

Dames & Moore Job Number 05467-030-06
Salt Lake City, Utah
March 5, 1982



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40-3453

REPORT OF PHASE I WORK
GROUND WATER MONITORING PROJECT
URANIUM MILL TAILINGS IMPOUNDMENT
MOAB, UTAH
FOR ATLAS MINERALS

Dames & Moore



8205030136

Dames & Moore



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Salt Lake City, Utah 84111
(801) 521-9255
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March 5, 1982

Atlas Minerals
P. O. Box 1207
Moab, Utah 84532

Attention: Mr. Richard E. Blubaugh

Gentlemen:

Report of Phase I Work
Ground Water Monitoring Project
Uranium Mill Tailings Impoundment
Moab, Utah
For Atlas Minerals

INTRODUCTION

This letter summarizes the results of our Phase I work for the ground water monitoring of your mill tailings impoundment near Moab, Utah. Phase I consisted of the drilling, logging, sampling and completion of four wells within a single boring located southeast of the tailings impoundment. This work was authorized by your Professional Services Agreement dated September 1, 1981.

PURPOSE AND SCOPE

The purpose and scope of our study were outlined in your Request For Proposal dated July 16, 1981 and our proposal dated July 30, 1981. The purpose of Phase I was to complete and test two deep monitor wells specifically requested by the U.S. Nuclear Regulatory Commission in their letter dated June 24, 1981 (Attachment 1). The primary objectives outlined by NRC

related to the Phase I work were evaluation of whether the Colorado River is a recharging or a discharging stream, whether any possible seepage is moving toward or away from the river or possibly downward and subsequently in some other direction and evaluation of the hydrogeologic characteristics of bedrock.

Specifically, the following Phase I scope of services has been conducted:

1. Recommendations on the drilling, construction and field procedures for installation of the Phase I monitor wells.
2. Inspection of the drilling and completion of the Phase I wells located between the tailings impoundment and the Colorado River.
3. Collection of lithologic samples and preparation of drilling and lithologic logs for the wells.
4. Supervision of geophysical logging and interpretation of the geophysical well logs.
5. Sieve analysis of lithologic samples and gravel pack; checking of screen slot sizes; and selection of screened, cased and grouted intervals.
6. Inspection of the completion operations of the wells, including grouting, surfacing casing placement, capping, and well development.
7. The pumping and sampling of water quality samples in accordance with NRC recommendations.

8. Preparation of this summary report listing, describing and interpreting all pertinent data collected in the Phase I program.

PHASE I FIELD STUDIES

The field program consisted of drilling, logging, completion and ground water quality sampling of four 2-inch diameter wells completed within a single boring located approximately 250 feet southeast of the tailings impoundment as shown on Plate 1, Plot Plan.

The NRC specifically requested, in their letter dated June 24, 1981 (see Attachment 1), that:

1. The well be drilled using a mud-rotary rig or hollow-stem augers.
2. The depth to bedrock be determined.
3. The first well be screened in bedrock.

As the use of hollow-stem augers is impractical in gravels and also impractical at the anticipated well depths, common rotary drilling methods were attempted. Rotary drilling methods failed to provide a clean stable boring below 40 feet (see Appendix 1) and, therefore, after consulting with the NRC the boring was redrilled using cable-tool methods. Due to the high cost of drilling and the questionable value of locating the bedrock surface at depths below 400 feet, drilling was discontinued at 406 feet and hence, the depth to bedrock was not determined and no well was completed within the bedrock. The NRC approved these changes subject to the provision that a new well would be drilled deeper if the deepest well showed any signs of contamination with tailing pond waters.

Water quality samples collected at the end of the field program were turned over to Atlas Minerals whereupon the samples were split and tested. For parameters with short hold times (ammonia, conductivity, nitrate, nitrite, and total dissolved solids) were flown to Salt Lake City for immediate analysis. The Phase I field studies and data obtained are summarized in detail in the attached Appendix A, Field Studies.

DATA INTERPRETATION

The lithologic character of the materials encountered in the well to the 406 feet of depth drilled was relatively uniform consisting of sand and gravel alluvial sediments. The upper 20 feet of alluvium was reddish brown fine to coarse sand and gravel probably derived from recent deposition from Moab Wash drainage. Below 20 feet, gray and brown fine to coarse gravel and coarse sand composed of diverse lithologies including limestone, granitic rock and sandstone predominate and represent deposition by the Colorado River rather than the Moab Wash. Somewhat finer materials generally consisting of well-graded sand with some gravel occur from 230 to 290 feet in depth and in several thin strata at greater depth. These apparently represent deposition under a lower velocity regime. Only a very few layers of reddish brown sand indicative of deposition from the Moab Wash drainage were noted in the boring below 20 feet. No low permeability confining stratum was noted over the entire interval drilled, either in the cuttings or the geophysical logs.

Table 1 summarizes average concentration determined by the various laboratories for the initial samples. The certificates of analyses listing the results from each of the laboratories are given in Attachment 2.

All monitor wells encountered ground water which would be classified as brine, having total dissolved solids concentrations exceeding 100,000 milligrams per litre (mg/l). The principal constituent in the ground water is sodium chloride.

Atlas Minerals

March 5, 1982

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Waters from the three shallowest wells (Well S, Well IS and Well ID) are quite similar and reflect an extremely small, if any, increase in salinity with depth. Well D is somewhat more saline than the shallower wells. Major and trace ions show a similar relationship to that of general salinity and generally show no indication of downward or vertical mixing of tailings effluent. Radium-226 shows a slight decreasing concentration with depth in the analyses by Hazen Research which might be construed to suggest vertical mixing of tailings pond effluent. Analyses by Atlas Minerals of radium-226 do not show this trend; however, analyses by Atlas are all less than one-tenth the values reported by Hazen.

Values of lead reported by Hazen Research appear to be impossibly high for dissolved concentrations (samples were filtered with a 0.45 micron filter prior to preservation) even though the values were rechecked and confirmed by Hazen. In addition, the wide range in reported radium-226 values should be resolved. We recommend that the wells be resampled and analyzed at the end of the Phase II work when new wells are sampled.

Water level elevations measured in the four wells are all within 1.1 feet of each other and indicate a slightly lower elevation with increasing depth of the wells. However, to measure the true relative potentiometric surfaces in the three wells, corrections must be made for the differences in salinity of the water in the piezometers (wells). Considering salinity variations, there is negligible difference in the potentiometric surface of the four wells as would be expected from the lack of confining beds. Due to differences in the density of the brine and fresh water, there is no potential for Colorado River water to move vertically downward without completely mixing with the deeper waters. Similarly, tailings pond seepage with a total dissolved solids concentration on the order of 60,000 mg/l (as reported by Atlas Minerals) would not be able to move vertically downward without mixing with and attaining the same density as the brine.

Atlas Minerals
March 5, 1982
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CONCLUSIONS

We conclude there is negligible potential for Colorado River water or for seepage to move vertically downward into the brine without mixing with and attaining the same density as the brine. Water quality data show no indication of mixing with tailings pond seepage.

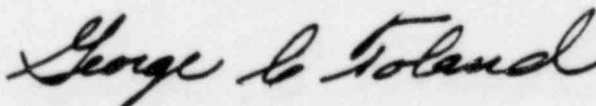
The presence of brines in the alluvial sediments and the resulting hydrologic relationships alleviate the need to locate the bedrock surface as well as the need to regularly monitor water qualities below the fresh water-brine interface. These needs are eliminated because of the low probabilities of downward migration of tailings pond waters through the brine and the low environmental hazard that would result from undetected tailings pond seepage water into the brine. All future monitoring programs should be directed towards the monitoring of the fresh water zone above the brine.

oOo

We have enjoyed performing these services for you. If you have any questions, please contact us.

Yours very truly,

DAMES & MOORE



George C. Toland
Partner

GCT/RLJ:f1

TABLE 1
SUMMARY OF WATER CHEMISTRY

	<u>Well S</u>	<u>Well IS</u>	<u>Well ID</u>	<u>Well D</u>
Screened Interval (ft)	145-155	219-229	300-310	394-396
Total dissolved solids (g/l) ¹⁾	111.1	112.6	112.6	125.2
Sodium (g/l) ²⁾	30.1	32.5	30.0	34.0
Chloride (g/l) ²⁾	51.8	54.3	54.5	55.7
Calcium (g/l) ³⁾	1.6	1.5	1.5	1.5
Sulfate (g/l) ²⁾	4.3	4.4	4.3	4.6
Field pH (units)	9.1	7.9	7.9	7.6
U-nat (pCi/l) ⁴⁾	<10 ⁴	<10 ⁴	<10 ⁴	<10 ⁴
Ra-226 (pCi/l) ³⁾	6.8	5.6	3.9	3.4
Ra-226 (pCi/l) ⁴⁾	0.36	0.19	0.36	0.34
Th-230 (pCi/l) ²⁾	0	0	0	0
Po-210 (pCi/l) ³⁾	0.4	0.0	1.7	0.9
Arsenic ³⁾ (mg/l)	< .001	< .001	< .001	< .001
Barium ³⁾ (mg/l)	1.99	1.58	1.44	1.61
Cadmium ³⁾ (mg/l)	0.298	0.311	0.311	0.334
Chromium ³⁾ (mg/l)	0.13	0.14	0.14	0.14
Lead ³⁾ (mg/l)	1.26	1.32	1.24	1.34
Manganese ³⁾ (mg/l)	0.49	0.560	0.550	0.475
Molybdenum ³⁾ (mg/l)	0.12	0.14	0.14	0.27
Nickel ³⁾ (mg/l)	3.51	3.84	3.85	4.10
Vanadium ³⁾ (mg/l)	0.38	0.36	0.44	0.18
Nitrate ⁵⁾ (mg/l)	1.20	0.90	0.60	5.90
Ammonia ⁵⁾ (mg/l)	3.32	3.16	3.12	3.13

-
- 1) Average of values reported by Hazen Research, Atlas Minerals and Ford Chemical.
2) Average values reported by Atlas Minerals and Hazen Research.
3) Values reported by Hazen Research.
4) Value reported by Atlas Minerals.
5) Value reported by Ford Chemical.



