indicate the presence of cyanide from 0.0058 to 0.10 mg/100 cm².

Finally, a three-inch concrete cap was placed over the asphalt surface, and metal lath covered by stucco was applied to the footing walls to provide a final seal. Arsenic and lead were not detected on six subsequent wipe samples. Cyanide was detected at 0.0050 and 0.0070 mg/100cm² on two wipe samples collected from the slab surface. Cyanide was not detected on four wipe samples collected from the footing surfaces. The measured concentrations of cyanide on the slab surface are at or near the laboratory detection limit. The cyanide is probably related to disturbance of the asphalt surface during placement of the concrete. In our opinion, these levels do not represent a significant environmental concern.

Soil samples collected from beneath the asphalt paving during the initial previous site investigation containing greater than 6 ppm total cyanide were analyzed for soluble cyanide using the WET method. Whereas total cyanide levels in these samples ranged from 7.0 to 160 mg/kg, measured soluble cyanide levels were reported to range from 0.24 to 5.8 ppm (mg/l). The results indicate that inplace soluble cyanide levels are below the Alameda County Department of Environmental Health specified level of 6 ppm (mg/l) based on the WET method analysis.

C. Contaminated Soil Disposal

Approximately 28 cubic yards of contaminated soil, along with the former process piping and stained wood flooring materials were removed from the site by a licensed waste hauler. Contaminated materials were transported under manifest to a licensed disposal of facility.

III. SCOPE OF SERVICES

The scope of work performed during this study consisted of the following tasks:

- 1. Preparation of site clean-up program specifications.
- Selection of remediation contractors.
- Observation of the site clean-up operations.

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- 4. Collection of soil and wipe samples to confirm clean-up effectiveness.
- 5. Analysis of samples by a contract analytical laboratory.
- 6. Analysis of building interior environmental quality by a Certified Industrial Hygenist.
- 7. Preparation of this report.

IV. CONTAMINATED SOIL CLEAN-UP PROGRAM A. Site Clean-Up Specifications

Clean-up levels for contaminated soils at the site were specified by ACDEH at the April 16, 1990 meeting at ACDEH offices. Based on toxicological considerations, ACDEH specified a soil clean-up level for cyanide of 6 parts per million as soluble cyanide. ACDEH specified that soluble cyanide concentration be determined by the Waste Extraction Test (WET) method as outlined in Title 22 of the Code of California Regulations (CCR). ACDEH also specified that acceptable levels of metals in soil be evaluated in accordance with Title 22 CCR specifications.

In their April 17, 1990 Site Remediation letter, ACDEH lists as acceptable, Kaldveer's recommendation to remove the top six inches of contaminated soil beneath the western portion of the building as the initial step to site clean-up. In addition, at the April 16, 1990 meeting, ACDEH approved the clean-up of the eastern portion of the building as outlined in Kaldveer's March 12, 1990 Work Plan for Partial Site Clean-Up.

B. Contaminated Soil Clean-Up - Etch Process Room Area, West End of Building

Clean-up of contaminated soil beneath the former etch process rooms at the western portion of the building was performed by IT Corporation of Martinez, California between April 16 and 20, 1990. Clean-up consisted of the manual removal of approximately 6 inches of surface soil from the area shown on Figure 3. Contaminated soil was hand-excavated by shovel, placed in buckets and transferred to a roll-off bin stored on the site. The thickness of contaminated soil beneath this portion of the building varied from about 3 inches to over 1 foot, and was easily identified on the basis of discoloration; contaminated soils were black to blue-green in color, whereas uncontaminated soils (as verified by analytical testing) were brown in color. Approximately 6 inches of non-discolored soil were also removed within an approximate 5-foot radius of former soil Boring B-6 (previously found to contain high lead at the 0.5 foot depth).

A hand auger was used to probe the soil surface at up to 20 locations within each of the three former etch-process rooms to check for soil discoloration to evaluate removal effectiveness. Following removal of the discolored soil, three soil samples were collected within 3 inches of the surface from each of three etch-process rooms. The nine samples were analyzed for hexavalent chromium, molybdenum, copper, and total cyanide. In addition, any sample containing greater than 6 ppm total cyanide was analyzed for soluble cyanide using both the Title 22 Waste Extraction Test and a similar extraction process test using deionized water. Sample locations are shown on Figure 3.

Soil sample analytical results are listed on Table 1 and are attached to this report as Appendix I. The results indicate that following initial removal of the discolored soil from beneath the western portion of the building, residual concentrations of copper at the surface range from 5.6 to 100 ppm, with the exception of Sample CS-9 which was found to contain 360 ppm. Analysis of sample CS-9 by the WET method indicated the sample contained 27 mg/l soluble copper. As this level exceeds the State STLC of 25 mg/l, additional soil was removed from this area on June 15, Analysis of three additional closure samples (CS-11, CS-12, CS-13) indicated the presence of 17 to 22 ppm total copper. In light of the copper analysis results from this investigation and the results from samples collected from the 0.5 to 6.0 foot depth outlined in our April 4, 1990 report (6.0 to 52 ppm), it is our opinion that soils containing significant levels of copper have been effectively removed from beneath the building. Molybdenum and hexavalent chromium were not detected in any of the closure samples analyzed.

Total cyanide analyses indicate residual cyanide levels of from non-detect to 8.3 ppm in surficial soils remaining beneath this portion of the building. The three samples containing greater than 6 ppm total cyanide were analyzed for soluble cyanide using the WET Soluble cyanide analyses on these samples indicated the presence of 0.18 to 0.78 ppm (milligrams per liter, mg/l) soluble cyanide when analyzed in accordance with the Title 22 WET method, and from non-detect to 1.7 ppm (mg/l) soluble cyanide when analyzed The three analyses do not with a deionized water extraction. indicate any consistent relationship between the two analytical In either case, residual concentrations of soluble methods. cyanide following excavation of discolored soil from beneath the former etch-process rooms are less than the ACDEH specified cleanup level of 6 ppm soluble cyanide.

Soil Sample CS-10 was collected from the ground surface following removal of about 6 inches of soil within an approximate five foot radius of previous Boring B-6. As outlined in our April 4, 1990 Soil Testing Report, a sample collected from a depth of 0.5 feet at this location was found to contain 1100 ppm lead. Sample CS-10, collected during this investigation, following removal of

approximately 6 inches of surface soils this area, was found to contain 12 ppm lead, indicating that soil in this area containing elevated levels of lead has been successfully removed from beneath the building.

C. Concrete Footing - Etch Process Room Areas, West End of Building

Two initial wipe samples were collected from the surface of the concrete footings at the western portion of the building following removal of contaminated soils. Wipe samples were collected by saturating a 9.0 cm diameter filter pad with deionized water and wiping a 10 cm by 10 cm area with both sides of the filter pad. One wipe sample was analyzed for hexavalent chromium, copper, and molybdenum. The second wipe sample was analyzed for cyanide. As shown on Table 2, hexavalent chromium and molybdenum were not found to be present in detectable quantities. Results for copper and cyanide are reported as 67 and 8.0 mg/100 cm², respectively.

Following the removal of contaminated soils and collection of wipe samples from the footing surfaces, the exposed footings were sprayed with latex enamel paint to provide a surface seal on the concrete. Following application of the paint sealer, four wipe samples were collected from the footing surface and analyzed for total cyanide. The analytical results, shown as Samples CFS-1 through CFS-4 on Table 2, indicate that three of the four wipe samples did not contain cyanide in detectable quantities. One of the samples contained 0.071 mg cyanide per 100 cm². In our opinion, application of the latex enamel paint has provided an effective seal on the concrete footing surface.

D. Site Clean-Up - Treatment and Storage Room Areas, East End of Building

Site clean-up beneath the former treatment and storage room areas at the eastern end of the building consisted of high-efficiency vacuuming of the underlying asphalt paving surface and adjacent concrete footing surfaces. Site clean-up was conducted between April 16 and May 24, 1990 by IT Corporation of Martinez, California. Vacuuming of the asphalt and concrete footing surfaces was performed using high-efficiency vacuum canisters of the type commonly used for asbestos removal operations.

Six initial wipe samples were collected from the footing and asphalt paving surfaces following the vacuuming by saturating a 9.0 cm diameter filter paper with deionized water and wiping down a 10 cm X 10 cm area using both sides of the filter pad. The location of the three samples collected from the footing walls and three samples collected from the asphalt paving surface are shown on Figure 4 (samples designated WS for asphalt paving samples and FS for footing wall samples). The samples were analyzed for

cyanide, arsenic, beryllium, copper, molybdenum and lead. Analytical results are presented on Table 3. The results indicated the presence of all constituents except beryllium.

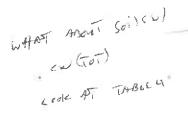
Following receipt of the wipe sample analytical data, the footing walls and asphalt paving surface beneath the building were sealed with two coats of latex enamel paint. Analytical results for cyanide analyses for six footing and asphalt surface wipe samples collected after application of the latex paint sealer are also listed on Table 3 (sample designation CFS on Figure 4). The results indicate the presence of only cyanide in all six samples.

Finally, a three-inch thick concrete topping slab was applied to the asphalt paving surface, and steel lath covered by stucco was applied to the footing stem walls. Analysis of final wipe samples (designated WPC on Figure 4) indicate that lead and arsenic are not present on the concrete or stucco surfaces above the laboratory detection limits. Cyanide was not detected on the four footing wall samples but was present in the two concrete slab samples at concentrations of 0.007 and 0.005 mg/100cm². These concentrations are at, or just barely exceed the laboratory detection limit of 0.0050 mg/wipe. The cyanide is probably related to disturbance of the asphalt surface during concrete placement. In our opinion, these levels do not represent a significant environmental concern, and it appears that the concrete and stucco toppings provide an effective seal of residual cyanide or metals on the asphalt and footing surfaces beneath this portion of the building.

In addition to analysis of wipe samples, soil and sediment samples collected during the initial investigation phase were reanalyzed for soluble cyanide using the Waste Extraction Test as specified by ACDEH. A variety of samples collected from both the eastern and western portions of the building were analyzed in an attempt to establish a relationship between total cyanide and soluble cyanide. As indicated on Table 4, no straight-forward relationship is apparent. However, it should be noted that based on the analytical results, all soil or sediment found to contain hazardous levels of metals, or greater than 6 ppm soluble cyanide as determined by the WET, has been removed from the site in accordance with ACDEH requirements.

E. Disposition of Contaminated Soil and Debris

Approximately 28 cubic yards of contaminated soil was removed from the site on May 29, 1990 by Sturgeon and Son Inc. of Bakersfield, California, a licensed waste hauler. In addition to removal of the contaminated soil, all former process piping and stained wood flooring removed from the building during the site clean-up were removed from the site. Contaminated soil and other debris were



transported under manifest to Envirosafe Services of Idaho, a licensed hazardous waste disposal facility.