E 3,000

E 4,000

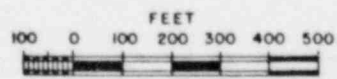
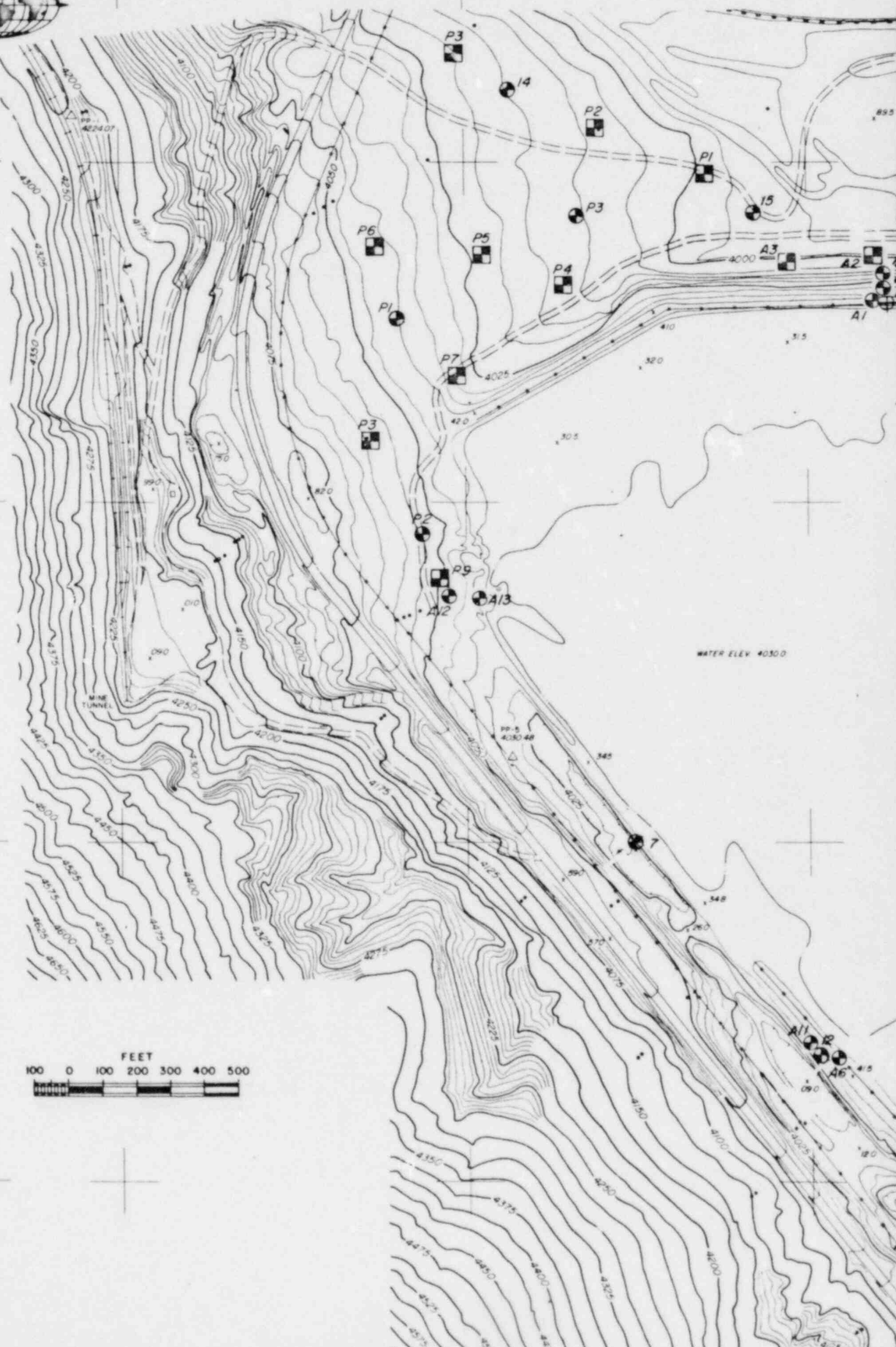
E 5,000

N 5,000

N 5,000

N 4,000

N 3,000



WATER ELEV 4030

BY CHECK



E 3,000

E 4,000

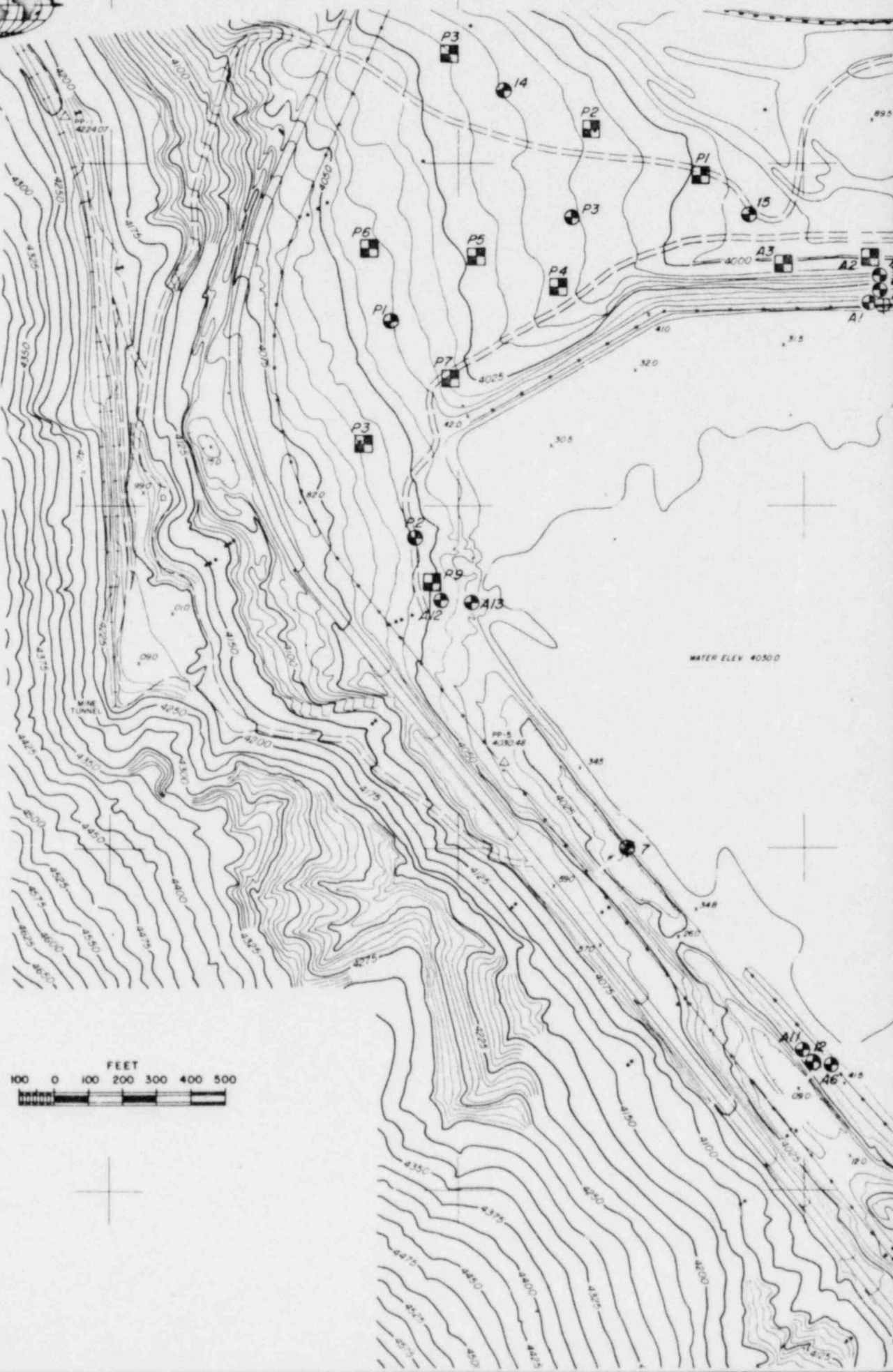
E 5,000

N 6,000

N 5,000

N 4,000

N 3,000



WATER ELEV 4030.0



BY: H. J. ...
CHECK: ...

ATTACHMENT 1

Letter From U.S. Nuclear Regulatory Commission

with the well cuttings. It is not sufficient simply to examine well cuttings as they are returned to the surface from drilling. It is not possible to obtain a reliable hydrostratigraphic section using well cuttings alone.

The alternative to the borehole geophysical logging program is to drill the well to bedrock with a hollow-stem auger and extract continuous samples with a shelly tube or a split-spoon sampler. However, this alternative will be difficult to implement if gravels are present as reported in the proposed study document. Gravels tend to plug the openings of split-spoon samplers or shelly tube samplers and prevent the acquisition of reliable undisturbed samples.

On the basis of the information obtained from the geophysical logs, the well cuttings and/or the shelly tube or split-spoon samples, decisions can be made regarding which sections of the casing should be slotted. Subsequently, decisions can be made regarding pump testing of the alluvial aquifer above the bedrock for purposes of defining the migration of the seepage plume. These data can be used also for the design of the remainder of the monitoring wells.

The second well should be drilled with mud rotary in the vicinity of the well described above. The purpose of this well should be to determine the hydrostratigraphy of the bedrock at the site and along with the first well, to establish the lateral hydraulic gradient and the extent of the seepage plume, if one exists, toward the river. This well should be drilled to bedrock and cased. The casing should be installed so as to seal off the alluvial aquifer above the bedrock. Drilling should continue into bedrock until it is determined that the bedrock is either an aquifer, an aquiclude, or an aquitard. Drilling should continue until an aquitard or an aquiclude has been encountered. This should be determined by an experienced hydrogeologist utilizing the geophysical logging techniques described above and observations during drilling. The aquiclude should be completely penetrated or drilled to a depth sufficient to verify the hypothesis that its thickness constitutes a barrier boundary to downward migration of seepage. A thickness of 30 to 40 feet should verify this hypothesis. If a bedrock aquifer exists either above or below a confining layer, the bedrock portion of the well should be designed so that the bedrock aquifer can be sampled separately from the alluvial aquifer.

These two wells should be developed and pumped and a sample withdrawn from each of them and submitted to a laboratory prior to continuation of drilling. It should be possible to utilize the specific electrical conductance, sulfate and pH to determine whether contamination exists in bedrock and whether additional monitor wells in bedrock are necessary.

The elevations of water levels in both wells, after casings have been slotted, should be measured so that the direction of the vertical component of the hydraulic gradient can be determined.

The location and design of additional monitoring wells along the southeast embankment of the impoundment should be based on the aforementioned resistivity survey, if one is conducted, and at the least, on the results of the two wells discussed above.

4. Due to the probability of seepage we recommend the placement of monitoring wells around the other sides of the pond as well. A resistivity survey might be used to locate initial monitoring wells on the other sides of the pond. These initial wells should also be designed and constructed according to Item 3 above so that they could also be used for aquifer pump tests, if appropriate. Such aquifer pump tests would provide the necessary hydraulic parameters prerequisite to predicting the size and rate of migration of the contaminated plume, if such a plume exists.
5. All well casing elevations must be determined so that the elevation of the water table in the alluvial aquifer can be contoured. Such a contour map will provide a description of the impact of the tailings pond on the water table and the direction and rate of groundwater movement in the vicinity of the site.
6. Prior to the collection of water quality samples from wells, all wells should be pumped until specific electrical conductance has stabilized during pumping.
7. Water quality samples shall be field filtered and preserved as indicated on Page 3 of the March 25, 1981 proposal, and uranium, vanadium, and gross beta added to the list, and gross gamma deleted. The first set of water quality samples should be sent to a laboratory that follows EPA recommended analytical procedures as well as to Atlas' laboratory, to ensure from the beginning of this study, that Atlas' water quality analysis techniques and data results are correct.
8. Atlas' response dated December 8, 1980 to NRC questions dated October 24 indicated that you were checking into the status of wells A, B, C, and the embankment well, which are listed as abandoned in table 2.3-4 of the report entitled "Safety Analysis Report," by Dames and Moore. Please indicate whether these wells are plugged and submit any available lithologic and hydrogeologic data on these wells.

The implementation of the aforementioned analytical hydrogeological procedures has become reasonably standard at other sites licensed by the NRC. Such studies, conducted by qualified hydrogeologists, have been successful in delineating contaminated plumes produced by uranium mill tailings ponds that were constructed prior to the implementation of groundwater protection technologies such as clay or synthetic liners. The probability of significant seepage at the Atlas-Moab site is reasonably high. The significance of the occurrence of seepage lies in the fact that seepage may be discharging into the Colorado River either directly southeast of the tailings pond or some

JUN 24 1981

distance downgradient therefrom. If this is the case, a pump back system to intercept seepage prior to its entry into the river may be required. If this is not the case (i.e., if the Colorado River is a losing stream at this location), then it is important to determine in what direction and at what depths the seepage plume is moving.

If you have any questions on our comments, please contact Mr. Pete Garcia of my staff.

Sincerely,

Harry J. Pettengill

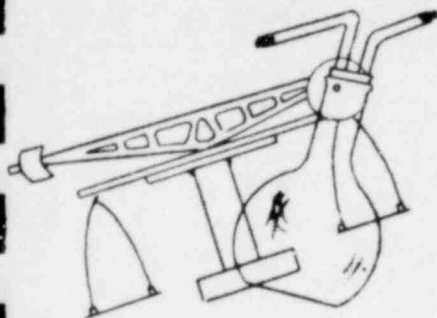
Harry J. Pettengill, Section Leader
Operating Facilities Section II
Uranium Recovery Licensing Branch
Division of Waste Management

File:

7/17/81 Discussed need for all logs and use of mud at any res
with H. Pettengill and P. Garcia. Pete to contact Ken Williams
and get back to me

ATTACHMENT 2

Certificates of Analyses



Ford Chemical

LABORATORY, INC.

Bacteriological and Chemical Analysis

40 WEST LOUISE AVENUE
SALT LAKE CITY, UTAH 84115

PHONE 466-8761

DATE: 01/27/82

CERTIFICATE OF ANALYSIS

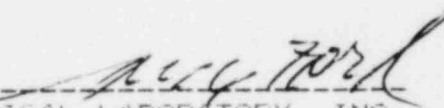
DAMES & MOORE, ENG.
250 E. BROADWAY
SALT LAKE CITY, UT
84111

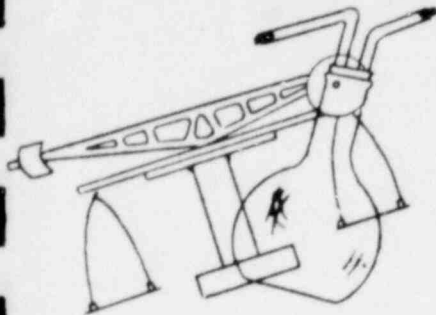
82-009050

SAMPLE: WELL WATER SAMPLES FROM ATLAS PROJECT, JOB #05467-030-06
COLLECTED 1-19-82, RECEIVED 1-20-82 FOR ANALYSIS.
UNDER P.O. #SL 1510.

WELL D WELL S
13:00 13:00

	=====	=====
Ammonia as NH ₃ -N mg/l	3.13	3.32
Conductivity umhos/cm	192,000	170,000
Nitrate as NO ₃ -N mg/l	5.90	1.20
Nitrite as NO ₂ -N mg/l	.32	.18
Total Diss. Solids Mg/l	144,500	125,380


FORD CHEMICAL LABORATORY, INC.



Ford Chemical

LABORATORY, INC.

Bacteriological and Chemical Analysis

40 WEST LOUISE AVENUE
SALT LAKE CITY, UTAH 84115

PHONE 466-8761

DATE: 01/26/81

CERTIFICATE OF ANALYSIS

DAMES & MOORE, ENG.
250 E. BROADWAY
SALT LAKE CITY, UT
84111

82-009060

SAMPLE: WELL WATER SAMPLE FROM ATLAS PROJECT, JOB #05467-030-06
RECEIVED 1-20-82 FOR ANALYSIS.

ATP	ATP
1-IS	1-ID
81	81

Ammonia as NH3-N mg/l	3.16	3.12
Nitrate as NO3-N mg/l	.90	.60
Nitrite as NO2-N mg/l	.12	.08
Total Diss. Solids Mg/l	125,000	127,500

[Signature]
FORD CHEMICAL LABORATORY, INC.

DAMES & MOORE		
SALT LAKE CITY		
RECEIVED		
JAN 27 '82		
GCT	GWS	PTC
WJG	WCH	LAC
LTB		
ADP		
SP		
SL		
DB		
EL		
CP		

HAZEN**Hazen Research, Inc.**4601 Indiana St • Golden, Colo 80401
Tel: (303) 279-4501 • Telex 45-860**REPORT OF ANALYSIS**

Atlas Minerals

Date February 11, 1982
HRI Project No. 4893
HRI Series No. 23111-2
Date Received 1/27/82
Customer P.O. A-10162

Element Found	S
Arsenic, recoverable, mg/l	<0.001 - <0.01 PPM
Barium, recoverable, mg/l	1.99 —
Cadmium, recoverable, mg/l	0.298 —
Calcium, recoverable, mg/l	1570 —
Chromium, recoverable, mg/l	0.13 —
Copper, recoverable, mg/l	0.18 - .19 PPM
Iron, recoverable, mg/l	1.22 - 1.2 PPM
Lead, recoverable, mg/l	1.26 —
Magnesium, recoverable, mg/l	472 - 615 PPM
Manganese, recoverable, mg/l	0.490 - .15 PPM
Molybdenum, recoverable, mg/l	0.12 —
Nickel, recoverable, mg/l	3.51 —
Potassium, recoverable, mg/l	749 - 1580 PPM
Sodium, recoverable, g/l	35.2 - 25 G/L
Vanadium, recoverable, mg/l	0.38 —
Zinc, recoverable, mg/l	0.21 —
Carbonate as CaCO ₃ , mg/l	<5 —
Bicarbonate as CaCO ₃ , mg/l	40 —
Chloride, g/l	59.2 44.5 G/L
Specific Conductance µmho/cm 25°C	152,600 - 143,600
Fluoride, mg/l	0.98 —
Solids, dissolved, g/l	100.3 - 107.6 G/L
Sulfate as SO ₄ , mg/l	4420 - 4150 ± 20 PPM
Selenium, recoverable, mg/l	0.002 - 2.1 PPM
Uranium, recoverable, mg/l	<0.002 - SAME AS BLANK

By: Robert RostadRobert Rostad
Manager, General Analytical

mem

REPORT OF ANALYSIS

Atlas Minerals

Date February 11, 1982
HRI Project No. 4893
HRI Series No. 23111-4
Date Received 1/27/82
Customer P.O. A-10162

Element Found	IS
Arsenic, recoverable, mg/l	<0.001 <i>2.01 PPM</i>
Barium, recoverable, mg/l	1.58 —
Cadmium, recoverable, mg/l	0.311 —
Calcium, recoverable, mg/l	1500
Chromium, recoverable, mg/l	0.14 —
Copper, recoverable, mg/l	0.19 <i>.213 PPM</i>
Iron, recoverable, mg/l	3.30 <i>2.5 PPM</i>
Lead, recoverable, mg/l	1.32 —
Magnesium, recoverable, mg/l	582 <i>578 PPM</i>
Manganese, recoverable, mg/l	0.560 — <i>.22 PPM</i>
Molybdenum, recoverable, mg/l	0.14 —
Nickel, recoverable, mg/l	3.84 —
Potassium, recoverable, mg/l	895 <i>1615 PPM</i>
Sodium, recoverable, g/l	37.6 <i>97.5 G/L</i>
Vanadium, recoverable, mg/l	0.36 —
Zinc, recoverable, mg/l	0.20 —
Carbonate as CaCO ₃ , mg/l	<5 —
Bicarbonate as CaCO ₃ , mg/l	119 —
Chloride, g/l	61.5 <i>46.9 G/L</i>
Specific Conductance µmho/cm 25°C	160,200 — <i>120,000</i>
Fluoride, mg/l	1.67 —
Solids, dissolved, g/l	105.1 <i>107.9 G/L</i>
Sulfate as SO ₄ , mg/l	4690 — <i>4174 PPM</i>
Selenium, recoverable,	0.001 <i>1.1 PPM</i>
Uranium, recoverable, mg/l	<0.002 <i>Same as Blank</i>

By: Robert Rostad
Manager, General Analytical

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Hazen Research, Inc.
4601 Indiana St • Golde Jolo 80401
Tel (303) 279-4501 • Telex 45-860

REPORT OF ANALYSIS

Atlas Minerals

Date February 11, 1982
HRI Project No. 4893
HRI Series No. 23111-3
Date Received 1/27/82
Customer P.O. A-10162

Element Found	ID
Arsenic, recoverable, mg/l	<0.001 - .01 PPM
Barium, recoverable, mg/l	1.44 -
Cadmium, recoverable, mg/l	0.311 -
Calcium, recoverable, mg/l	1510
Chromium, recoverable, mg/l	0.14 -
Copper, recoverable, mg/l	0.20 - .205 PPM
Iron, recoverable, mg/l	3.20 - 3.0 PPM
Lead, recoverable, mg/l	1.24 -
Magnesium, recoverable, mg/l	577 - 857 PPM
Manganese, recoverable, mg/l	0.550 - .17 PPM
Molybdenum, recoverable, mg/l	0.14 -
Nickel, recoverable, mg/l	3.85 -
Potassium, recoverable, mg/l	789 - 1450 PPM
Sodium, recoverable, g/l	39.1 - 25.0 PPM
Vanadium, recoverable, mg/l	0.44 -
Zinc, recoverable, mg/l	0.21 -
Carbonate as CaCO ₃ , mg/l	<5 -
Bicarbonate as CaCO ₃ , mg/l	98 -
Chloride, g/l	62.0 - 47.0 PPM
Specific Conductance, umho/cm 25°C	161,200 - 135,000
Fluoride, mg/l	1.37 -
Solids, dissolved, g/l	106.3 - 103.9 G/L
Sulfate as SO ₄ , mg/l	4600 - 3986 PPM
Selenium, recoverable, mg/l	0.002 - .1 PPM
Uranium, recoverable, mg/l	<0.002 - SAME AS BLANK

By: Robert Rostad
Robert Rostad
Manager, General Analytical

mem



Hazen Research, Inc.
 4601 Indiana St • Golden, Colo 80401
 Tel (303) 279-4501 • Telex 45-860

REPORT OF ANALYSIS

Atlas Minerals

Date February 11, 1982
 HRI Project No. 4893
 HRI Series No. 23111-1
 Date Received 1/27/82
 Customer P.O. A-10162

Element Found	D	
Arsenic, recoverable, mg/l	<0.001	- .01 PPM
Barium, recoverable, mg/l	1.61	—
Cadmium, recoverable, mg/l	0.334	—
Calcium, recoverable, mg/l	1550	- 879 PPM
Chromium, recoverable, mg/l	0.14	—
Copper, recoverable, mg/l	0.21	- .22 PPM
Iron, recoverable, mg/l	9.48	- 9.6 PPM
Lead, recoverable, mg/l	1.34	—
Magnesium, recoverable, mg/l	604	- 857 PPM
Manganese, recoverable, mg/l	0.475	- .16 PPM
Molybdenum, recoverable, mg/l	0.27	—
Nickel, recoverable, mg/l	4.10	—
Potassium, recoverable, mg/l	891	- 1555 PPM
Sodium, recoverable, g/l	39.6	- 28.5 G/L
Vanadium, recoverable, mg/l	0.18	—
Zinc, recoverable, mg/l	0.34	—
Carbonate as CaCO ₃ , mg/l	<5	—
Bicarbonate as CaCO ₃ , mg/l	107	—
Chloride, g/l	62.4	- 48.97 G/L
Specific Conductance, umho/cm 25°C	166,600	- 150,400
Fluoride, mg/l	2.02	—
Solids, dissolved, g/l	110.3	- 120.7 G/L
Sulfate as SO ₄ , mg/l	4720	- 4441 PPM
Selenium, recoverable, mg/l	<0.001	- 2.1 PPM
Uranium, recoverable, mg/l	<0.002	- Same As Blank

By: Robert Rostad
 Robert Rostad
 Manager, General Analytical

mem



Hazen Research, Inc.
 4601 Indiana St • Golden, Colo 80401
 Tel (303) 279-4501 • Telex 45-860

REPORT OF ANALYSIS

Atlas Minerals
 North Highway 163 Mill Site
 Moab, Utah 84532

Date February 11, 1982
 HRI Project No. 4893
 HRI Series No. 23111
 Date Received 1/27/82
 Customer P.O. A-10162

Analysis No.	Sample Designation	pCi/l Ra ²²⁶	recoverable ± Precision*	
23111-1	D	3.4	± 2.2	= 2.34×10^{-8} uCi/ml
-2	S	6.8	± 2.5	= 0.19×10^{-8} uCi/ml
-3	ID	5.6	± 2.7	= 0.36×10^{-8} uCi/ml
-4	IS	3.9	± 1.9	= 0.26×10^{-8} uCi/ml

By: Robert Rostad
 Robert Rostad
 Manager, General Analytical

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*Variability of the radioactive disintegration process (counting error) at the 95% confidence level, 1.96σ.

NOTE: Lead 210 results to follow.



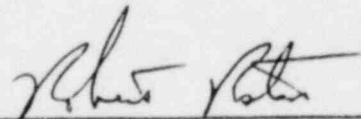
Hazen Research, Inc.
4601 Indiana St • Golden, Colo. 80401
Tel (303) 279-4501 • Telex 45-860

REPORT OF ANALYSIS

Atlas Minerals

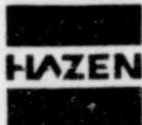
Date February 11, 1982
HRI Project No. 4893
HRI Series No. 23111
Date Received 1/27/82
Customer P.O. A-10162

Analysis No.	Sample Designation	pCi/l Po ²¹⁰	recoverable ± Precision*
23111-1	D	0.9	± 2.4 0.0 pCi/ml
-2	S	0.4	± 2.2 0.0 pCi/ml
-3	ID	1.7	± 2.7 .00015 pCi/ml
-4	IS	0.0	± 1.9 0.0 pCi/ml

By: 
Robert Rostad
Manager, General Analytical

mem

*Variability of the radioactive disintegration process (counting error) at the 95% confidence level, 1.96σ.