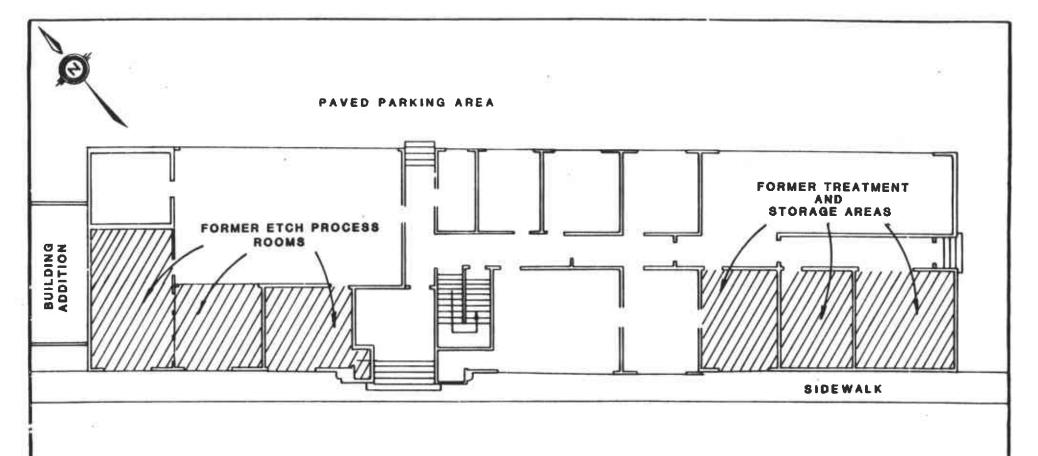
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KE1179-1

July 1990

Figure 1



Clement Avenue -

APPROXIMATE SCALE IN FEET

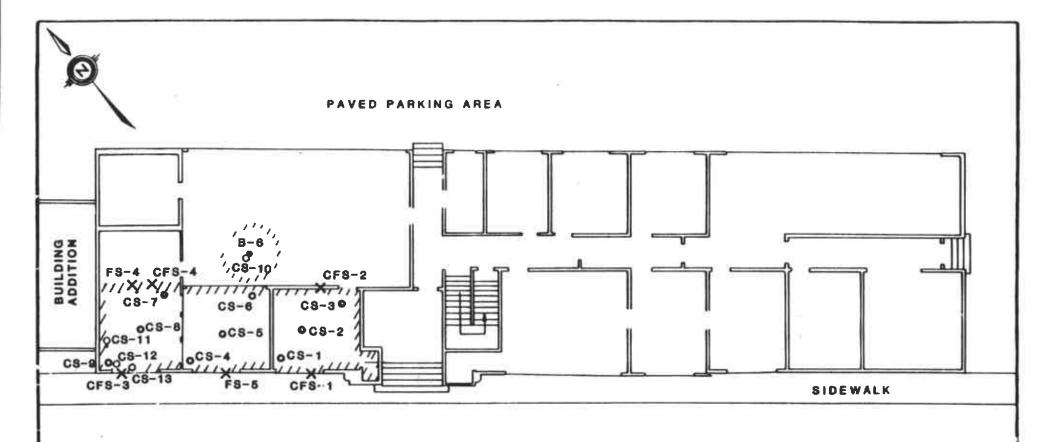


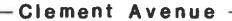


SITE PLAN

1829 CLEMENT AVENUE Alameda, California

PROJECT NO	ROJECT NO DATE		2
KE1179-1	July 1990	Figure	~





LEGEND

SOIL REMOVAL AREA ENCLOSED BY CROSS- HATCHING

B-6.

PREVIOUS BORING B-6

C8-5

LOCATION OF SURFACE

SOIL SAMPLE

F8-3 X CFS-3

FOOTING SURFACE WIPE SAMPLE LOCATIONS



Kaldveer Associates Geoscience Consultants

A California Corporation

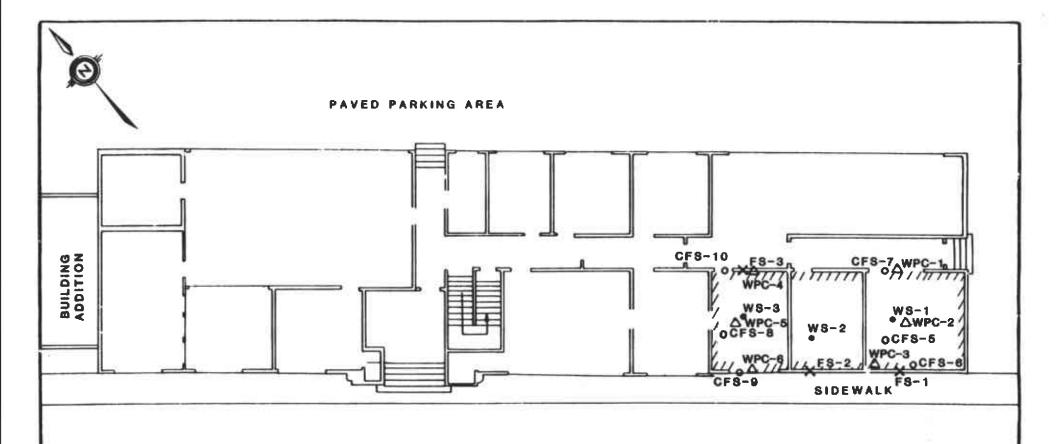
APPROXIMATE SCALE IN FEET



FORMER ETCH PROCESS ROOMS SOIL REMOVAL AREA

1829 CLEMENT AVENUE Alameda, California

PROJECT NO	DATE	Sie 3
KE1179-1	July 1990	Figure 3



Clement Avenue

LEGEND

APPROXIMATE SCALE IN FEET



77777 SEDIMENT REMOVAL AREA ENCLOSED BY CROSS-HATCHING

FINAL WIPE SAMPLE LOCATIONS, ASPHALT SURFACE AND FOOTING WALLS

WS-1 WIPE SAMPLE LOCATION

FROM ASPHALT PAVING

F8-1 WIPE SAMPLE LOCATION FROM FOOTING WALL

CFS-6 FOLLOW-UP WIPE SAMPLE LOCATIONS, ASPHALT SURFACE AND FOOTING WALLS



Kaldveer Associates Geoscience Consultants

A California Corporation

FORMER TREATMENT AND STORAGE ROOM SEDIMENT REMOVAL AREA

> **1829 CLEMENT AVENUE** Alameda, California

PROJECT NO	DATE	- A
KE1179-1	July 1990	Figure 4

TABLE 1

SOIL SAMPLE ANALYTICAL RESULTS
FORMER ETCH PROCESS ROOM AREA - WEST END OF BUILDING

						Soluble	Cyanide
			Hexavalent		Total	Title 22	- DI
Sample	Copper	Molybdenum	Chromium	Lead	Cyanide	WET	Water
Number	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/1)	(mg/l)
CS-1	36	ND	ND		2.1		
CS-2	5.6	ND	ND		8.3	0.78	0.41
CS-3	9.7	ND	ND		1.4		
_ CS-4	28	ND	ND		1.3		
CS-5	97	ND	ND		2.8		
CS-6	100	ND	ND		6.5	0.49	ND
CS-7	55	ND	ND		6.3	0.18	1.7
CS-8	91	ND	ND		ND		
CS-9	360	ND	ND '		ND	27	
CS-10				12			
CS-11	17						
CS-12	21						
CS-13	22						

TABLE 2

CONCRETE FOOTING WIPE SAMPLE RESULTS FORMER ETCH PROCESS ROOM AREA - WEST END OF BUILDING (all results reported as mg/100 cm²)

1. Samples collected prior to application of paint seal.

Sample Designation	Cyanide	Chromium VI	Copper	Molybdenum
FS-4		ND	67	ND
FS-5	8.0			
Detection Limit	1.0	0.050	0.50	2.5

2. Samples collected after application of paint seal.

Sample Designation	Cyanide
CFS-1	ND
CFS-2	ND
CFS-3	0.071
CFS-4	ND
Detection Limit	0.020

TABLE 3

CONCRETE FOOTING AND ASPHALT PAVING WIPE SAMPLES FORMER TREATMENT ROOM AREA- EAST END OF BUILDING (all results reported as mg/100 cm²)

1. Samples collected prior to application of paint seal.

Sample						
Designation	CN	As	Ве	Cu	Mo	Pb
WS-1	1.7	0.24	ND	0.096	0.23	0.88
WS-2	7.9	0.064	ND	0.16	0.11	0.031
WS-3	2.0	1.6	ND	0.078	0.059	0.053
FS-1	0.079	0.099	ND	0.026	ND	0.013
FS-2	1.6	0.019	ND	0.033	ND	0.0088
FS-3	0.83	0.25	ИĎ	0.0051	0.033	0.010
Detection						
Limit	0.02	0.0005	0.001	0.001	0.001	0.0005

2. Samples collected after application of paint seal.

Sample	
Designation	Cyanide
CFS-5	0.087
CFS-6	0.0067
CFS-7	0.10
CFS-8	0.0058
CFS-9	0.0065
CFS-10	0.0065
Detection Limit	0.005

3. Samples collected after application of concrete seal and stucco.

Sampler			
Designation	As	Pb	CN
WPC-1	ND	ND	ND
WPC-2	ND	ND	0.007
WPC-3	ND	ND	ND
WPC-4	ND	ND	ND
WPC-5	ND	ND	0.005
WPC-6	ND	ND	ND
WPC-B (Blank)	ND	ND	ND
Detection			
Limit	0.005	0.005	0.005

SUMMARY OF TOTAL vs. TABLE 4
SOLUBLE CYANIDE TEST RESULTS

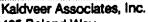
Sample Designation	Total Cyanide (mg/kg)	Soluble Cyanide+ (mg/l)
B1,B2-0.5*	160	4.2
B1,B2-3*	24	5.8
B3-0.5*	26	2.2
B3-6*	22	0.24
B4-0.5*	7.1	0.05
S-2**	510	8.8
S-3**	120	2.7
S-4**	650	2.0
S-7**	1100	18

Notes:

- + = Soluble cyanide determined by Waste Extraction Test, CCR Title 22.
- * = These soils left in-place beneath asphalt at east end of building.
- ** = These soils removed from the west end of building.



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233



425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

#KE1179-1, Clement

Sample Descript: Analysis for:

First Sample #:

Wipes Cyanide

004-2429

Sampledo R Received:

Reported:

Apr 21, 1990

LABORATORY ANALYSIS FOR:

Cyanide

Sample Number	Sample Description	Detection Limit mg/100 cm ²	Sample Result mg/100 cm²
0042429 A-B	WS-1	0.020	1.7
0042430 A-B	WS-2	0.020	7.9
0042431 A-B	WS-3	0.020	2.0
0042432 A-B	FS-1	0.020	0.079
0042433 A-B	FS-2	0.020	1.6
0042434 A-B	FS-3	0.020	0.83

Analytes reported as N.D. were not present above the stated limit of detection.

aurer

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

42429.KAL <1>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

#KE1179-1, Clement

Sample Descript:

Analysis for: First Sample #:

Wipes Arsenic

004-2429

Sampled:

Apr 17, 1990 Apr 17, 1990

Received:

Reported: Apr 21, 1990

LABORATORY ANALYSIS FOR:

Arsenic

Sample Number	Sample Description	Detection Limit mg/100 cm²	Sample Result mg/100 cm²
004-2429	WS-1	0.0005	0.24
004-2430	WS-2	0.0005	0.064
004-2431	WS-3	0.0005	1.6
004-2432	FS-1	0.0005	0.099
004-2433	FS-2	0.0005	0.019
004-2434	FS-3	0.0005	0.25

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

) lane Diane Elich Lawvel Project Manager

42429.KAL <2>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

oject ID: #KE1179-1, Clement

Sample Descript: Analysis for: First Sample #:

Wipes Beryllium · 004-2429 Sampled:

Apr 17, 1990

Received:

Apr 17, 1990

Reported:

Apr 21, 1990

LABORATORY ANALYSIS FOR:

Beryllium

Sample Number	Sample Description	Detection Limit mg/100 cm ²	Sample Result mg/100 cm²
004-2429	WS-1	0.0010	N.D.
004-2430	WS-2	0.0010	N.D.
004-2431	ws-3	0.0010	N.D.
004-2432	FS-1	0.0010	N.D.
004-2433	FS-2	0.0010	N.D.
004-2434	F\$-3	0.0010	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

auver

SEQUOIA ANALYTICAL,

lane

Diane Elich Lawyer Project Manager

42429.KAL <3>



Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

#KE1179-1, Clement

Sample Descript: Wipes Analysis for:

Copper First Sample #: 004-2429 Sampled:

Apr 17, 1990

Received:

Apr 17, 1990

Reported: Apr 21, 1990

LABORATORY ANALYSIS FOR:

Copper

Sample Number	Sample Description	Detection Limit mg/100 cm ²	Sample Result mg/100 cm²
004-2429	WS-1	0.0010	0.096
004-2430	WS-2	0.0010	0.16
004-2431	ws-3	0.0010	0.078
004-2432	FS-1	0.0010	0.026
004-2433	FS-2	0.0010	0.033
004-2434	FS-3	0.020	0.0051

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

42429.KAL <4>

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Remarks:												425 Oal	Rola	nd Wa Cali	iy fornia	Koldveer Associal Geoccience Consulta ACaltimus Corporation

Yellow - Analytical Laboratory

White - Kaldveer Associates



Kaldveer Associates, Inc.