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REPORT OF ANALYSIS

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Date February 11, 1982
 HRI Project No. 4893
 HRI Series No. 23111
 Date Received 1/27/82
 Customer P.O. A-10162

Analysis No.	Sample Designation	pCi/l Th ²³⁰	recoverable ± Precision*	
23111-1	D	0	± 16	0 ± 0 × 10 ⁻⁶ dpm/ml
-2	S	0	± 20	0 ± 0 × 10 ⁻⁶ dpm/ml
-3	ID	0	± 16	0 ± 0 × 10 ⁻⁶ dpm/ml
-4	IS	0	± 20	0 ± 0 × 10 ⁻⁶ dpm/ml

117132

By: Robert Rostad
 Robert Rostad
 Manager, General Analytical

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*Variability of the radioactive disintegration process (counting error) at the 95% confidence level, 1.96σ.

MONITOR WELL REPORTS

Quarter

Well # Well 5

Date and time sample was collected January 20, 1982

Location of sample _____

Sampling method used (bailed or pumped) Pumped

The amount of water to be removed prior to sampling _____

<u>Radionuclide</u>	<u>M.P.C.</u>	<u>Concentration uci/ml</u>	<u>Error Estimate</u>
Gross Beta-Gamma		<u>Nil x 10⁻⁶ uci/ml</u>	<u>±0</u>
U-Nat	<u>3 x 10⁻⁵ uci/ml</u>	<u>Nil x 10⁻⁵ uci/ml</u>	<u>±0</u>
Ra ²²⁶	<u>3 x 10⁻⁸ uci/ml</u>	<u>.026 x 10⁻⁸ uci/ml</u>	<u>±.06</u>
Th ²³⁰	<u>2 x 10⁻⁶ uci/ml</u>	<u>0 x 10⁻⁶ uci/ml</u>	<u>±0</u>
Pb ²¹⁰	<u>1 x 10⁻⁷ uci/ml</u>	_____	_____
Po ²¹⁰	<u>7 x 10⁻⁷ uci/ml</u>	<u>0.0 Pci/ml</u>	<u>±.0009</u>

<u>Common Ions and Trace Metals</u>	<u>Concentration</u>	<u>Error Estimate</u>
K ⁺	<u>1580 PPM</u>	<u>±0</u>
Na ⁺	<u>25000 PPM</u>	<u>±0</u>
Cl ⁻	<u>44,485 PPM</u>	<u>±481</u>
SO ⁼	<u>4150 PPM</u>	<u>±200</u>
NO ⁴⁻	<u>.011 G/L</u>	<u>±.009</u>
Fe ³	<u>1.2 PPM</u>	<u>±0</u>
Mn	<u>.15 PPM</u>	<u>±0</u>
As	<u><.01 PPM</u>	<u>±0</u>
Se	<u><.1 PPM</u>	<u>±0</u>
TDS	<u>107,638</u>	<u>±4236</u>
Conductivity	<u>143,600</u>	<u>umhos</u>
Ph	<u>9.0</u>	_____
Cu	<u>.193 PPM</u>	<u>±.02</u>

MONITOR WELL REPORTS

_____Quarter_____

Well # Well S1

Date and time sample was collected January 20, 1982

Location of sample _____

Sampling method used (bailed or pumped) Pumped

The amount of water to be removed prior to sampling _____

<u>Radionuclide</u>	<u>M.P.C.</u>	<u>Concentration uci/ml</u>	<u>Error Estimate</u>
Gross Beta-Gamma		<u>Nil x 10⁻⁶ uci/ml</u>	<u>±0</u>
U-Nat	<u>3 x 10⁻⁵ uci/ml</u>	<u>Nil x 10⁻⁵ uci/ml</u>	<u>±0</u>
Ra ²²⁶	<u>3 x 10⁻⁸ uci/ml</u>	<u>.019 x 10⁻⁸ uci/ml</u>	<u>±035</u>
Th ²³⁰	<u>2 x 10⁻⁶ uci/ml</u>	<u>0 x 10⁻⁶ uci/ml</u>	<u>±0</u>
Pb ²¹⁰	<u>1 x 10⁻⁷ uci/ml</u>	_____	_____
Po ²¹⁰	<u>7 x 10⁻⁷ uci/ml</u>	<u>.00015 Pci/ml</u>	<u>±.0004</u>

<u>Common Ions and Trace Metals</u>	<u>Concentration</u>	<u>Error Estimate</u>
K ⁺	<u>1615 PPM</u>	<u>±45</u>
Na ⁺	<u>27,500 PPM</u>	<u>±4492</u>
Cl ⁻	<u>46,985 PPM</u>	<u>±319</u>
SO ⁼	<u>4174 PPM</u>	<u>±314</u>
NO ⁴⁻	<u>.015 g/L</u>	<u>±.0046</u>
Fe ³	<u>2.5 PPM</u>	<u>±.44</u>
Mn	<u>.22 PPM</u>	<u>±.04</u>
As	<u><.01 PPM</u>	<u>±0</u>
Se	<u><.01 PPM</u>	<u>±0</u>
TDS	<u>107,859</u>	<u>±2412</u>
Conductivity	<u>121,250</u>	<u>umhos</u>
Ph	<u>7.9</u>	_____
Cu	<u>.213 PPM</u>	<u>±.02</u>

MONITOR WELL REPORTS

Quarter

Well # Well D1

Date and time sample was collected January 20, 1982

Location of sample _____

Sampling method used (bailed or pumped) _____

The amount of water to be removed prior to sampling _____

<u>Radionuclide</u>	<u>M.P.C.</u>	<u>Concentration uci/ml</u>	<u>Error Estimate</u>
Gross Beta-Gamma		<u>1.13 x 10⁻⁶ uci/ml</u>	<u>±.17</u>
U-Nat	<u>3 x 10⁻⁵ uci/ml</u>	<u>Nil x 10⁻⁵ uci/ml</u>	<u>█</u>
Ra ²²⁶	<u>3 x 10⁻⁸ uci/ml</u>	<u>.036 x 10⁻⁸ uci/ml</u>	<u>±.058</u>
Th ²³⁰	<u>2 x 10⁻⁶ uci/ml</u>	<u>0 x 10⁻⁶ uci/ml</u>	<u>±0</u>
Pb ²¹⁰	<u>1 x 10⁻⁷ uci/ml</u>	<u>_____</u>	<u>_____</u>
Po ²¹⁰	<u>7 x 10⁻⁷ uci/ml</u>	<u>.0.0 Pci/ml</u>	<u>±0</u>

<u>Common Ions and Trace Metals</u>	<u>Concentration</u>	<u>Error Estimate</u>
K ⁺	<u>1450 PPM</u>	<u>±90</u>
Na ⁺	<u>25,000 PPM</u>	<u>±0</u>
Cl ⁻	<u>47,017 PPM</u>	<u>±1182</u>
SO ⁼	<u>3986 PPM</u>	<u>±136</u>
NO ⁴⁻	<u>.010 g/l</u>	<u>±0</u>
Fe ³	<u>3.0 PPM</u>	<u>±.45</u>
Mn	<u>.17 PPM</u>	<u>±0</u>
As	<u><.01 PPM</u>	<u>±0</u>
Se	<u><.01 PPM</u>	<u>±0</u>
TDS	<u>103,982 PPM</u>	<u>±10,927</u>
Conductivity	<u>120,000</u>	<u>umhos</u>
Ph	<u>7.9</u>	<u>_____</u>
Cu	<u>.205 PPM</u>	<u>±0</u>

APPENDIX A

FIELD PROGRAM

INTRODUCTION

The field program consisted of drilling, logging, completion, and ground water quality sampling of four wells completed within a single boring. The boring was drilled using both rotary and cable-tool drilling methods and was located approximately 250 feet southeast of the tailings impoundment. The boring was drilled to a total depth of 406 feet in the Colorado River alluvium. Each of the wells installed within the boring were completed using 2-inch PVC casing and screen and were hydraulically isolated from each other through the use of grout seals. All field operations were conducted and supervised by an experienced Dames & Moore hydrogeologist.

DRILLING METHODS

GENERAL

The drilling was done by Zimmerman Drilling, Inc. of Moab, Utah, using both a Speedstar SS-15 rotary and a Backus Erie cable-tool drilling rig.

ROTARY DRILLING

As open-hole geophysical logs of the boring were desired, our first attempt at drilling the boring was made using rotary-wash drilling techniques. Prior to drilling, a mud pit was dug and an organic polymer drilling fluid was mixed. The mixing of drilling fluids and the control of mud viscosities were supervised by drilling mud engineers furnished by Nova Mud Corporation.

The Marsh Funnel Viscosity of the drilling mud was adjusted to approximately 60 seconds prior to the start of drilling. Drilling to 30 feet proceeded without difficulty; however, drilling after 30 feet failed to produce a clean stable hole. The Marsh Funnel Viscosity was then increased to over 100 seconds and drilling continued to 40 feet.

During the drilling within the interval between 40 and 60 feet, the problems of hole stability increased. The drilling mud viscosity was raised to 230 seconds (Marsh Funnel). The sample cuttings returned to the surface indicated that the sediments drilled were very coarse gravels and that the drilling mud was of sufficient viscosity to carry the drill cuttings. Caving of the hole, however, continued. In an attempt to stabilize the caving, a solution was added to the drilling mud that caused the mud to gel within the formation. This procedure also failed to stabilize the boring. At 60 feet the driller started to lose circulation to the formation. At this point the use of rotary drilling methods was discontinued.

It was the opinion of the personnel from Zimmerman Drilling, Nova Mud Corporation, and Dames & Moore that the use of other drilling mud types such as bentonite would also fail to produce a clean, stable boring and that the use of cable-tool drilling methods was required.

CABLE-TOOL DRILLING

Drilling using the cable-tool method was continued at the same location. A 10-3/4-inch surface casing was driven to 12 feet and bailed clean. The drive casing was 8-7/8-inch with a 8-inch drive shoe. The casing was alternately driven and bailed clean. Casing drive rates varied from approximately one foot per minute to one foot per hour. The bailing of cuttings from the boring was generally done within the casing as the hole generally caved when the boring was advanced more than a foot beyond the bottom of the casing. The casing was still capable of being advanced when the decision was made to stop drilling at 406 feet.

GEOLOGIC AND GEOPHYSICAL LOGGING

GENERAL

During the drilling phase, cutting samples were collected and saved at five foot intervals. Rotary drill cuttings were obtained by settling the cuttings obtained in the returned drilling muds. Cuttings from the cable-tool drilling were obtained from the bailer. After completion of the boring, the hole was geophysically logged through the steel casing.

GEOLOGIC LOG

The cutting samples collected during the drilling phase were visually analyzed and then described. These cuttings are the basis for the geologic log presented in Plate A-1, Log of Boring, and the following discussion. The sample classifications and descriptions given on Plate A-1 are based on the terminology and methods of classification recommended in the Unified Soil Classification System. A summary of this classification system is presented in Plate A-2, Unified Soil Classification System.

The boring penetrated only alluvial sediments composed primarily of gravels and coarse sands. As both rotary and cable-tool methods tended to crush the individual grains of gravel, few grains larger than two centimeters were found in the drill cuttings. However, based on observations of fractured and unfractured surfaces, the approximate grain size could be estimated.

The upper 25 feet of the boring was composed of sands and gravels. In this interval the sediments were generally reddish brown and individual gravel grains tended to be angular. The grains consisted primarily of reddish brown sandstone and shale fragments. Below 25 feet the sediments were generally light brownish gray and individual gravel grains tended to be rounded to well-rounded and were composed primarily of granite, gniess, quartzite, and limestone rock fragments. A few beds of coarse sand were also present.