425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

Lab Number:

#KE1179-1, Clement

Soil, CS-2

Sampled:

Apr 18, 1990

Sample Descript:

004-2616

Received:

Apr 18, 1990

Analyzed: Reported:

Apr 19, 1990 Apr 21, 1990

LABORATORY ANALYSIS

Analyte **Detection Limit** Sample Results mg/kg mg/kg

Cyanide	1.0		5.3
Hexavalent Chromium	0.050		N.D.
Molybdenum	2.5	*****************************	N.D.
Copper	0.50		5.5

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Client Project ID:

#KE1179-1, Clement

Sampled:

Apr 18, 1990

425 noiand way Oakland, CA 94621 Sample Descript:

Soil, CS-1

Received:

Apr 18, 1990

Attention: Dennis Laduzinski

Lab Number:

Analyzed: Reported: Apr 19, 1990 Apr 21, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit mg/kg

Sample Results mg/kg

Cyanida	1.0	******************************	1.4
Hexavalent Chromium	0.050	***************************************	N.D.
Molybdenum	2.5		N.D.
<u> Соружника и принципалника и</u>		***********************	9.7

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Lane

Diane Elich Lawver¹ Project Manager



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID: Sample Descript:

Lab Number:

#KE1179-1, Clement

Soil, CS-4

Sampled:

Apr 18, 1990

Received:

Apr 18, 1990

Analyzed:

004-2618

Reported:

Apr 19, 1990 Apr 21, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit

Sample Results mg/kg

mg/kg

Cyarios	1.0	************************	1.3
Hexavalent Chromium	0.050	****************************	N.D.
Molybdenum	2.5	400400400000000000000000000000000000000	N.D.
Соррег	0.50	***********	28

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

lane

Diane Elich Lawver Project Manager

42615.KAL <4>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way

Client Project ID:

#KE1179-1, Clement

Sampled:

Apr 18, 1990

Sample Descript:

Soil, CS-5

Received:

Apr 18, 1990

Oakland, CA 94621

Attention: Dennis Laduzinski Lab Number: 004-2619

Analyzed: Reported:

Apr 19, 1990 Apr 21, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit mg/kg

Sample Results mg/kg

Cyanida	1.0	************************	2.8
Hexavalent Chromium	0.050	***************************************	N.D.
Molybdenum	2.5	************************	N.D.
Copper	0.59		97

Analytes reported as N.D. were not present above the stated limit of detection.

wow

SEQUOIA ANALYTICAL

ane

Diane Elich Lawver **Project Manager**

42615.KAL <5>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, inc. 425 Roland Way

Client Project ID:

#KE1179-1, Clement

Sampled:

Apr 18, 1990

Oakland, CA 94621

Sample Descript:

Soil, CS-6

Received:

Apr 18, 1990

Attention: Dennis Laduzinski

Lab Number:

004-2620

Analyzed: Reported: Apr 19, 1990 Apr 21, 1990

LABORATORY ANALYSIS

Detection Limit Sample Results Analyte mg/kg mg/kg

Cyanida	1.0	***************************************	5.5
Hexavalent Chromium	0.050	***************************************	N.D.
Molybdenum	2.5	******************************	N.D.
Соррегиничний	0,50	************	100

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

42615.KAL <6>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

Matrix Descript:

Analysis for: First Sample #: #KE1179-1, Clement

Liquid

STLC, Cyanide 0042616 R

Sampled:

Apr 18, 1990

Received: Apr 18, 1990

Reported: Apr 25, 1990

LABORATORY ANALYSIS FOR:

STLC, Cyanide

Sample Number	Sample Description	Detection Limit mg/kg	Date Analyzed	Sample Result mg/kg
0042616 R	CS-2	0.010	4/24/90	0.78
0042620 R	CS-6	0.010	4/24/90	0.49

Method of Analysis:

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

R42616.KAL <2>





680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinski

Client Project ID:

Matrix Descript:

Analysis for: First Sample #: #KE1179-1, Clement

Liquid

Soluble Cyanide

R42616

Reported:

Apr 25, 1990

LABORATORY ANALYSIS FOR:

Soluble Cyanide

Sample Number	Sample Description	Detection Limit mg/kg	Date Analyzed	Sample Result mg/kg
0042616 R	CS-2	1.0	4/24/90	0.41
0042620 R	CS-6	1.0	4/24/90	N.D.

Method of Analysis:

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

R42616.KAL <1>

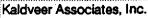
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White - Kaldveer Associates

Yellow - Analytical Laboratory



680 Chesapeake Drive . Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233



425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID:

#KE1179-1, Clement

Sample Descript:

Soil, CS-7

Sampled:

Apr 20, 1990

Receive APR AP 80. 1990

Lab Number:

004-3028

Analyzed:

Apr 24, 1990

Reported:

Apr 25, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit mg/kg

Sample Results mg/kg

Cyanida	1.0	***********	6.3
Hexavalent Chromium	0.050	41344	N.D
Copper	0.60		56
Molybdenum	2.5	******************************	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

43028.KAL <1>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Client Project ID:

#KE1179-1, Clement

Sampled:

Apr 20, 1990

Oakland, CA 94621

Sample Descript:

Soil, CS-8

Received:

Apr 20, 1990

Attention: Dennis Laduzinsky

Lab Number:

004-3029

Analyzed: Reported:

Apr 24, 1990 Apr 25, 1990

LABORATORY ANALYSIS

Analyte	Detection Limit mg/kg		Sample Results mg/kg
Cyanide	1.0	***************************************	N.D.
Hexavalent Chromium	0.050	***************************************	N.D.
Copper	0.63	************************************	. 91
Molybdenum	2.5	***************************************	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver **Project Manager**

43028.KAL < 2>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript: #KE1179-1, Clement Soil, CS-9 Sampled: Received: Apr 20, 1990 Apr 20, 1990

Oakland, CA 94621

Lab Number:

004-3030

Analyzed: Reported: Apr 24, 1990 Apr 25, 1990

LABORATORY ANALYSIS

Analyte	Detection Limit mg/kg		Sample Results mg/kg	
Cyanide	1.0	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	N.D.	
Hexavalent Chromium		***************************************	N.D	
Copper	G(G)		. 360	
Molybdenum	2.5		N.D.	

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawer Project Manager

43028.KAL <3>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Lab Number:

Kaldveer Associates, Inc.

425 Roland Way

Oakland, CA 94621 Attention: Dennis Laduzinsky Client Project ID:

#KE1179-1, Clement

Soil, CS-7

Sampled:

Apr 20, 1990

Sample Descript:

Received:

Apr 20, 1990

004-3028

Analyzed:

May 1, 1990

Reported: May 3, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit mg/L

Sample Results mg/L

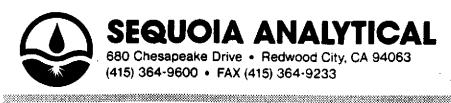
Cyanida, STLO	010 D 18
C/SING, SINGERCOMMINENTAL AND ARTHUR STATE	<u>\ </u>
	010
Soluble Cyanide	VIJ E-E-P-P-F-F-F-F-F-F-F-F-F-F-F-F-F-F-F-F-

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawyer Project Manager

43028.KAL <1>



Kaldveer Associates, Inc. 425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript:

Lab Number:

#KE1179-1, Clement

Soil, CS-10

004-3031

Sampled:

Apr 20, 1990

Received:

Apr 20, 1990

Analyzed: Reported:

Apr 23, 1990 Apr 25, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit mg/kg

Sample Results mg/kg

Lead.....

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

43028.KAL <4>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way

Oakland, CA 94621 Attention: Dennis Laduzinsky Client Project ID:

Lab Number:

#KE1179-1, Clement

Sample Descript: Soil, CS-9

004-3030

Sampled:

Apr 20, 1990

Relogged:

Apr 20, 1990

Analyzed: Reported: Jun 5, 1990

Jun 12, 1990

LABORATORY ANALYSIS

Analyte

Detection Limit

Sample Results

Copper, STLC, mg/L	
Copper, TLC, ng/kg	

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL>

Diane Elich Lawver Project Manager

43028.KAL <2>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript: #KE1179-1, Clement

Sampled: Received: Apr 20, 1990 Apr 20, 1990

Oakland, CA 94621

Lab Number:

Wipes, FS-4.

004-3032

Reported:

Apr 25, 1990

LABORATORY ANALYSIS

Sample Results **Detection Limit** Analyte mg/kg mg/kg N.D. 0.050 Hexavalent Chromium..... 57 Copper..... 2.5 N.D. Molybdenum.....

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

43028.KAL <5>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Lab Number:

Kaldveer Associates, inc. 425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID:

#KE1179-1, Clement

Sample Descript: Soil, FS-5

Sampled:

Apr 20, 1990

Received:

Apr 20, 1990

Analyzed: Reported: Apr 24, 1990 Apr 25, 1990

LABORATORY ANALYSIS

004-3033

Analyte

Detection Limit mg/kg

Sample Results mg/kg

Cyanide.....

87.3

* 37.3

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

43028.KAL <6>



Kaldveer Associates, Inc.

Client Project ID: #KE1179-1, Clement

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinsky

QC Sample Group: 43028 - 31

Reported: Apr 25, 1990

QUALITY CONTROL DATA REPORT

ANALYTE	Hexavalent Chromium	Copper	Molybdenum	Lead	
Method: Analyst: Reporting Units: Date Analyzed: QC Sample #:	EPA 7196 R. Sharma mg/kg Apr 20, 1990 004-3126	EPA 6010 B. Oliver mg/L Apr 23, 1990 004-3029	EPA 6010 B. Oliver mg/L Apr 23, 1990 004-3029	EPA 6010 B. Oliver mg/L. Apr 23, 1990 004-1608	
Sample Conc.:	N.D.	1.8	N.D.	3.6	
Spike Conc. Added:	0.50	1.0	1.0	50	
Conc. Matrix Spike:	0.50	2.9	0.79	56	
Matrix Spike % Recovery:	100	110	79	100	
Conc. Matrix Spike Dup.:	0.50	2.9	0.75	61	
Matrix Spike Duplicate % Recovery:	100	110	75	110	
Relative % Difference:	0.26	0.0	5.2	8.5	

SEQUOIA ANALYTICAL

Diane aurer

Diane Elich Lawyer Project Manager

% Recovery:

Conc. of M.S. - Conc. of Sample x 100

Spike Conc. Added

Relative % Difference:

Conc. of M.S. - Conc. of M.S.D. x 100

(Conc. of M.S. + Conc. of M.S.D.) / 2

43028.KAL <7>

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Yellow - Analytical Laboratory

White - Kaldveer Associates

MID-PACIFIC ENVIRONMENTAL LABORATORY



Kaldveer Associates 425 Roland Way Oakland, CA 94621

MAY 4 1990

April 24, 1990 MPELI ID: 9004074 Client PO: Ke1179-1 Page 1 of 4

Attention: Dennis Laduzinsky

Subject: Analysis of 9 Soil Samples, Received 3/21/90.

The samples were prepared by extracting with a citrate buffer for 48 hours. The resulting extract was filtered and analyzed for Cyanide. The resulting values are the soluble threshold limit concentrations for the requested compound. The results are presented in Table 1.

If you should have any technical questions, please contact the undersigned at (415)964-0844.

Approved by:

M. Claire Ferguson

Client Services Manager

These results were obtained by following standard laboratory procedures; the liability of Mid-Pacific Environmental Laboratory, Inc. shall not exceed the amount paid for this report. In no event shall Mid-Pacific be liable for special or consequential damages.

Table 1. General Chemical Results

Kaldveer Sample ID

		B1,B2 0.5	B1,B2 -3	B3-0.5	B3-6 mg/L	
Parameter	EPA Method	mg/L	mg/L	mg/L		
Cyanides, total	335.2 C	4.2	5.8	2.2	0.24	0.05

C - colorimetric

Table 1. General Chemical Results

Kaldveer Sample ID

	EPA	S2	S3	S4	S 7	Spike
Parameter	Method	mg/L	mg/L	mg/L	mg/L	% Recov
Cyanides, total	335.2 C	8.8	2.7	2.0	18	85

C - colorimetric

Table 1. General Chemical Results

Kaldveer Sample ID

	ED3	Dup	Control Sample	Method Blank	Method Detection Limit
Parameter	EPA Method	RPD	% Recov	mg/L	mg/L
Cyanides, total	335.2 C	17	88.8	<0.01	0.01

C - colorimetric

5682205 P.08

MID-PROIFIC End Lab

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MED-TOX ASSOCIATES, INC. ANALYTICAL REQUEST/CHAIN OF CUSTODY FORM

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Lient	Kaldueer Associates	(Complete Information on Opposite Side)	Date: 2-12-10
LIENT	JOB REF.: KE1171-1		SAMPLER(S): Deunis Ladurinsky.

LAB PROJECT NO: (lab use only) ANALYSES * Tubes with arrows should be analyzed from the arrow end ATR SAMPLE CLIENT SAMPLE DATE Lab Number VOLUME TYPE NO. IDENTIFICATION (leb use only) (Liters) CONT. COMMENTS! 9002 074 INTERFERENC 5-1 2-11-70 014 B1-0.5 Soil 024 XIXXII composite 8 2-0.5 81-3 83-3 1034 82-6.81-6 CHA composite of ; 83-0.5 65% 83-3 CGA 83-6 (D 77) B4-0.5 0874 B4-3 OPA 04-7 LUA X + H19190: Add'l work requested 12 Dhadusins to be done OMPBLE -S

ONE WEEK TAT.

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*SAMPLE TYPE (SPECIFY): (1) 37 mm 0.8 um HCEP; 2) 25 mm 0.8 um HCEP; (3) 25 mm 0.4 um polycarb. filter; (4) PVC filter. diam. pore size ___; (5) Charcoal tube; (6) Silica gel tube (7) Water: (8) Soil: (9) Bolk Samola.



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript:

Clement, KE 1179-1

Wipe

Analysis for: First Sample #: Cyanide 005-1299 Sampled:

May 8, 1990

Received:

May 9, 1990

Reported:

May 15, 1990

LABORATORY ANALYSIS FOR:

Cyanide

Sample Number	Sample Description	Detection Limit mg/wipe	Sample Result mg/wipe
005-1299	CFS-1	0.020	N.D.
005-1300	CFS-2	0.020	N.D.
005-1301	CFS-3	0.020	0.071
005-1302	CFS-4	0.020	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

51299.KAL <1>



Kaidveer Associates, Inc.

Client Project ID: Clement, KE 1179-1

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinsky

QC Sample Group: 0051299-302

Reported: May 15, 1990

QUALITY CONTROL DATA REPORT

ANALYTE Cyanide

Method:

EPA 335,2

Analyst:

A. Chu

Reporting Units:

mg/L

Date Analyzed: QC Sample #:

May 14, 1990 004-3926

Sample Conc.:

N.D.

Spike Conc.

Added:

1.0

Conc. Matrix

Spike:

0.92

Matrix Spike

% Recovery:

92

Conc. Matrix

Spike Dup.:

0.93

Matrix Spike Duplicate

% Recovery:

93

Relative

% Difference:

1.1

SEQUOIA ANALYTICAL

-) rane

Diane Elich Lawver Project Manager % Recovery:

Conc. of M.S. - Conc. of Sample

x 100

Spike Conc. Added

Relative % Difference:

Conc. of M.S. - Conc. of M.S.D.

x 100

(Conc. of M.S. + Conc. of M.S.D.) / 2

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Yellow - Analytical Laboratory

White - Kaldveer Associates



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID:

Sample Descript:

Analysis for:

#KE1179-1, Alameda

Wipe Cyanide

First Sample #: 005-0429

Sampled: May 24, 1990 Received: May 25, 1990

Analyzed: May 30, 1990

Reported: May 31, 1990

LABORATORY ANALYSIS FOR:

Cyanide

Sample Number	Sample Description	Detection Limit mg/wipe	Sample Result mg/wipe
005-0429	CFS - 5	0.0050	0.087
005-0430	CFS-6	0.0050	0.0067
005-0431	CFS - 7	0.0050	0.10
005-0432	CFS - 8	0.0050	0.0058
005-0433	CFS - 9	0.0050	0.0065
005-0434	CFS - 10	0.0050	0.0065

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

50429.KAL <1>



Kaldveer Associates, Inc.

Client Project ID: #KE1179-1, Alameda

425 Roland Way

Oakland, CA 94621

Attention: Dennis Laduzinsky

QC Sample Group: 0050429 - 0050434

Reported: May 31, 1990

QUALITY CONTROL DATA REPORT

ANALYTE Cyanide

Method:

EPA 335.2

Analyst:

A. Chu

Reporting Units:

mg/L

Date Analyzed:

May 30, 1990

QC Sample #:

DI Water

Sample Conc.:

N.D.

Spike Conc.

Added:

0.10

Conc. Matrix

Spike:

0.10

Matrix Spike

% Recovery:

100

Conc. Matrix

Spike Dup.:

0.094

Matrix Spike

Duplicate % Recovery:

94

Relative

% Difference:

6.2

SEQUOIA ANALYTICAL

rane aurver

Diane Elich Lawver Project Manager

% Recovery:

Conc. of M.S. - Conc. of Sample

Spike Conc. Added

Relative % Difference:

Conc. of M.S. - Conc. of M.S.D.

x 100

x 100

(Conc. of M.S. + Conc. of M.S.D.) / 2

50429.KAL <2>

Page of Lab Job #

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680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript: #KE1179-1, Clement - Alameda

Soil

Analysis for: First Sample #: Copper

006-2476

Sampled:

Jun 15, 1990

Received:

Jun 15, 1990:

Reported:

Jun 19, 1990;

LABORATORY ANALYSIS FOR:

Copper

Sample Number	Sample Description	Detection Limit mg/kg	Sample Result mg/kg
006-2476	CS-11	0.50	17
006-2477	CS-12	0.50	21
006-2478	CS-13	0.50	22
006-2479	EX-1	0.50	24
006-2480	EX-2	0.50	35

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

62476.KAL <1>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript:

#KE1179-1, Clement - Alameda

Wipe

Analysis for: First Sample #: Cyanide

006-2481 A - B Sampled: Jun 15, 1990

Received: Jun 15, 1990

Analyzed: Jun 19, 1990 Reported: Jun 19, 1990

LABORATORY ANALYSIS FOR:

Cyanide

Sample Number	Sample Description	Detection Limit mg/wipe	Sample Result mg/wipe	
0062481 A-B	WPC-1	0.0050	N.D.	
0062482 A-B	WPC-2	0.0050	0.0070	
0062483 A-B	WPC-3	0.0050	N.D.	
0062484 A-B	WPC-4	0.0050	N.D.	
0062485 A-B	WPC-5	0.0050	0.0050	
0062486 A-B	WPC-6	0.0050	N.D.	
0062487 A-B	WPC-B	0.0050	N.D.	

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

62476.KAL <2>



Kaldveer Associates, Inc.

425 Roland Way Oakland, CA 94621

Attention: Dennis Laduzinsky

Client Project ID: Sample Descript:

#KE1179-1, Clement - Alameda

Wipe

Analysis for: First Sample #:

Arsenic 006-2481

A-B

Sampled: Jun 15, 1990

Received: Jun 15, 1990

Analyzed: Jun 19, 1990 Reported: Jun 19, 1990

LABORATORY ANALYSIS FOR:

•

Arsenic

Sample Number	Sample Description	Detection Limit mg/kg	Sample Result mg/kg
0062481 A-B	WPC-1	0.0050	N.D.
0062482 A-B	WPC-2	0.0050	N.D.
0062483 A-B	WPC-3	0.0050	N.D.
0062484 A-B	WPC-4	0.0050	N.D.
0062485 A-B	WPC-5	0.0050	N.D.
0062486 A-B	WPC-6	0.0050	N.D.
0062487 A-B	WPC-B	0.0050	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawvet **Project Manager**

62476.KAL <3>



680 Chesapeake Drive • Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Kaldveer Associates, Inc. 425 Roland Way

Oakland, CA 94621 Attention: Dennis Laduzinsky Client Project ID: #KE1 Sample Descript: Wipe

Analysis for: Lead

First Sample #: 006-2481 A - B

#KE1179-1, Clement - Alameda

Lead

a

Sampled: Jun 15, 1990 Received: Jun 15, 1990

Analyzed: Reported:

Jun 19, 1990 Jun 19, 1990

LABORATORY ANALYSIS FOR:

Sample Number	Sample Description	Detection Limit mg/wipe	Sample Result mg/wipe
0062481 A-B	WPC-1	0.0050	N.D.
0062482 A-B	WPC-2	0.0050	N.D.
0062483 A-B	WPC-3	0.0050	N.D.
0062484 A-B	WPC-4	0.0050	N.D.
0062485 A-B	WPC-5	0.0050	N.D.
0062486 A-B	WPC-6	0.0050	N.D.
0062487 A-B	WPC-B	0.0050	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

SEQUOIA ANALYTICAL

Diane Elich Lawver Project Manager

62476.KAL <4>



680 Chesapeake Drive . Redwood City, CA 94063 (415) 364-9600 • FAX (415) 364-9233

Oakland, CA 94621 Attention: Dennis Laduzinsky QC Sample Group: 0062476 - 0062487 Reported: Jun 19, 1990 **QUALITY CONTROL DATA REPORT**

ANALYTE	Copper	Arsenic	Lead	
Method: Analyst: Reporting Units: Date Analyzed: QC Sample #:	EPA 6010 B. Oliver mg/kg Jun 18, 1990 006-0966	EPA 7050 S. Foster rng/wipe Jun 19, 1990 006-2614	EPA 7421 R. Sharma mg/wipe Jun 19, 1990 008-2481	
Sample Conc.:	20	5.6	N.D.	
Spike Conc. Added:	500	50	0.050	
Conc. Matrix Spike:	500	54	0.052	
Matrix Spike % Recovery:	96	97	100	
Conc. Matrix Spike Dup.:	390	52	0.051	
Matrix Spike Duplicate % Recovery:	74	93	100	
Relative % Difference:	25	3.8	2.0	

SEQUOIA ANALYTICAL

Diane Elich Lawyer Project Manager

% Recovery: Conc. of M.S. - Conc. of Sample x 100 Spike Conc. Added

Conc. of M.S. - Conc. of M.S.D. x 100 Relative % Difference:

(Conc. of M.S. + Conc. of M.S.D.) / 2

Lab Job # _____

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Cx-2		*	1		<u> </u>						-	 	_		<u> </u>	Copper	
WPC-I		6/15	+	with	wip E Somple						+		×	×	X		
WPC-Z		1			from								×	×	X		
wPC-3					10 CM & 10 CM								×	×	×		
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WP C-5				Ш	12 wipes								×	×			
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APPENDIX II

REPORT ON AIR AND WIPE SAMPLING
1829 CLEMENT AVENUE
BUILDING EVALUATION
BY
ENVIRONMENTAL HEALTH CONSULTANTS

REPORT ON AIR AND WIPE SAMPLING 1829 CLEMENT AVENUE BUILDING EVALUATION

Prepared for:

Loren Smith

Prepared by:

Environmental Health Consultants, Inc.

TABLE OF CONTENTS

1.0	Introduction	3
2.0	Sampling Strategy	4
3.0	Selection of Analytes	5
4.0	4.1 Sampling and Analytical Methods 4.2 Recordkeeping	
5.0	Analytical Results	11
6.0	Recommendations	12

FIGURE

APPENDICES

1.0 Introduction

This report addresses the air monitoring and wipe sampling conducted at the building at 1829 Clement Avenue in Alameda, California on April 26, 1990. The monitoring program was conducted in order to characterize the potential exposure of future occupants the building to chemical constituents known to be used in various processes by former tenants of the building. These chemicals were known to exist in the materials beneath the building. However, it was not known to what extent they were present inside the building, and what exposure pathways may exist for future tenants. Through the sampling it was found that the site constituents are present inside the building at very low levels, if at all, and then only on uncovered or unpainted surfaces. No airborne constituents were found at any amount above detectable levels.

2.0 Sampling Strategy

In order to characterize the average concentrations present in the building, a strategy was chosen in which the existing site chemical data and site observations were used to determine the most likely potential constituents for exposure. Appropriate sampling/analysis methods were then chosen, samples collected and analyzed, and results matched with their respective building locations.

3.0 Selection of Analytes

Analytes for the samples were chosen according to: their prevalence on site; the likelihood of exposure of building occupants; and chemical volatility, toxicity, and carcinogenicity.

Metals (specifically arsenic, beryllium, chromium, copper, molybdenum, and lead), and cyanides were chosen as parameters because site data showed that both were used by the former tenants of the building during the plating and etching operations, and that both are present in the materials beneath the building.

4.0 Sampling Protocol

4.1 Sampling and Analytical Methods .

The sampling was conducted in accordance with standardized NIOSH and OSHA methods for sampling and analysis, or a suitable equivalent. Specifically, the methods used for the sampling were: NIOSH Method 7300 for airborne metals; NIOSH Method 7904 for airborne cyanides; and a modified OSHA method for wipe sampling using the appropriate wetting solution. These methods specify QA/QC provisions for instrument calibration. sample collection parameters, packaging, storage, preparation analysis, and analytical procedures. Both air and wipe metals samples were analyzed via inductively-coupled argon spectrophotometry. All cyanide samples were analyzed via ionspecific electrode. A copy of the NIOSH methods cited above are included in Appendix A.