The differences between the sediments of the upper 25 feet and the rest suggests that two different source areas and alluvial systems have provided sediments into this area. The upper 25 feet appear to have been the result of sediment deposition in the Moab Wash alluvial system. The sediments deeper than 25 feet appear to be related to sediment deposition along the Colorado River alluvial system.

GEOPHYSICAL LOGS

Three cased hole geophysical logs were run in the boring by Professional Logging Services, Inc. These logs were natural gamma, gamma-gamma density, and neutron-neutron logs. These geophysical logs are shown on Plate A-1.

NATURAL GAMMA RAY LOG

The natural gamma ray log responds to the presence of gamma ray radiation in the formational materials surrounding the bore hole. In alluvial materials, such as those present at this site, the majority of the natural gamma radiation is produced by the radioactive elements commonly found in and absorbed to the clay minerals, in particular, radioactive potassium found in illitic clays. Hence, in general, the presence of clay layers are represented by high counting rates, while clean sands and gravels are generally associated with low gamma counting rates.

The natural gamma log for this well shows little, if any, significant change in the natural gamma intensity for sediments located below the water table (13 feet). This lack of change indicates that there are no clay layers of sufficient thickness to be considered a confining bed.

GAMMA-GAMMA DENSITY LOG

The gamma-gamma density log provides a measure of the insitu density of formational materials. In this log gamma rays are emitted into the formation from a gamma ray source in the logging tool. The percentage of this gamma ray radiation returning from the formation to the detector, is measured. In general, the percentage of radiation returned to the tool is a function of the density of formational materials, as the absorption of gamma radiation is a function of density. However, because the borehole is cased, the density of the casing is also interpreted by the logging tool as part of the formation. Hence, quantitative measurement of formation density cannot be made with the log.

The gamma-gamma density log obtained from this boring shows little, if any, significant change over its entire depth. This probably results from both the lack of any significant formational changes and also the attenuating effects of the steel casing.

NEUTRON-NEUTRON LOG

The neutron-neutron log provides a measure of the formation porosity. In this tool, epithermal neutrons are emitted from a neutron source into the formation and the number of neutrons returning from the formation to the detector, are counted. As the hydrogen atom is the principal absorber of epithermal neutrons, the number of neutrons returning is related principally to the concentration of hydrogen atoms in the formational material. For alluvial sediments such as those found in this boring, the majority of the hydrogen atoms present are located within the sediment pore water, and hence, the neutron-neutron log is an indirect measurement of the formation porosity.

The neutron-neutron log of this boring shows few significant changes, if any, in the intensity of the returned neutron as a function of depth. The changes in intensity appear to be related to changes in the sediment porosity; however, the actual change in porosity of the granular aquifer is likely only a few percent.

WELL COMPLETION

The 8-7/8-inch steel drive casing was perforated at four separate intervals and four monitor wells were installed in these perforated zones. Each well was separately sand-packed and sealed to prevent the migration of ground water between wells. The location and construction details for each of these wells are shown graphically on Plate A-1 and are discussed below.

The 8-7/8-inch steel drive casing inplaced while drilling the well, was perforated at selected intervals using a mechanical knife casing perforator. Each perforation cut by the perforating knife was approximately three inches long and one-quarter inch wide. The locations of the perforations are shown graphically on Plate A-1. Development of the well was done by bailing the perforated casing after the perforations were cut.

Four 2-inch diameter casing and screen assemblies were then sequentially set, sand-packed and grouted into place. The screen and casing pipe used was 2-inch Schedule 40 PVC pipe. The screen was manufactured by Hydrophyllic, Inc. of Tacoma, Washington and had 0.020 inch wide saw-cut perforations. Each pipe joint was properly glued prior to installation. Glued pipe joints that were not allowed more than one hour to set were given additional strength through the installation of two, 3/16-inch stainless steel screws, screwed through the glued joint.

Each casing-screen assembly was sand-packed and grouted immediately after its installation. The sand-pack used was a Number 4-10 sand manufactured by Colorado Silica Sand, Inc., Pueblo, Colorado. The grout mixture was a neat cement grout made by mixing 5.2 gallons of water and 2 pounds of calcium chloride with each bag of Type V portland cement. The calcium chloride was added to accelerate the grout setting time. The grout was tremied into place and allowed a minimum of eight hours to set prior to the installation of the next casing-screen assembly. The locations of all casings, screens, sand-pack, and grout seals are shown graphically on Plate A-1.

Following the construction of all wells, each well was individually tested for the presence of a hydraulic interconnection between other casing-screen assemblies. These tests were conducted by measuring the water levels of each piezometer and then pumping one of the piezometers for approximately 30 minutes and again measuring the water levels. No significant changes in the water levels of the unpumped wells were noted while pumping any of the other wells. The pumping level in the pumped wells varied from 2 to 8 feet.

WATER QUALITY SAMPLING

Each of the wells were pumped prior to sampling until it was determined that there was no longer any significant change in the specific conductivity of the ground water being pumped. The wells were pumped until three successive measurements of the ground water's specific conductivity showed no significant change. A minimum of 1,500 gallons was pumped from each well. The volumes of water pumped, specific conductivity, pH, temperature and water level measurements for each of the wells prior to sampling are given in Table A-2A through A-2D, Ground Water Sampling Forms.

Samples for parameters with short-hold times (ammonia, nitrate, nitrite, and total dissolved solids) were immediately shipped by air for analysis by Ford Chemical Laboratories, Inc., Salt Lake City, Utah. Ground water samples for other parameters were split and analyzed by Hazen Research, Inc., Golden,

Colorado, and by Atlas Minerals, Moab, Utah. The results of the analyses are given in Attachment 2.

GROUND WATER QUALITY SAMPLING FORMWell No.: ATP-1-S-81Owner: Atlas MineralsDate/Time 1/19/82 / 1:00 Sampled By: Dale Edwards (Atlas Minerals)Appearance: Clear
(clear, colored, turbid, sediment, etc.)Remarks: Well had been pumped on the previous day at 12 gpm for approximately two hours (pumped 1,500 gal).Field Measurements and Sample Collection (a)Static Water Level: 14.26 (16.86 from TOC) (in feet below land surface)Pumping Rate: 8 gpm (in gallons per minute)Duration of Pumping: from 10:30 to 1:00, Total Time = 170 (in minutes)Volume of Water Pumped Prior to Sampling (b): 1,400 (in gallons)Method of Pumping: Centrifugal pumpElectrical Conductivity (E.C.) MeasurementsMeter No.: YSI #1Time Sampled: 10:43 pH 9.9 (E.C.) 158,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 11:01 pH 9.2 (E.C.) 172,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 11:05 pH 9.0 (E.C.) 158,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 11:15 pH 9.1 (E.C.) 130,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 12:15 pH 9.1 (E.C.) 130,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 12:45 pH 9.1 (E.C.) 145,000 (in micromhos/cm, corrected to 25°C)Meter No.: Digital pH Meter #609 standardized to pH 10Temperature: 18 °C

(a) Field measurements and sample collection made in accordance with Dames & Moore Technical Memorandum for the pre-operational ground water.

(b) A minimum of two (2) casing volumes of water shall be pumped from the well prior to sample collection.

GROUND WATER QUALITY SAMPLING FORMWell No.: ATP-1-IS-81Owner: Atlas MineralsDate/Time 1/20/82 / 11:00 Sampled By: Dale Edwards (Atlas Minerals)Appearance: Clear
(clear, colored, turbid, sediment, etc.)Remarks: 3,000 gallons pumped from the well on 1-19-82Field Measurements and Sample Collection^(a)Static Water Level: 14.71 (17.46 from TOC) (in feet below land surface)Pumping Rate: 15 (in gallons per minute)Duration of Pumping: from 9:06 to 11:15, Total Time = 129 (in minutes)Volume of Water Pumped Prior to Sampling^(b): 1,935 (in gallons)Method of Pumping: Centrifugal pumpElectrical Conductivity (E.C.) MeasurementsMeter No.: Yellow Spring Instrument #1Time Sampled: 9:55 pH 7.9 (E.C.) 115,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 10:15 pH 7.9 (E.C.) 120,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 10:25 pH 7.9 (E.C.) 125,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 10:45 pH 7.9 (E.C.) 125,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 11:00 pH 7.9 (E.C.) 125,000 (in micromhos/cm, corrected to 25°C)

Time Sampled: _____ pH _____ (E.C.) _____ (in micromhos/cm, corrected to 25°C)

Meter No.: Digital pH Meter #609 standardized to pH 7Temperature: 15 °C

(a) Field measurements and sample collection made in accordance with Dames & Moore Technical Memorandum for the pre-operational ground water.

(b) A minimum of two (2) casing volumes of water shall be pumped from the well prior to sample collection.

GROUND WATER QUALITY SAMPLING FORMWell No.: ATP-1-ID-81Owner: Atlas MineralsDate/Time 1/20/81 / 10:30 Sampled By: Dale Edwards (Atlas Minerals)Appearance: Clear
(clear, colored, turbid, sediment, etc.)Remarks: Well had been pumped on the previous day at approximately 10 gpm for 1.5 hours
(volume pumped was 1,500 gallons)Field Measurements and Sample Collection^(a)Static Water Level: 14.65 (17.55 from TOC) (in feet below land surface)Pumping Rate: 12 (in gallons per minute)Duration of Pumping: from 8:50 to 10:31, Total Time = 101 (in minutes)Volume of Water Pumped Prior to Sampling^(b): 1,200 (in gallons)Method of Pumping: Centrifugal pumpElectrical Conductivity (E.C.) MeasurementsMeter No.: Yellow Springs Instrument #1Time Sampled: 9:45 pH 7.4 (E.C.) 135,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 10:00 pH 7.7 (E.C.) 135,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 10:16 pH 7.9 (E.C.) 130,000 (in micromhos/cm, corrected to 25°C)

Time Sampled: _____ pH _____ (E.C.) _____ (in micromhos/cm, corrected to 25°C)

Time Sampled: _____ pH _____ (E.C.) _____ (in micromhos/cm, corrected to 25°C)

Time Sampled: _____ pH _____ (E.C.) _____ (in micromhos/cm, corrected to 25°C)

Meter No.: Digital pH Meter #609 standardized to pH 7.0Temperature: 15 °C

(a) Field measurements and sample collection made in accordance with Dames & Moore Technical Memorandum for the pre-operational ground water.

(b) A minimum of two (2) casing volumes of water shall be pumped from the well prior to sample collection.