4.2 Recordkeeping

Detailed sample documentation is an integral part of the overall sampling program. This documentation is necessary in order to demonstrate the accuracy and validity of sample data. All sampling data was recorded in ink in the sampling log. Any problems encountered during the sample period were recorded in the log. The following information was also recorded for the samples;

- Date of sampling;
- Sample location;
- 3. Sample identification number;
- 4. Start and stop times, total minutes sampled, and sample volume (air samples);
- 5. Pump number (air samples);
- 6. Pump calibration data and sample flow rate (air samples);

In addition to preserving accurate sampling data, shipping and handling of the samples were documented through the use of a chain-of-custody form which accompanied the samples to the laboratory.

4.3 Sample Media

Several types of media were utilized for the sampling. These media included mixed cellulose ester fiber (MCEF) filters for airborne metals; MCEF filters in conjunction with sodium hydroxide (NaOH) solution in an impinger for airborne cyanides; and Watman ashless filter paper for the wipe sampling. The Watman filters were prewetted by the laboratory with NaOH for the cyanides analysis, and were wetted in the field with purified distilled water for the metals analysis.

4.4 Air Sample Collection

4.4.1 Pump Calibration

All sample pumps were calibrated before and after use to verify the accuracy of the flowrate at which the samples were collected. Pumps were calibrated utilizing a precision rotameter, which was in turn calibrated to a Gilibrator primary standard. The lowest of the pre- and post-calibration flowrates was used in the volume determination. The use of the lower air volume results in a higher calculated airborne concentration, and is therefore more conservative.

4.4.2 Sample Placement

The air samples were collected by placing the sample at a location near the middle of each room, attached to a stanchion which allowed the sample to be collected at a height of approximately three feet above the floor. These locations were chosen in order to obtain the average as far as airborne chemical constituents in the breathing zone throughout the building. Air sample locations are shown on Figure 1. Samples were collected for cyanides at all locations. Metals samples were taken at locations 1, 3, 5, 6, 8, and 9.

4.4.3 Sample Collection

For both the metals and the cyanides air samples, the sampling pump was operated in the high-flow configuration. The pump senses changes in resistance across the filter and adjusts its speed correspondingly, thus assuring a consistent flow rate. The metals samples were collected at a flow rate of approximately two liters/minute (1/min.), while the cyanides samples were run at one 1/min. The plastic plugs were removed from the filter cassette and saved for resealing it after the sampling event. For the cyanides samples, the bubbler impinger was put in-line between the filter and the pump. The sampling trains were attached to the pump by a length of tubing such that the air sample was drawn in through the port on the cassette marked "INLET".

4.4.4 Sampling Control

The sample start and stop times were recorded in order to calculate the total sample volume. The pumps were observed periodically throughout the sampling period to verify the consistency of the flowrate, and any problems which arose were noted. No tampering with the samples was tolerated. All samples ran for a period of 480 minutes, except Sample A-4.26C-3. On this sample, the pump faulted out after 317 minutes due to battery failure. Another battery was installed and the pump restarted. The total sampling time, and hence the sample volume, was adjusted to reflect the period during which the pump was not operating. At the end of the sampling period, the shut-off time and the pump timer reading were recorded for all the pumps. The total air volume for each ambient air sample was calculated by multiplying the flowrate by the total

number of minutes sampled.

4.4.5 Sample Handling

The MCEF filter samples for metals were left in the filter cassette for shipment to the laboratory. The cassette plugs were replaced, and the cassettes were placed in a Ziplock bag and stored in a safe location until shipment. The filters for the cyanides samples were removed from the cassettes with tweezers, transferred to a vial supplied by the laboratory, and labelled with the label from the cassette. The impinger solution was transferred to a separate vial, and the impinger was rinsed into the vial with a small amount of fresh NaOH solution. The label was then placed on the vial.

4.5 Wipe Sample Collection

4.5.1 Sample Locations

Sample locations were chosen in order to obtain an average characterization of concentrations of chemicals on exposed surfaces within the building. An effort was made to obtain a worst-case excenario by sampling only those areas which had not recently been painted or covered with new materials. As such, the areas sampled were the sections of original flooring throughout the building, with nearly every room that was used in the former operations being sampled. Each cyanides and metals samples were taken at every exception.

4.5.2 Sample Collection

At each location, two adjacent 10-centimeter (cm) by 10-cm squares were marked on the surface to be sampled, one square for the cyanides sample and one for the metals. A clean pair of latex gloves was used for each sample location in order to prevent cross-contamination of samples.

4.5.3 Sample Handling

After each wipe sample was collected, it was placed in a separate pre-labelled vial and the lid was securely fastened. At the end of the sampling, the vials were placed in the cooler for shipment to the laboratory.

4.6 Field Blanks

One field blank was prepared for each type of sample collected. One of each of the sample media was handled as though it were being used to sample. The plugs from the filter cassettes for the metals samples were removed and then replaced. The MCEF filter for a cyanide air sample was removed from the cassette and placed in a vial. An amount of NaOH solution was poured directly into another vial as a blank for the cyanides impinger samples. The Watman filter for the wipe samples for cyanides was briefly taken from its vial and placed back in the same vial. A wipe filter for metals was moistened with the same distilled water used for the other metals samples and placed in a vial. The blank samples were labelled, stored, and shipped in the same manner as the other samples.

4.7 Chain of Custody and Shipment

Each sample number was listed on the chain-of-custody form with the analysis to be performed, along with other pertinent information.

The samples were shipped via overnight delivery to the laboratory. The samples and chain-of-custody were placed in a cooler with blue ice and packing material (the metals samples do not require blue ice.) The cooler was then taped shut to assure evidence of any tampering, and to prevent accidental loss of the samples from the cooler. All samples were sent to Clayton Environmental Consultants, Inc., for analysis. Clayton is a laboratory

accredited by the American Industrial Hygiene Association (AIHA). Chain-of-custody forms for the samples are included in Appendix B.

5.0 Analytical Results

All sample data was reviewed for the sampling procedures followed, recordkeeping, and validity and accuracy of the data. The actual analytical results reported by Clayton are included in Appendix C. The number at the end of each sample identification number corresponds to the location numbers delineated on Figure 1.

Volumes for the air samples ranged from 434 to 480 liters for the cyanides samples, and all metals samples volumes were 960 liters. Detection limits for metals ranged from 1 to 5 micrograms of metal per cubic meter of air (ug/m^3) , and for cyanides from 13 to 14 ug/m^3 . All of these detection levels are well below the current OSHA Permissible Exposure Limits (PEL)s.

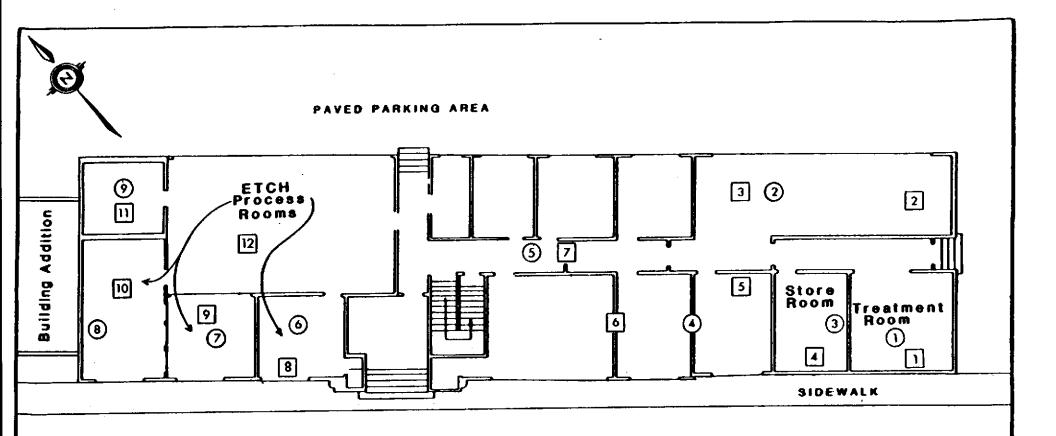
The detection limits for the wipe samples for metals ranged from 2 to 7 ug, while the evenides detection level was 3 ug.

were below the detection limits, as were all of the cyanides wipe samples. Two of the wipe samples for metals contained amounts just at or above the detection level. The wipe sample from the "Treatment Room", Location 1, contained copper at 7 ug, right at the detection level. The sample from the corner storage room, Location 11, contained lead at 4 ug, just above the detection level of 3 ug.

6.0 Recommendations

As of the writing of this document, the intended use for the building at 1829 Clement Avenue is for business office space. The walls have been painted, some of the floors have been replaced, and other floors have been or are going to be covered over with plywood and carpet. All of these measures will tend to reduce or eliminate any potential exposures which may occur to the occupants through a contact route of exposure. Because of the non-detectable to low concentrations found on the building surfaces, and because the samples were taken as a worst-case example, such measures should be sufficient.

Air quality is not of concern, since the samples were all below the limits of detection, which are all well below OSHA permissible levels.



----- Clement Avenue

APPROXIMATE SCALE IN FEET

LEGEND

■ Wipe

O Air



|--|

1829 CLEMENT AVENUE Alameda, California

PROJECT NO	DATE	Figure 1
KE1179-1	March 1990	1.00.0

APPENDIX A

NIOSH ANALYTICAL METHODS

CYANIDES, aerosol and gas

FORMULA: HCN and salts

METHOD: 7904

H.W.: 27.03 (HCN); 65.11 (KCN)

ISSUED: 2/15/84

OSHA: 11 mg/m³; skin (HCN)

PROPERTIES: HCN: gas, 8P 26 °C

5 mg/m³; skin (cyanides, as CNT)

KCN: solid, d 1.52 g/mL, MP 634 °C

NIOSH: $5 \text{ mg/m}^3/10 \text{ min (as CN}^-)$ [1] ACGIH: C 10 mg/m3; skin (HCN);

5 mg/m³; skin (cyanides, as CN)

SYNONYMS: HCM: hydrocyanic acid, prussic acid, formonitrile, CAS #74-90-8.

cyanides: CAS #151-50-8; CAS #143-33-9.

MEASUREMENT

SAMPLER: FILTER + BUBBLER

(0.8-um cellulose ester membrane

SAMPL ING

+ 10 mL 0.1 N KOH)

!TECHNIQUE: ION-SPECIFIC ELECTRODE

!ANALYTE: cyanide ion (CNT)

FLOW RATE: 0.5 to 1 L/min

!EXTRACT FILTER: 25 mL 0.1 N KOH; 30 min

VOL-MIN: 10 L @ 5 mg/m³ (as CN-)-MAX: 180 L @ 11 mg/m3 (as CNT)

!RINSE BUBBLER: 2 mL 0.1 N KOH; dilute to 25 mL

with 0.1 N KOH

SHIPMENT: routine

!MEASURE: mV reading of cyanide ion electrode vs.

reference electrode

SAMPLE STABILITY: HCN stable in 0.1 N KOH

at least 1 week [1];

!CALIBRATION: solutions of KCN in 0.1 N KOH

particulate on filter may liberate HCN gas [2]

!RANGE: 0.05 to 2 mg CNT

BLANKS: 2 to 10 field blanks per set

!ESTIMATED LOD: 2.5 µg CNT [2]

ACCURACY

!PRECISION (sr): 0.043 (HCN) [3];

0.038 (KCN) [2]

RANGE STUDIED: 5 to 21 mg/m3 (HCN) [3];

2.6 to 10 mg/m3 (KCN) [2]

BIAS: not significant [2,3]

OVERALL PRECISION (sr): 0.081 (HCN) [3];

0.103 (KCN) [2]

APPLICABILITY: The working range (as CN-) is 0.5 to 15 mg/m 3 for a 90-L air sample or

5 to 20 mg/m³ for a 10-L air sample. INTERFERENCES: Sulfide, chloride, iodide, bromide, cadmium, zinc, silver, nickel, cuprous iron and mercury interfere. In humid atmospheres, some particulate cyanide collected on the filter will liberate hydrogen cyanide which will be trapped in the bubbler [2]. The method cannot distinguish between HCN formed in this manner and HCN originally present in air.

OTHER METHODS: This method combines and replaces Methods S288 [4], S250 [5], and P&CAM 116 [6].