GROUND WATER QUALITY SAMPLING FORMWell No.: ATP-1-D-81Owner: Atlas MineralsDate/Time 1/19/82 / 1:05 Sampled By: Dale Edwards (Atlas Minerals)Appearance: Clear
(clear, colored, turbid, sediment, etc.)Remarks: 1,500 gallons pumped from the well on 1-18-82Field Measurements and Sample Collection (a)Static Water Level: 14.95 (18.40 from TOC) (in feet below land surface)Pumping Rate: 2.5 gpm (in gallons per minute)Duration of Pumping: from 10:15 to 1:05, Total Time = 110 (in minutes)Volume of Water Pumped Prior to Sampling^(b): 425 (in gallons)Method of Pumping: Centrifugal pumpElectrical Conductivity (E.C.) MeasurementsMeter No.: YSI #1Time Sampled: 10:30 pH 7.6 (E.C.) 150,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 11:15 pH 7.6 (E.C.) 130,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 12:00 pH 7.6 (E.C.) 130,000 (in micromhos/cm, corrected to 25°C)Time Sampled: 12:30 pH 7.6 (E.C.) 150,000 (in micromhos/cm, corrected to 25°C)

Time Sampled: _____ pH _____ (E.C.) _____ (in micromhos/cm, corrected to 25°C)

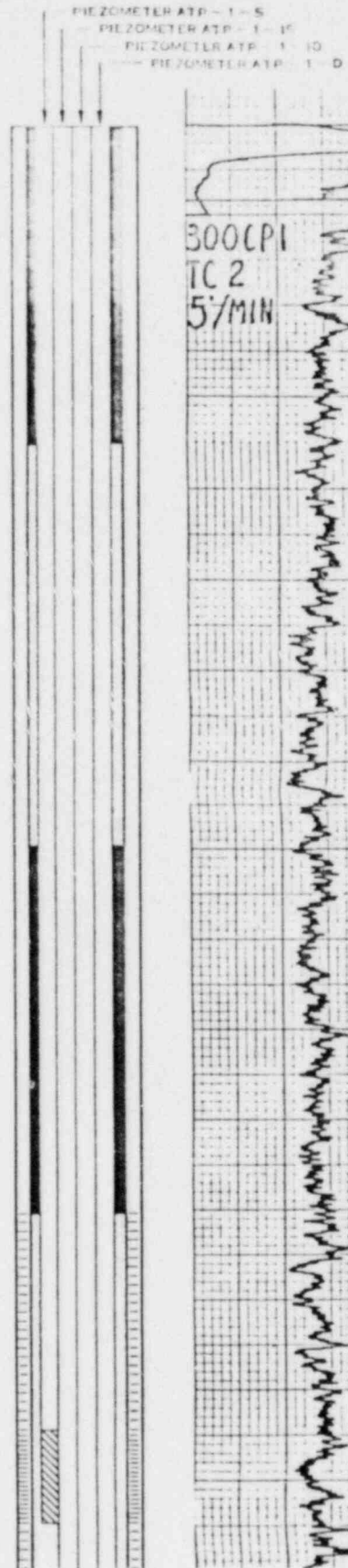
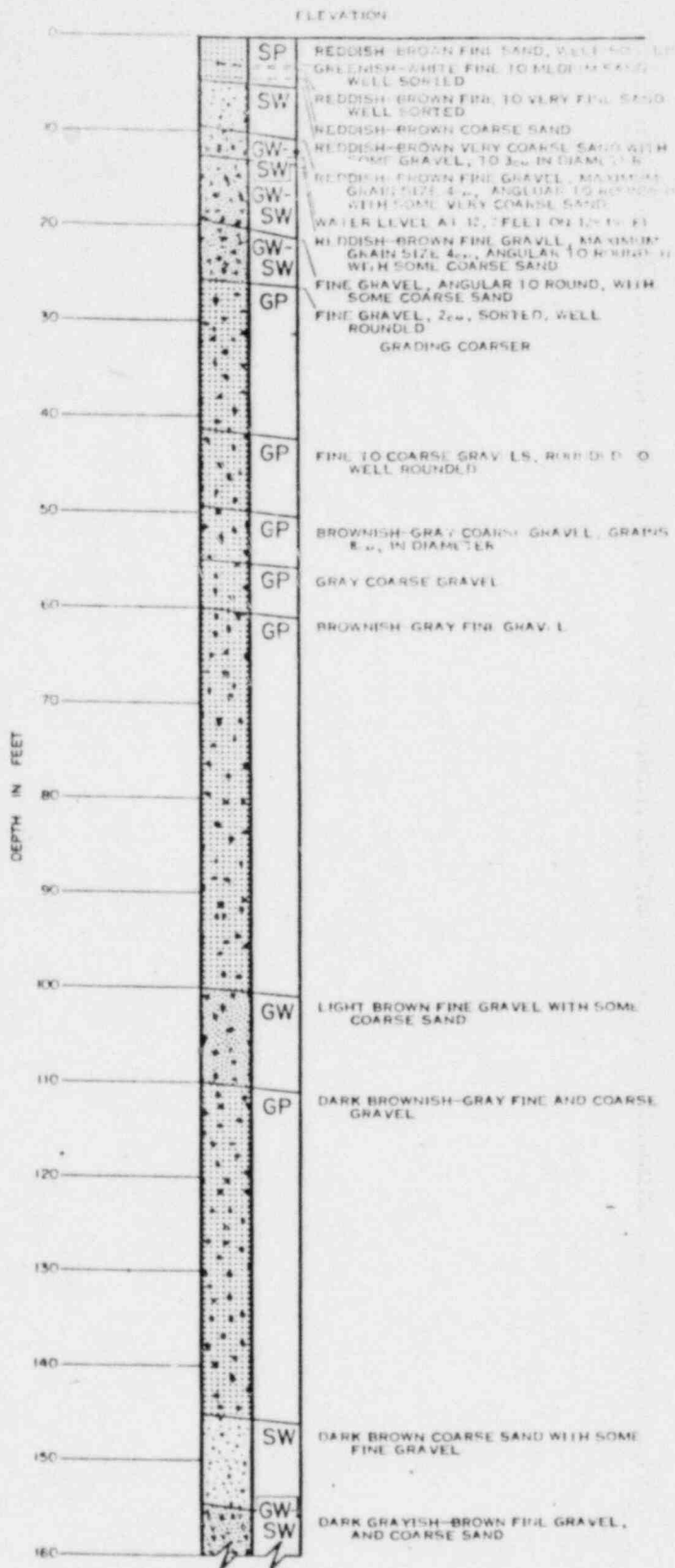
Time Sampled: _____ pH _____ (E.C.) _____ (in micromhos/cm, corrected to 25°C)

Meter No.: Digital pH Meter #609 standardized to pH 10Temperature: Buffer solution
18 °C

(a) Field measurements and sample collection made in accordance with James & Moore Technical Memorandum for the pre-operational ground water.

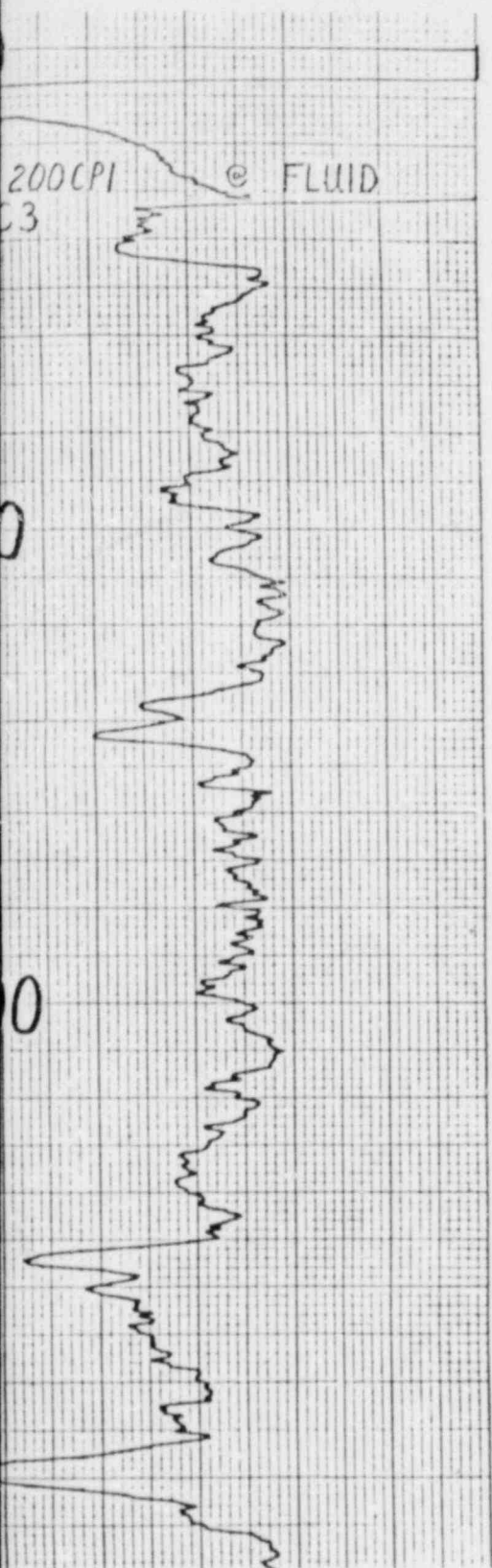
(b) A minimum of two (2) casing volumes of water shall be pumped from the well prior to sample collection.