REAGENTS:

- 1. Double distilled (d.d.) water.
- 2. Potassium cyanide.*
- Calibration stock solution, 1000 µg CNT/mL. Dissolve 0.250 g KCN in 0.1 M KOH to make 100 mL solution. Stable for at least 1 week in polyethylene bottle.
- Potassium hydroxide (KOH), 0.1 N. Dissolve 5.6 g KOH in d.d water; dilute to 1000 mL.
- 5. lead acetate paper.
- 6. Cadmium carbonate (if sulfide present).
- 7. Hydrogen peroxide, 30% (if sulfide present).
- Sodium sulfite, 1 M (if sulfide present).

*See Special Precautions.

EQUIPMENT:

- Sampler: mixed cellulose ester membrane filter, 37-mm diameter, 0.8-um pore size, followed by a glass midget bubbler containing 15 mL 0.1 <u>H</u> KOH.
- Personal sampling pump, 0.5 to 1 L/min, with splashover protection and flexible connecting tubing.
- Vials, polyethylene, with screw caps, 20-mL, and plastic tape for sealing.
- 4. Cyanide ion electrode, (Orion 94-06 or equivalent).
- 5. Reference electrode.
- 6. pH meter, readable to 0.1 mV.
- 7. Magnetic stirrer and stirring bars.
- 8. Jars, ointment, 60-mL, squat-form with aluminum-lined screw caps.
- 9. Pipets, 0.05- to 2-and 25-mL, with pipet bulb.
- 10. Volumetric flasks, 25-mL.
- 11. Beakers, 50-mL.
- 12. Analytical balance, readable to 0.1 mg.

SPECIAL PRECAUTIONS: Hydrogen cyanide gas and the cyanide particulates may be fatal if swallowed, inhaled or absorbed through the skin. Work in a hood.

Amyl nitrite is the antidote for cyanide poisoning [1].

SAMPLING:

- 1. Calibrate each personal sampling pump with a representative sampler in line.
- Sample at 0.5 to 1 L/min for a total sample size of 10 to 180 L.
 NOTE: Maintain bubblers in a vertical position during sampling. Do not allow the solution level to fall below 10 mL.
- 3. Remove the bubbler stem and tap it gently against the inside wall of the bubbler. Rinse the bubbler stem with 1 to 2 mL of unused 0.1 \underline{N} KOH. Add the rinse to the bubbler.
- 4. Quantitatively transfer the contents of the bubbler to a 20-mL vial. Close cap tightly and wrap with plastic tape to avoid sample loss during transit. Label each vial.

SAMPLE PREPARATION:

- 5. Transfer the filter from the cassette filter holder to a 60-mL ointment jar.
- 6. Pipet 25.0 mL 0.1 N KOH into the jar. Cap and allow to stand for at least 30 min with occasional shaking to complete extraction. Analyze within 2 hrs after extraction.
- 7. Empty the contents of the vial into a 25-mL volumetric flask using 0.1 N KOH to rinse the vial. Add rinse to the volumetric flask. Dilute to the mark with 0.1 N KOH.
 - NOTE: Sulfide ion irreversibly poisons the cyanide ion specific electrode and must be removed if present. Check for the presence of sulfide ion by touching a drop of sample to a piece of lead acetate paper; the paper will discolor in the presence of sulfide ion. If this test is positive, remove sulfide by one of the following
 - a. Add 1 mL 1 $\underline{\text{M}}$ H₂O₂ and 1 mL 1 $\underline{\text{M}}$ Na₂SO₃ to sample solutions prior to diluting to volume; or

b. Add a small amount (spatula tip) of powdered cadmium carbonate to the sample. Swirl to disperse the solid and recheck the liquid with lead acetate paper. If sulfide ion has not been removed completely, add more cadmium carbonate. Avoid a large excess of cadmium carbonate and long contact time with the solution. When a drop of liquid no longer discolors a strip of lead acetate paper, filter the sample through a small plug of glass wool in a Pasteur pipette and proceed with the analysis.

CALIBRATION AND QUALITY CONTROL:

- 8. Prepare at least five working standards fresh daily to cover the range 50 to 2000 μg CNT per sample by diluting aliquots of 1000 μ g/mL calibration stock solution with 0.1 \underline{N} KOH (e.g., 0.05 to 2.0 mL calibration stock solution diluted to 25 mL).
- 9. Analyze the working standards according to steps 11 and 12 together with the samples and blanks.
- 10. Prepare a calibration graph on semilog paper by plotting cyanide ion concentration on the logarithmic axis and mW on the linear axis.

MEASUREMENT:

- 11. Transfer the solution to be measured to a 50-mL beaker. Immerse the cyanide ion electrode and reference electrode in the sample and start the magnetic stirrer.
- 12. With the magnetic stirrer on, allow the potential reading to stabilize. Record the mV reading.
 - NOTE 1: Potential readings are a function of temperature. Measure samples and working standards at the same temperature (\pm 2 °C).
 - NOTE 2: The cyanide electrode will malfunction if chloride, lodide and bromide ions, which form insoluble silver salts, are present in sufficient quantity. Several metal ions are also known to complex with cyanide such as cadmium, zinc, silver, nickel, cuprous iron and mercury. Consult the electrode instruction manual for the procedure to use when such ions are present.

CALCULATIONS:

- 13. Read the mass, μg , of cyanide ion present in the sample filter ($W_{\vec{r}}$), sample bubbler $(W_{\rm b})$, average media blank filter $(B_{\rm f})$ and media blank bubblers $(B_{\rm b})$ from the calibration graph.
- 14. Calculate the concentration (mg/m^2) of particulate cyanide, Cp, and hydrogen cyanide, C_{HCN}, in the air volume sampled, V (L):

$$Cp = \frac{W_f - B_f}{V}$$
, mg/m³ and $C_{HCN} = \frac{(W_0 - B_0)_{+1.04}}{V}$, mg/m³

where 1.04 is the stoichiometric conversion factor from CNT to HCN.

NOTE: Particulate cyanides will be collected on the filter. In humid atmospheres, however, it has been observed that during the collection of particulate cyanide, HCN is gradually liberated [2]; therefore, particulate cyanide interference is not completely removed.

EVALUATION OF METHOD:

HCN: Method \$288 was issued on September 2, 1977 [4]. Test atmospheres of HCN were generated by calibrated flow from a compressed mixture of HCN in nitrogen [3,7]. The range of HCN concentrations in air was 5 to 21 mg/m³ for 12-L air samples. Eighteen HCN samples collected at 0.2 L/min for 60 min indicated overall precision of 6.2%, with a 96.7% recovery. An eight-day storage stability study involving six samples at the OSHA standard concentration

level indicated a 92.4% average recovery for the one-day old samples and a 92.6% for eight-day old samples. A collection efficiency study at twice the OSHA standard level, which included backup bubblers; indicated that an average of 99.8% of HCN was collected in the first bubbler. The HCN air generated concentrations were independently confirmed by a titration method [3].

<u>KCN</u>: Method S250 was issued on January 30, 1976 [5]. A set of six weighed KCN samples in the range of 1.8 to 2.5 mg KCN per filter indicated a 97% recovery and a 3.8% measurement precision [2]. Spiking with aqueous or basic solutions of KCN proved unsuccessful (low recovery) because of the cyanide instability in the presence of water and CO_2 . Test atmospheres of KCN were generated by atomization of an aqueous solution (162 g/L) of KCN into a dry airstream. Eighteen KCN samples collected in 0.1 N NaOH at 1.5 L/min for 60 min indicated overall precision, s_p , of 0.09. Collection was accomplished with cellulose ester membrane filters followed with backup bubblers. The collection efficiency at twice the OSHA level was 100.0% on the filters. Cyanide salts are known to decompose in moist air with liberation of HCN. This instabilty was determined with two sets of six samples at the one and two times the OSHA level. Each of the samples which were twice the OSHA level were connected with two backup bubblers. Both sets indicated a loss of 16.5%.

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- [2] Documentation of the NIOSH Validation Tests, S250, U.S. Department of Health, Education, and Welfare, Publ. (NIOSH) 77-185 (1977).
- [3] Backup Data Report for Hydrogen Cyanide, S288, available as "Ten NIOSH Analytical Methods, Set 5," Order No. 8P 287-499 from NTIS, Springfield, VA 22161.
- [4] NIOSH Manual of Analytical Methods, 2nd. ed., V. 4, \$288, U.S. Department of Health, Education, and Welfare, Publ. (NIOSH) 78-175 (1978).
- [5] Ibid, V. 3, \$250, U.S. Department of Health, Education, and Helfare, Publ. (NIOSH) 77-157-C (1977).
- [6] Ibid, V. 1, P&CAM 116, U.S. Department of Health, Education, and Welfare, Publ. (NIOSH) 77-157-A (1977).
- [7] NIOSH Research Report-Development and Validation of Methods for Sampling and Analysis of Workplace Toxic Substances, U.S. Department of Health and Human Services, Publ. (NIOSH) 80-133 (1980).

METHOD REVISED BY: J. Palassis, NIOSH/DPSE; S250 and S288 originally validated under NIOSH Contracts COC-99-74-45 and 210-76-0123, respectively.

ELEMENTS (ICP)

METHOD: 7300

M.W.: Table 1

ISSUED: 2/15/84

OSHA/NIOSH/ACGIH: Table 1

PROPERTIES: Table 1

ELEMENTS: aluminum

cobalt arsenic copper beryllium iron cadmium lead lithium calcium

manganese mp) ybdenum nickel phosphorus platinum

selenium

silver sod i um tellurium thallium.

tunosten vanadium yttrium zinc zirconium

chromium

SYNONYMS: vary depending upon the compound.

magnesium

SAMPLING

HEASUREMENT

titanium

SAMPLER: FILTER

(0.8-um, cellulose ester membrane)

FLOW RATE: 1 to 4 L/min

VOL-MIN: Table 1 -MAX: Table 1

SHIPMENT: routine

SAMPLE STABILITY: stable

BLANKS: 2 to 10 field blanks per set

ACCURACY

!TECHNIQUE: INDUCTIVELY COUPLED ARGON PLASMA, ATOMIC EMISSION SPECTROSCOPY

tin

!ANALYTE: elements above

!ASHING REAGENTS: conc. HNO3, 4 mL;

and conc. HC104, 1 mL

CONDITIONS: room temperature, 30 min;

150 °C to near dryness

FINAL SOLUTION: 4% HNO3, 1% HC104, 10 mL

:WAVELENGTH: depends upon element; Table 2

!BACKGROUND CORRECTION: spectral wavelength shift

!CALIBRATION: elements in 45 HNO3, 15 HC104

!RANGE: 2.5 to 1000 ug per sample [1]

!ESTIMATED LOO: 1 µg per sample [1]

!PRECISION (s_r): Table 2

RANGE STUDIED: not studied

BIAS: none identified

OVERALL PRECISION (sp): not evaluated

APPLICABILITY: The working range of this method is 0.005 to 2.0 mg/m3 for each element in a 500-L air sample. This is simultaneous elemental analysis, not compound specific. Verify that the types of compounds in the samples are soluble with this ashing procedure.

INTERFERENCES: Spectral interferences are the primary interferences encountered in ICP-AES analysis. These are minimized by judicious wavelength selection, interelement correction factors and background correction [1,2].

OTHER METHODS: This method replaces PSCAM 351 [2] for trace elements. Atomic absorption spectroscopy (e.g., Methods 70xx) is an alternate analytical technique for many of these <u>elements.</u>

2/15/84

1300-1

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REAGENTS:

- 1. Nitric acid, conc.
- 2. Perchloric acid. conc.*
- Ashing acid: 4:1 (v/v) HNO₃:HC10₄.
 Mix 4 volumes conc. HNO₃ with 1 volume conc. HC10₄.
- Calibration stock solutions, 1000 µg/mL. Commercially available, or prepared per instrument manufacturer's recommendation (see step 12).
- Dilution acid, 4% HNO₃, 1% HClO₄.
 Add 50 mL ashing acid to 600 mL water; dilute to 1 L.
- 6. Argon.
- 7. Distilled, deionized water.

*See Special Precautions.

EQUIPMENT:

- Sampler: cellulose ester membrane filter,
 O.8-mm pore size, 37-mm diameter; in cassette filter holder.
- Personal sampling pump, 1 to 4 t/min, with flexible connecting tubing.
- Inductively coupled plasma-atomic emission spectrometer, equipped as specified by the manufacturer for analysis of elements of interest.
- 4. Regulator, two-stage, for argon.
- 5. Beakers, Phillips, 125-mL, or Griffin, 50-mL, with watchglass covers.*
- 6. Volumetric flasks, 10- and 100- mL.*
- 7. Assorted volumetric pipets as needed.*
- 8. Hotplate, surface temperature 150 °C.