BORING ATP-1-82

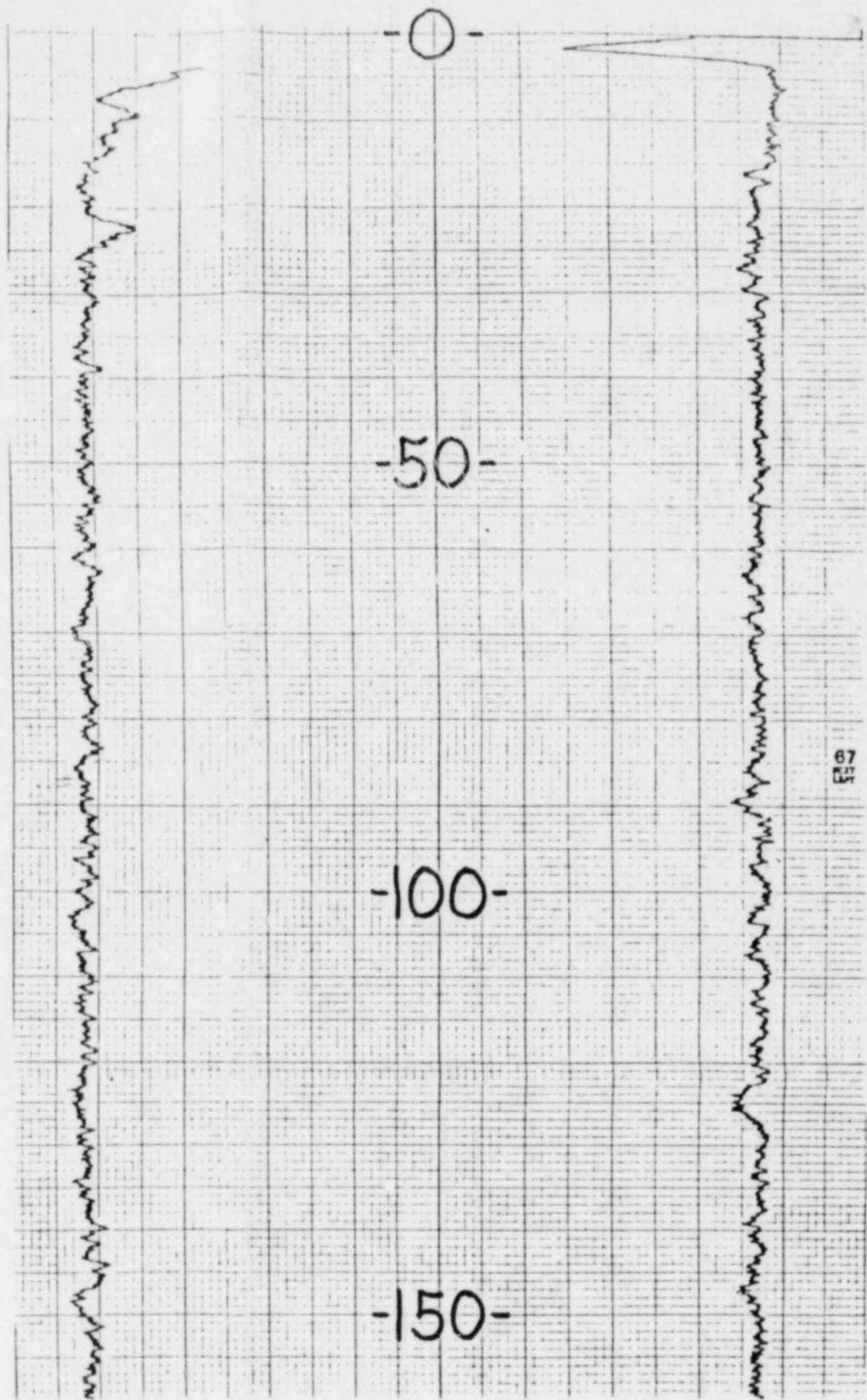


WELL COMPLETION LOG

LOG OF BORING



NEUTRON-NEUTRON LOG

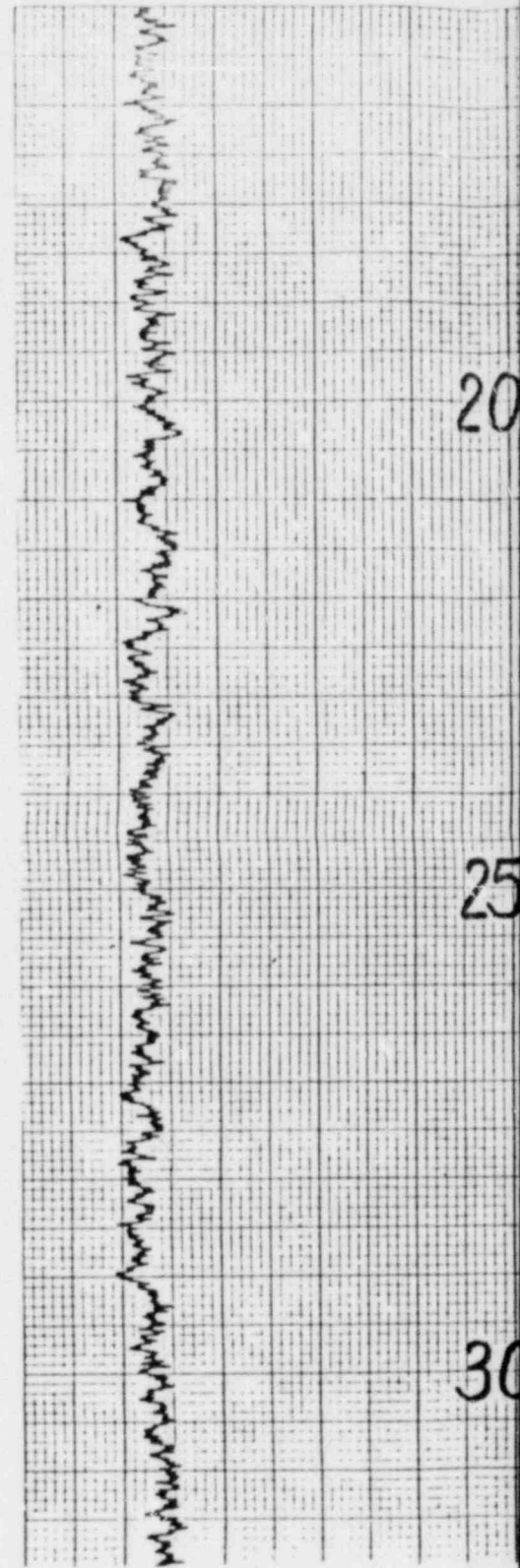
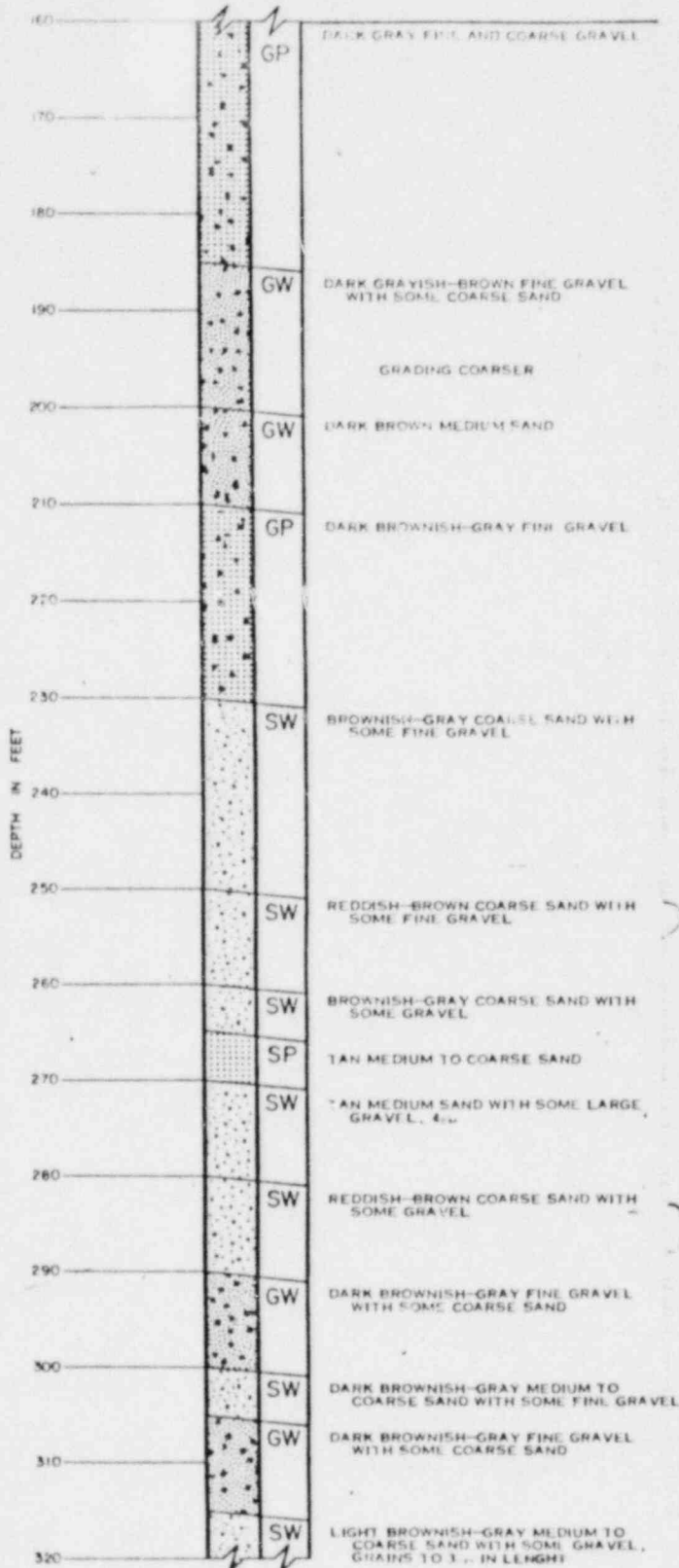


GAMMA RAY LOG

GAMMA-GAMMA DENSITY

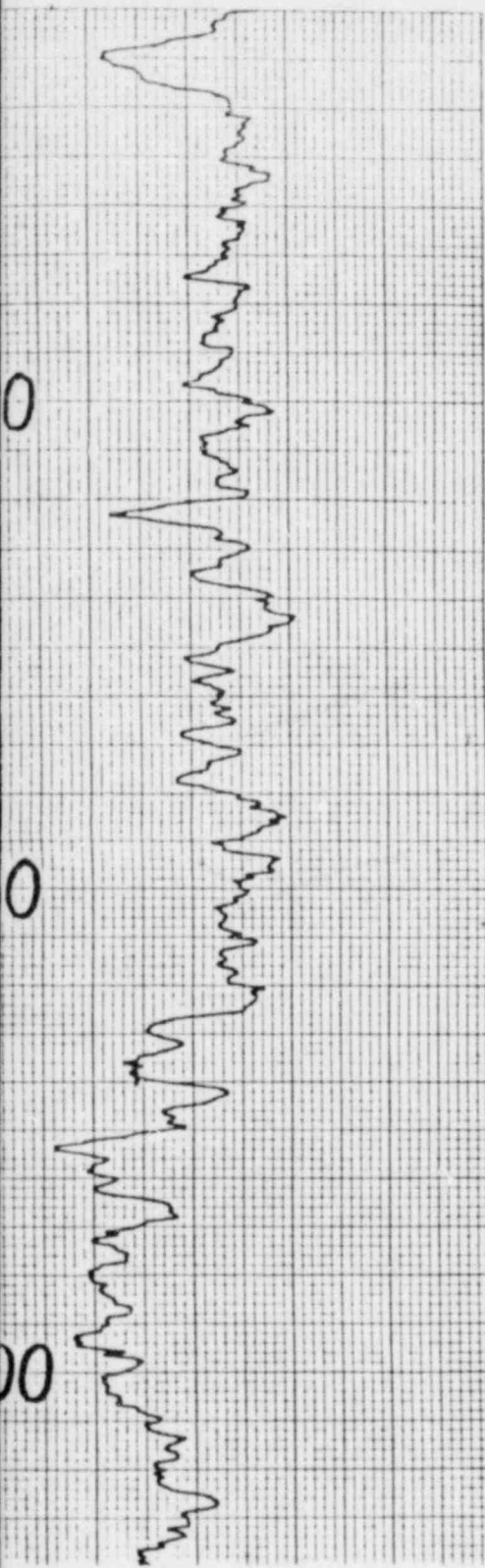
GEOPHYSICAL LOGS

BORING ATP-I-82 (CONTINUED)