*Clean all glassware with conc. nitric acid and rinse thoroughly in distilled water before use.

SPECIAL PRECAUTIONS: Perform all perchloric acid digestions in a perchloric acid hood.

SAMPLING:

- 1. Calibrate each personal sampling pump with a representative sampler in line.
- Sample at an accurately known flow rate between 1 and 4 L/min for a total sample size of 200 to 2000 L (see Table 1) for TWA measurements. Do not exceed a filter loading of approximately 2 mg total dust.

SAMPLE PREPARATION:

- 3. Open the cassette filter holders and transfer the samples and blanks to clean beakers.
- Add 5 mL ashing acid. Cover with a watchglass. Let stand 30 min at room temperature.
 MOTE: Start a reagent blank at this step.
- 5. Heat on hotplate (120 °C) until ca. 0.5 mL remains.
 - NOTE: Some species of Li, Mn, Mo, Sn, W, and Zr will not be completely solubilized by this procedure. Alternative solubilization techniques for most of these elements can be found elsewhere [2,3,4,5,6,7].
- 6. Add 2 mL ashing acid and repeat step 5. Repeat this step until the solution is clear.
- Remove watchglass and rinse into the beaker with distilled water.
- 8. Increase the temperature to 150 °C and take the sample to dryness.
- 9. Dissolve the residue in 2 to 3 mL dilution acid.
- Transfer the solutions quantitatively to 10-mL volumetric flasks.
- 11. Dilute to volume with dilution acid.

CALIBRATION AND QUALITY CONTROL:

- 12. Calibrate the spectrometer according to the manufacturers recommendations.
 - MOTE: Typically, an acid blank and 10 $\mu g/mL$ multielement working standards are used. The following multielement combinations are chemically compatible in 4% HNO₃/1% HClO₄:
 - a. Ag, Ca, Co, Mn, Pb, V, Zn;
 - b. Al, Be, Cd, La, Li, Ni, Tl;
 - : c. As, B, Ba, Mg, Mo, P, Sn;

d. Cu, Fe, Na, Pt, Sr, Te, Y;

e. Cr, K, Sb, Se, Ti, Zr; and

f. Si, W (distilled water only)

13. Analyze a standard for every ten samples.

14. Check recoveries with at least two spiked media blanks per ten samples.

MEASUREMENT:

15. Set spectrometer to conditions specified by manufacturer.

16. Analyze standards and samples.

NOTE: If the values for the samples are above the range of the standards, dilute the solutions with dilution acid, reanalyze and apply the appropriate dilution factor in the calculations.

CALCULATIONS:

17. Obtain the solution concentrations for the sample, C_g (µg/mL), and the average media blank, C_b (µg/mL), from the instrument.

18. Using the solution volumes of sample, V_S (mL), and media blank, V_D (mL), calculate the concentration, C (mg/m³), of each element in the air volume sampled, V (L):

EVALUATION OF METHOD:

Method P&CAM 35) was evaluated in 1981 [1,2]. The precision and recovery data were determined at 2.5 and 1000 µg of each element per sample on spiked filters. The precision and recovery data, instumental detection limits, sensitivity, and analytical wavelengths are listed in Table 2. The values in Table 2 were determined with a Jarrell-Ash Model 1160 ICP operated according to manufacturer's instructions.

REFERENCES:

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- [3] Ibid, S341 (Lead).
- [4] Ibid, V. 2, S5 (Manganese), U.S. Department of Health, Education, and Welfare, Publ. (NIOSH) 77-157-8 (1977).
- [5] Ibid, V. 4, P&CAM 271 (Tungsten), U.S. Department of Health, Education, and Welfare, Publ. (NIOSH) 78-175 (1978).
- [6] Ibid, V. 5, P&CVM 173 (Metals by Atomic Absorption), U.S. Department of Hamilton, Education, and Welfare, Publ. (NIOSH) 79-141 (1979).
- [7] Ibid, V. 3, \$183 (Tin), \$185 (Zirconium), and \$376 (Molybdenum), U.S. Department of Health, Education, and Welfare, Publ. (NIOSH) 77-157-C (1977).

METHOD REVISED BY: R. DeLon Hull and Mark Millson, NIOSH/DPSE.

Table 1. Properties and sampling volumes.

Arsenic (As) 74.92 817# 0.5/C 0.002/ 0.2 5 2000 Beryllium (Be) 9.01 1278 0.002/ 0.0005/ 0.002 1250 2000 Calcium (Ca) 40.08 842 5 (b)/ — / 2 (b) 5 200 Cadmium (Cd) 112.40 321 0.2/ 0.04/ 0.05 13 2000 Cobalt (Co) 58.93 1495 0.1/ — / 0.1 25 2000 Chromium (Cr) 52.00 1890 1.0 (c)/ 0.025/ 0.5 (c) 5 1000 Copper (Cu) 63.54 1083 1.0/ — / 1.0 5 1000 Iron (Fe) 55.85 1535 10 (b)/ — / 5 (b) 5 1000 Lithium (Li) 6.94 179 0.025 (d)/ — / 0.025 (d) 100 2000 Magnesium (Mg) 24.31 651 15 (b)/ — / 10 (b) 5 6 Manganese (Mn) 54.94 1244 C 5/ — / C 5 5 200 Mclybdenum (Mg) 95 94 251 15 (e)/ — / 10 (e) 5 6 Sodium (Na) 22.99 98 2 (c)/ 2 2 (f)/ C 2 (f) 13 2000 Nickel (Ni) 58.71 1453 1/ 0.015/ 1 (c) 5 1000	
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Lead (Pb) 207.19 328 0.05/ 0.1/ 0.15 50 200	
	(g)
Platinum (Pt) 195.09 1769 0.002 (a)/ / 1 (c) 1250 200	
Selenium (Se) 78.96 217 0.2/ — / — 13 200	
Tin (Sn) 118.69 232 2/ / 2 (c) 5 50	
Tellurium (Te) 127.60 450 0.1/ — / 0.1 25 200	
Titanium (Ti) 47.90 1675 $-/-/10$ (b) 5 10	
Thallium (T1) 204.37 304 0.1 (a) $-$ / 0.1 (a) 25 200	
Vanadium (V) 50.94 1890 C 0.5/ 1 (c)/ 0.05 (V ₂ O ₅) 5 200	
Tungsten (W) 183.85 3410 — / 5 (e) / 5 (e) 5 (g) 20	(g)
Yttrium (Y) 88.91 1495 1/ / 1 5 100	-
Zinc (Zn) 65.37 419 5 (b) / 5 (b) / 5 (b) 5 20	
Zirconium (Zr) 91.22 1852 5/ / 5 5 20	

⁽a) soluble

⁽b) oxide

⁽c) metal

⁽d) hydride

⁽e) insoluble (f) hydroxide

⁽g) at the ACGIH TLV

Table 2. Heasurement procedures and data (a).

		Instrumental	Sensitivity	Recove	ry (\$)	Prec (N	ision (s _r) = 3)	
Element	Wavelength (nm)	LOD (ng/mL)	(Intensity/ µg/mL)	# 2.5 µg/ filter (b)	# 1000 ug/ filter	6 2.5 µg/ filter	€ 1000 µg/ filter	
Ag	328.3	26	0.65	113	91	0.02	0.075	
ΑĪ	308.2	14	0.23	93	100	0.092	0.023	
As	193. <i>7</i>	13	0.57	103	99	0.062	0.026	
Be	313.0	1.5	1.29	107	90	0.040	0.034	
Ca	315.9	10	0.49	99	95	0.036	0.014	
Cd	226.5	1.6	0.83	107	99	0.032	0.020	
Co	231.2	7.4	0.38	101	95	0.040	0.005	
Cr	205.6	1.3	0.50	98	106	0.053	0.016	
Cu	324.8	2.1	0.72	98	99	0.036	0.022	
Fe	259.9	3.9	0.13	94	97	0.068	0.016	
Li	670.B	2.8	0.48	89	95	0.171	0.043	
Mg	279.6	24	0.22	105	106	0.084	0.027	
Mn	257.6	0.4	0.74	84	93	0.062	0.035 0.049	
Mo	281.6	7.0	0.18	94	88	0.023		
Na	589.0	10	0.76	(c)	101	(c)	0.045	
Ni	231.6	3.4	0.41	105	97	0.027	0.020	
P	214.9	22	0.17	(c)	91	(c)	0.056	
Pb	220.4	17	0.42	105	95	0.060	0.011	
Pt	203.7	15	0.60	106	91	0.041	0.075	
Şe	190.6	21	0.28	105	9.7	0.068	0.049	
\$n	190.0	64	0.49	74	67	0.33	0.16	
Te	214.3	29	0.41	102	94	0.050	0.063	
Ti	334.9	1.2	0.55	96	108	0.051	0.029	
TI	190.9	17	0.22	103	99	0.043	0.017	
٧	310.2	3.2	0.88	99	94	0.043	0.014	
₩	207.9	13	2.58	35	23	0.053	0.60	
Y	371.0	0.8	2.35	99	100	0.015	0.013	
Zn	213.9	0.6	0.60	101	94	0.013	0.013	
Zr	339.2	1.9	0.88	75	98	0.049	0.008	

⁽a) Values reported were obtained with a Jarrell-Ash Model 1160 ICP; performance may vary with instrument and should be independently verified.

⁽b) 2.5 μ g/filter corresponds to 5 μ g/m³ for a 500-L air sample.

⁽c) Blank levels too high to make accurate determinations

APPENDIX B

CHAIN-OF-CUSTODY FORMS

Clayton ENVIRONMENTAL CONSULTANTS

REQUEST FOR LABORATORY ANALYTICAL SERVICES

For Clayton Use Only Page	a <u>3</u>
Project No.	
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Date Logged In	By

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- 74444 11	war completed form and semples to	ON OF THE C	HITION ENVIRON	ineuro Cou	suπants, Inc. labe	iisled	below:] [JISTR	IBUTI	ON:		

22345 Roethel Drive Novi, MI 48050 (313) 344-1770

Reritan Center 160 Fieldcrest Ave. Edison, NJ 08837 (201) 225-6040 400 Chastain Center Blvd., N.W. Suite 490

Kennesaw, GA 30144

(404) 499-7500

1252 Quarry Lane Pleasanton, CA 94566 (415) 426-2600 WHITE - Clayton Laboratory
YELLOW - Clayton Accounting
PINK - Client Retains

Clayton ENVIRONMENTAL CONSULTANTS

REQUEST FOR LABORATORY ANALYTICAL SERVICES

For Clayton Use Only	Page _	2	_ a_2_	
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Please r	eturn completed form and samples to one of the		mental Con	sultants, Inc. lab	s listed	below:		<u> </u>	DISTRIB	UTION:		

22345 Roethel Drive Novi, MI 48050 (313) 344-1770 Raritan Center 160 Fieldcrest Ave. Edison, NJ 08837 (201) 225-6040 400 Chastain Center Blvd., N.W. Suite 490

Kennesaw, GA 30144

(404) 499-7500

Vd., N.W. 1252 Quarry Lane Pleasanton, CA 94566 (415) 426-2600 WHITE - Clayton Laboratory YELLOW - Clayton Accounting

PINK - Client Retains

Clayton **ENVIRONMENTAL**

REQUEST FOR LABORATORY

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For Clayton Use Only	Page_	of ,	<u> </u>	
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Dept.

Title

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ANALYSIS REQUESTED

(Enter an X' in the box below to indicate request; Enter a 'P' if Preservative added")

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Rush Charges Authorized?

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Please return completed form and samples to one of the Clayton Environmental Consultants, Inc. labs listed below:

22345 Roethel Drive Novi, MI 48050 (313) 344-1770

Authorized by:

(If required) Method of Shipment:

Raritan Center 160 Fieldcrest Ave. Edison, NJ 08837 (201) 225-6040

(Client Signature Must Accompany Request)

Special Instructions: (method, limit of detection, phone results, rush results, etc.)

400 Chastain Center Blvd., N.W.

Suite 490 Kennesaw, GA 30144 (404) 499-7500

1252 Quarry Lane Pleasanton, CA 94566 (415) 426-2600

4-26-90

Sample condition upon receipt:

DISTRIBUTION:

WHITE Clayton Laboratory **Clayton Accounting YELLOW** -**Client Retains** PINK

Clayton ENVIRONMENTAL CONSULTANTS

REQUEST FOR LABORATORY ANALYTICAL SERVICES

For Clayton Use Only Page	<u>/ a 3 </u>
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Please return completed form and samples to one of the Clayton Environmental Consultants, Inc. labs listed below:

22345 Roethel Drive Novi, MI 48050 (313) 344-1770

Raritan Center 160 Fieldcrest Ave. Edison, NJ 08837 (201) 225-6040

400 Chastain Center Blvd., N.W. Suite 490

1252 Quarry Lane Pleasanton, CA 94566 Kennesaw, GA 30144 (415) 426-2600 (404) 499-7500

Clayton Laboratory WHITE YELLOW -**Clayton Accounting Client Retains** PINK



REQUEST FOR LABORATORY ANALYTICAL SERVICES

For Clayton Use Only Page	2.	<u>-3</u>
Project No.		
Batch No.		
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A Marsh & McLennan Company

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Please return completed form and samples to one of the Clayton Environmental Consultants, Inc. labs listed below

22345 Roethel Drive Novi, MI 48050 (313) 344-1770 Reritan Center 160 Fieldcrest Ave. Edison, NJ 08837 (201) 225-6040 400 Chastain Center Blvd., N.W. Suite 490

Kennesaw, GA 30144

(404) 499-7500

1252 Quarry Lane Pleasanton, CA 94566 (415) 426-2600 WHITE - Clayton Laboratory
YELLOW - Clayton Accounting
PINK - Client Retains

Clayton ENVIRONMENTAL CONSULTANTS

REQUEST FOR LABORATORY **ANALYTICAL SERVICES**

For Clayton Use Only Pag	. <u>3</u> . <u>3</u>
Project No.	
Batch No.	
Client No.	
Date Received	Ву
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A Marsh & McLennan Company

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Date Re	sults Required:	Rush Charg	es Authorized	? 🗆 Yes	₩		(Enter	en X' in ti	e box bek			REQUE		• P # P	reservi	stive added")
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	CLIENT SAMPLE	DENTIFICATION	DATE SAMPLED	MATRIX/ MEDIA	AIR VOLUME (specify units)		/C	<u> </u>		\angle						FOR LAB USE ONLY
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22345 Roethel Drive Novi, MI 48050 (313) 344-1770

Raritan Center 160 Fieldcrest Ave. Edison, NJ 08837 (201) 225-6040

400 Chastain Center Blvd., N.W. Suite 490

1252 Quarry Lane Pleasanton, CA 94566 Kennesaw, GA 30144 (415) 426-2600 (404) 499-7500

WHITE **Clayton Laboratory** Clayton Accounting YELLOW -**Client Retains** PINK

APPENDIX C

LABORATORY ANALYTICAL REPORTS

1252 Quarry Lane Pleasanton, CA 94566 (415) 426-2600 Fax (415) 426-0106



May 1, 1990

Ms. Irene Fanelli ENVIRONMENTAL HEALTH CONSULTANTS. INC. P.O. Box 117910 Burlingame, CA 94011-7910

> Client Ref. No. 89-055 Work Order No. 9004182 Lab Client Code 100662

Dear Ms. Fanelli:

Attached is our analytical laboratory report for the samples received on April 27, 1990. A copy of the Chain-of-Custody form acknowledging receipt of these samples is attached.

Please note that any unused portion of the samples will be disposed of 30 days after the date of this report, unless you have requested otherwise.

We appreciate the opportunity to be of assistance to you. If you have any questions, please contact Maryann Gambino, Client Services Representative, at (415) 426-2657.

Sincerely,

Ronald H. Peters, CIH

Manager, Laboratory Services

Western Operations

RHP/dt

Attachments

INDUSTRIAL HYGIENE METALS ANALYSIS

Sample I.D.: See below

Client: ENVIRONMENTAL HEALTH CONS.INC

Sample Received: 04/27/90

Client Ref. No.: 89-055

Samples Analyzed: 04/27/90

Lab Client Code: 100662

Sample Matrix: FILTER

Project No.: 9004182

Lab No. Sample I.D.	Volume (Liters)	Analyte	Amount (mg)	Conc. (mg/m3)	Detection Limit (mg)
-01 A-4.26M-1		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
		Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001
-02 A-4.26M-3		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
		Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001
-03 A-4.26M-5		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
		Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001
-04 A-4.26M-6		Arsenic	<0.001		0.001
		Beryllium	<0.0005	-,	0.0005
•	•	Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001

< = Less than the indicated limit of detection (LOD)</pre>

^{-- =} Information not available or not applicable

INDUSTRIAL HYGIENE METALS ANALYSIS

Sample I.D.: See below

Client: ENVIRONMENTAL HEALTH CONS.INC

Sample Received: 04/27/90

Client Ref. No.: 89-055

Samples Analyzed: 04/27/90

Lab Client Code: 100662

Sample Matrix: FILTER

Project No.: 9004182

Lab No. Sample I.D.	Volume (Liters)	Analyte	Amount (mg)	Conc. (mg/m3)	Detection Limit (mg)
-05 A-4.26M-8		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
		Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001
-06 A-4.26M-9		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
		Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001
-20 BLANK		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
		Chromium	<0.002		0.002
		Copper	<0.002		0.002
-		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001
-21 METHOD BLANK		Arsenic	<0.001		0.001
		Beryllium	<0.0005		0.0005
·		Chromium	<0.002		0.002
		Copper	<0.002		0.002
		Molybdenum	<0.001		0.001
		Lead	<0.001		0.001

< = Less than the indicated limit of detection (LOD)
-- - Information not available or not applicable</pre>

METHOD REFERENCE: NIOSH 7300

^{-- =} Information not available or not applicable

Page 4 of 7

INORGANIC LABORATORY ANALYSES

Sample I.D.:

See below

Client: ENVIRONMENTAL HEALTH

IND HUNDIN

Sample Received:

04/27/90

Client Ref. No.:

89-055

Sample Analyzed:

05/01/90

Lab Client Code:

100662

Sample Matrix:

Wipe

Lab No.:

9004182

Batch Sub. No.	Sample Identification	Arsenic (mg/wipe)	Beryllium (mg/wipe)	Chromium (mg/wipe)
-07A	W-4.26M-1	<0.003	<0.002	<0.007
-08A	W-4.26M-2	<0.003	<0.002	<0.007
-09A	W-4.26M-3	<0.003	<0.002	<0.007
-10A	W-4.26M-4	<0.003	<0.002	<0.007
-11A	W-4.26M-5	<0.003	<0.002	<0.007
-12A	W-4.26M-6	<0.003	<0.002	<0.007
-13A	W-4.26M-7	<0.003	<0.002	<0.007
-14A	W-4.26M-8	<0.003	<0.002	<0.007
-15A	W-4.26M-9	<0.003	<0.002	<0.007
Limit of D	etection:	0.003	0.002	0.007
Method Ref	erence:	NIOSH 7300	(Modified)	

< = less than, below limit of detection</pre>

INORGANIC LABORATORY ANALYSES

Sample I.D.: See below Client: ENVIRONMENTAL HEALTH

Sample Received: 04/27/90 Client Ref. No.: 89-055

Sample Analyzed: 05/01/90 Lab Client Code: 100662

Sample Matrix: Wipe __ Lab No.: 9004182

				*
Batch Sub. No.	Sample Identification	Arsenic (mg/wipe)	Beryllium (mg/wipe)	Chromium (mg/wipe)
-16A	W-4.26M-10	<0.003	<0.002	<0.007
-17A	W-4.26M-11	<0.003	<0.002	<0.007
-18A	W-4.26M-12	<0.003	<0.002	<0.007
-19A	W-4.26M-Blank	<0.003	<0.002	<0.007
-21B	Method Blank	<0.003	<0.002	<0.007
Limit of D	etection:	0.003	0.002	0.007
Method Ref	erence:	NIOSH 7300	(Modified)	

< = less than, below limit of detection</pre>



INORGANIC LABORATORY ANALYSES

Sample I.D.:

See below

Client: ENVIRONMENTAL HEALTH

Sample Received:

04/27/90

Client Ref. No.:

89-055

Sample Analyzed:

05/01/90

Lab Client Code:

100662

Sample Matrix:

Wipe

Lab No.:

9004182

Batch Sub. No.	Sample Identification	Copper (mg/wipe)	Molybdenum (mg/wipe)	Lead (mg/wipe)
-07A	W-4.26M-1	0.007	<0.003	<0.003
-08A	W-4.26M-2	<0.007	<0.003	<0.003
-09A	W-4.26M-3	<0.007	<0.003	<0.003
-10A	W-4.26M-4	<0.007	<0.003	<0.003
-11A	W-4.26M-5	<0.007	<0.003	<0.003
-12A	W-4.26M-6	<0.007	<0.003	<0.003
-13A	W-4.26M-7	<0.007	<0.003	<0.003
-14A	W-4.26M-8	<0.007	<0.003	<0.003
-15A	W-4.26M-9	<0.007	<0.003	<0.003
Limit of D	etection:	0.007	0.003	0.003
Method Ref	erence:	NIOSH 7300) (Modified)	

< = less than, below limit of detection</pre>

Page 7 of 7

INORGANIC LABORATORY ANALYSES

Sample I.D.:

See below

Client: ENVIRONMENTAL HEALTH

Sample Received:

04/27/90

Client Ref. No.:

89-055

Sample Analyzed:

05/01/90

Lab Client Code:

100662

Sample Matrix:

Wipe

Lab No.:

9004182

Batch Sub. No.	Sample Identification	Copper (mg/wipe)	Molybdenum (mg/wipe)	Lead (mg/wipe)
-16A	W-4.26M-10	<0.007	<0.003	<0.003
-17A	W-4.26M-11	<0.007	<0.003	0.004
-18A	W-4.26M-12	<0.007	<0.003	<0.003
-19A	W-4.26M-Blank	<0.007	<0.003	<0.003
-21B	Method Blank	<0.007	<0.003	<0.003
Limit of D	etection:	0.007	0.003	0.003
Method Ref	erence:	NIOSH 7300	(Modified)	

< = less than, below limit of detection

CLAYTON ENVIRONMENTAL CONSULTANTS, INC. 22345 Roethel Drive Novi, Michigan 48050

Ms. Irene Fanelli President ENVIRONMENTAL HEALTH CONSULTANTS P.O. Box 117910 Burlington, CA 94011-7910

Date Reported: 7-MAY-90 Date Received: 28-APR-90 Clayton Project No. 65309-17

Client Job No. 89-055

0.006 mg

7904

0.003 mg 0.003 mg

7904

7904

Dear Ms. Fanelli:

Limit of Detection:

Analytical Method (NIOSH):

The following is our report on the samples submitted for analysis.

Table 1

		Cyanide							
Lab Number	Sample Description	Filter (mg)	Impinger (mg)	Total					
819325	AA-104291/A-4.26C-1	<0.003	<0.003	<0.006					
819326	AA-109841/A-4.26C-2	<0.003	<0.003	<0.006					
819327	AA-109845/A-4.26C-3	<0.003	<0.003	<0.006					
819328	AA-109840/A-4.26C-4	<0.003	<0.003	<0.006					
819329	AA-109849/A-4.26C-5	<0.003	<0.003	<0.006					
819330	AA-109842/A-4.26C-6	<0.003	<0.003	<0.006					
819331	AA-109843/A-4.26C-7	(0.003	<0.003	<0.006					
819332	AA-109848/A-4.26C-8	<0.003	<0.003	<0.006					
819333	AA-109846/A-4.26C-9	<0.003	<0.003	<0.006					
819334	AA-109847/A-4.26C-BLANK	<0.003	<0.003	<0.006					

ENVIRONMENTAL HEALTH CONSULTANTS Clayton Project No. 65309-17

Table 2

Lab Number	Sample Description	Cyanide (mg)
819335	W-4.26C-1	<0.003
819336	W-4.26C-2	<0.003
819337	W-4.26C-3	<0.003
819338	W-4.26C-4	(0.003
819339	W-4.26C-5	(0.003
819340	W-4.26C-6	⟨0.003
819341	W-4.26C-7	(0.003
819342	W-4.26C-8	<0.003
819343	W-4.26C-9	<0.003
819344	W-4.26C-10	(0.003
819345	W-4.26C-11	<0.003
819346	W-4.26C-12	<0.003
819347	W-4.26C-BLANK	<0.003

Limit of Detection: 0.003 mg Analytical Method (NIOSH): 7904

We appreciate the opportunity to be of assistance to you. Please call our Client Services Department at (313) 344-2650 or me if you have any questions.

Robert Lieckfield Jr., C.I.H. Manager, Laboratory Services