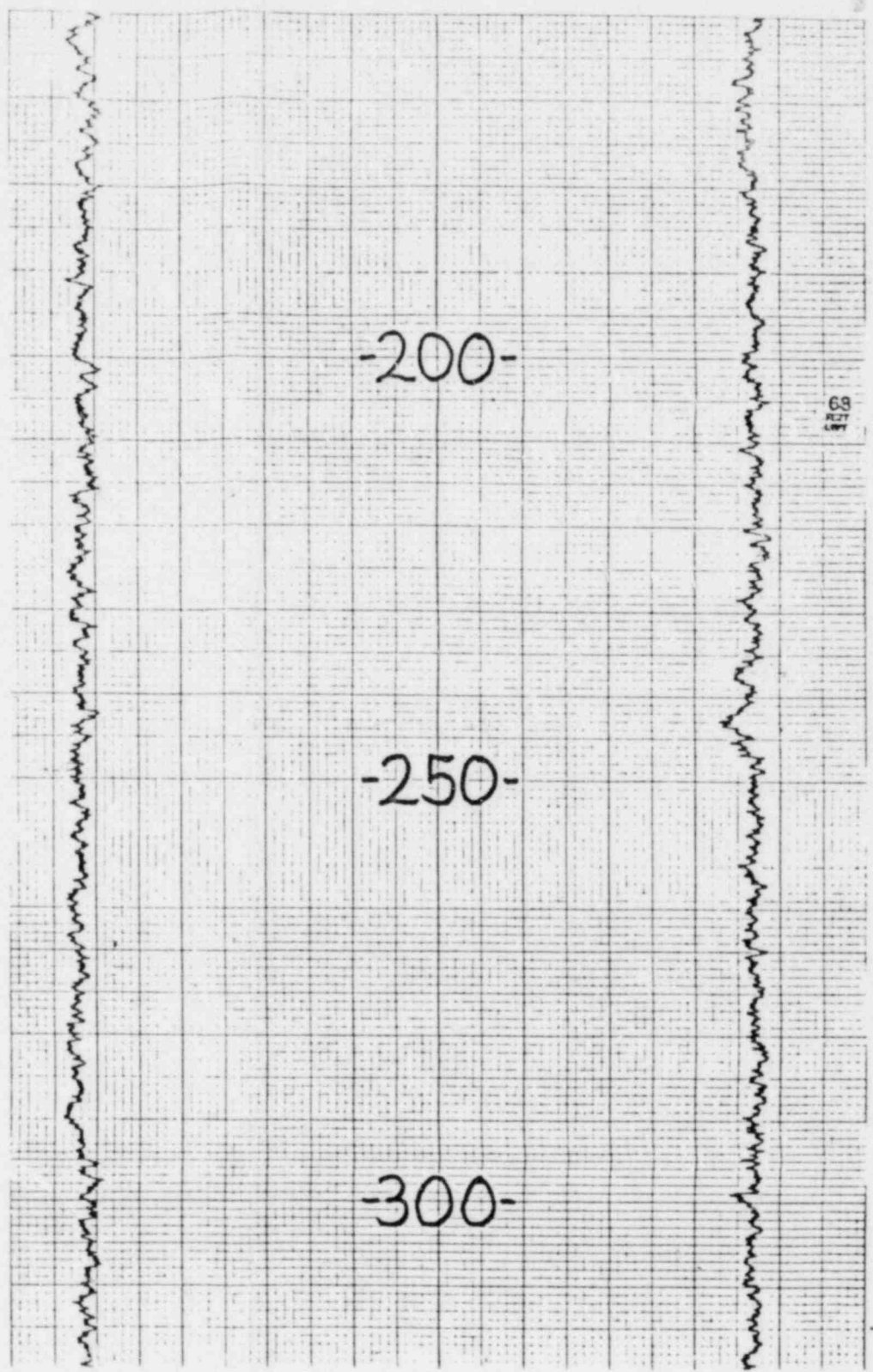


LOG OF BORING

WELL COMPLETION LOG



NEUTRON-NEUTRON LOG

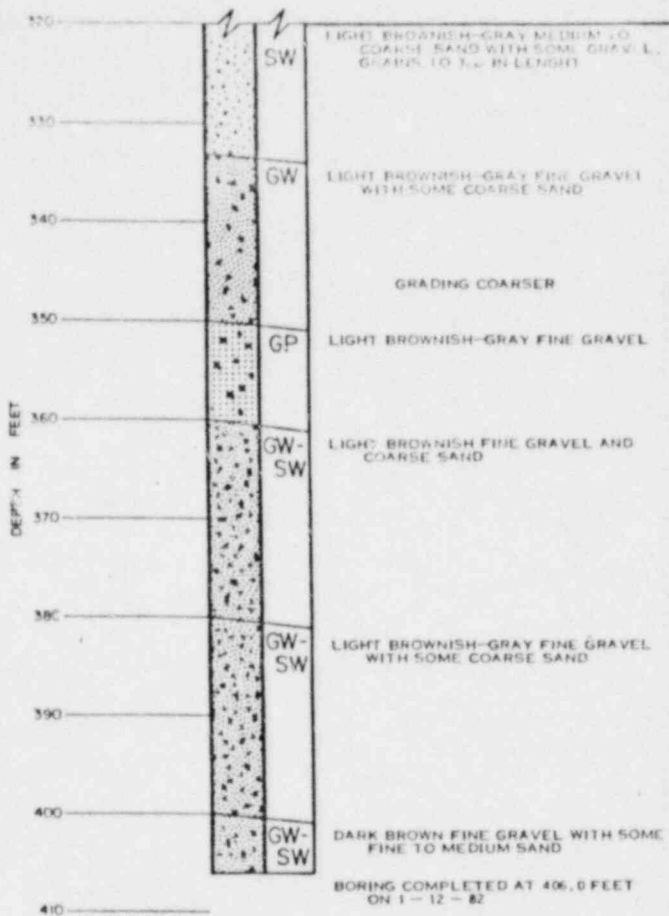


GAMMA RAY LOG

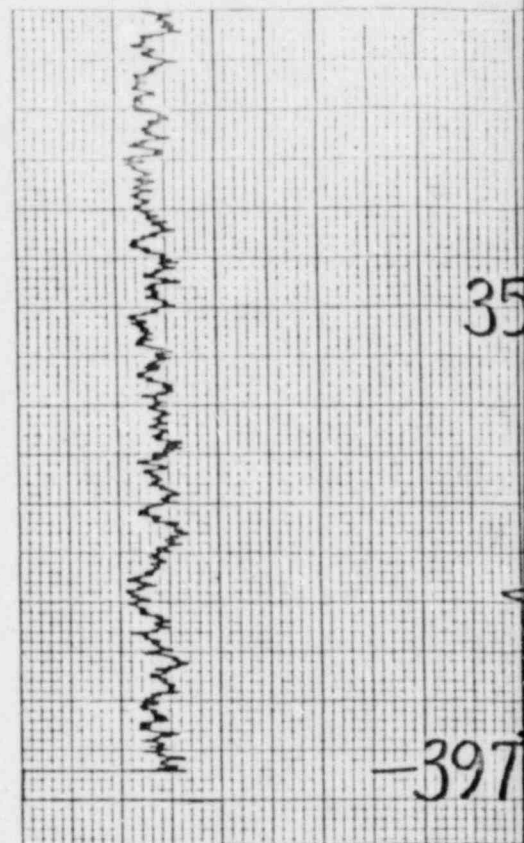
GAMMA-GAMMA DENSITY

GEOPHYSICAL LOGS

BORING ATP-1-82 (CONTINUED)



WELL COMPLETION LOG



GAMMA RAY LOG

LOG OF BORING

KEY :

- 8 1/2 INCH STEEL CASING
- 8 1/2 INCH STEEL CASING WITH 2 PERFORATIONS PER FOOT
- 8 1/2 INCH STEEL CASING WITH 4 PERFORATIONS PER FOOT
- 2 INCH SCHEDULE 40 PVC CASING
- 2 INCH SCHEDULE 40 PVC SLOTTED SCREEN
- GROUT SEAL
- SAND PACK

PLS PROFESSIONAL LOGGING SERVICES, INC.
Clifton, Colorado 81520

HOLE NO.

ATP-1-82

CLIENT: ATLAS MINERALS
AREA: MILL
COUNTY: GRAND
SECTION:

DATE: 01-15

STATE: UTAH
RANGE:

ELEVATION:

DEPTH REFERENCE: XTAL @ GROUND LEVEL

GEOLOGIST:

PHONE NO:

1809

INITIAL PLAN:

K FACTOR:

1.19 x 10⁻⁵

TO DRILLED:

406

WATER FACTOR:

1.15

TO LOGGED:

397

DETECTOR TYPE AND SIZE:

XTAL VAX 1 3/8

NATURAL GAMMA RANGE:

25 CPS / IN

CALIBRATION:

DOE GJO

TC:

2

RATE:

20 / MIN

SOURCE:

NA

SP RANGE:

NA

RES RANGE:

NA

OTHER:

BOTTOM @ 405'

Y-000:

GAMMA

RANGE:

300 CPS / IN

TC:

2

FROM:

10'

TO:

0'

TOTAL:

10'

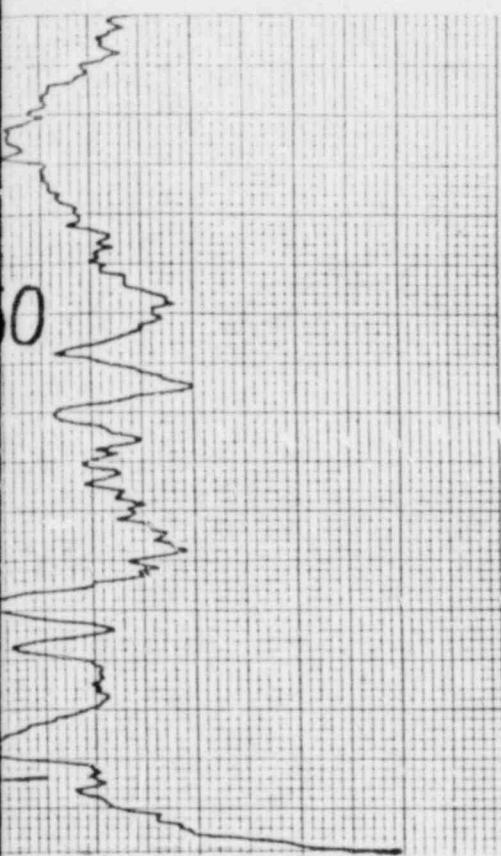
TAPE:

01-25-04A

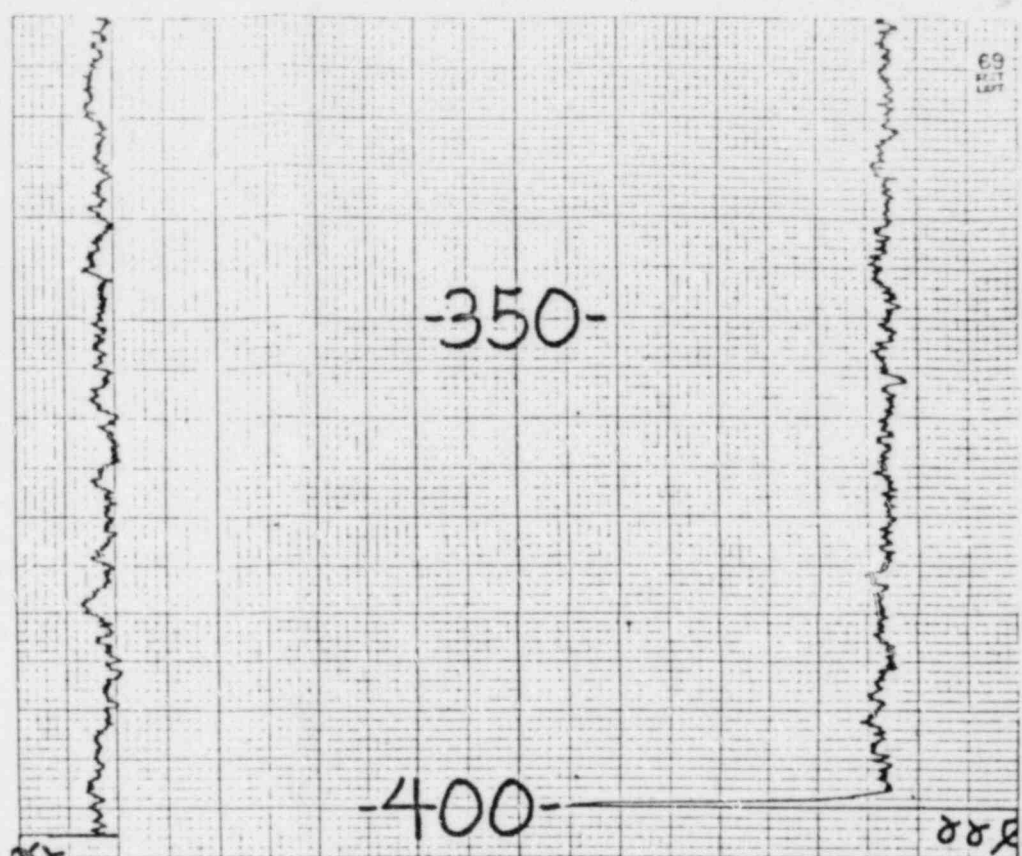
TOTAL FOOTAGE LOG:

416

11-30-81



NEUTRON-NEUTRON LOG



GAMMA RAY LOG

GEOPHYSICAL LOGS

ACRYLIC
DEN 1.2

GM/CC

MG GM/CC
DEN 2.1

AI GM/CC
DEN 2.7

DRILLING CONTRACTOR
ZIMMERMAN
I.D. UNITS
NA
TRUE DEPTH
NA
AZIMUTH
DISTANCE
NA
BIT SIZE
8 1/2 IN
CASING
8" STEEL 5/16" THK
CASING FACTOR
1.67
BOREHOLE FLUID
H₂O - BRINE
FLUID LEVEL
16'
VISCOSITY
DENSITY
ROUND TRIP MILEAGE

OPERATOR
JIM GENTRY
UNIT
25
LOCATION
MOAB
DRIVING HOURS
STANDBY HOURS
TIME IN
TIME OUT
LOGGING HOURS
TOTAL HOURS
ROUND TRIP MILEAGE

PLS PROFESSIONAL LOGGING SERVICES, INC.
Clifton, Colorado 81520

HOLE NO
ATP-1-82
CLIENT
ATLAS
AREA
ATLAS TAILINGS POND
COUNTY
GRAND
SECTION
TOWNSHIP
ELEVATION
DEPTH REFERENCE
88 XTL @ GL
STATE
UTAH
RANGE
GEOLOGIST
DATE
01-13-82

DRILLING CONTRACTOR
ZIMMERMAN
I.D. UNITS
NA
TRUE DEPTH
NA
AZIMUTH
DISTANCE
NA
BIT SIZE
8.5"
CASING
8" STEEL 5/16"
CASING FACTOR
1.67
BOREHOLE FLUID
H₂O . BRINE
FLUID LEVEL
16'
VISCOSITY
DENSITY
ROUND TRIP MILEAGE

OPERATOR
R. LARSON
UNIT
22
LOCATION
MOAB
DRIVING HOURS
STANDBY HOURS
TIME IN
TIME OUT
LOGGING HOURS
TOTAL HOURS
ROUND TRIP MILEAGE

LOG
HANDLE
NEUTRON
RANGE
100 CPS/IN
TC
3
FROM
404
TO
16
TOTAL
TAPE

LOG
HANDLE
NEUTRON
RANGE
200 CPS/IN
TC
3
FROM
16
TO
7
TOTAL
TAPE

PROF NO
2103-2228
CALIBRATION
1.49 x 10⁻³
WATER FACTOR
1.15
DETECTOR TYPE AND SIZE
XTL 2x2x2
CALIBRATION SOURCE
80 Den. 200 CPS/IN
SERIES
B319 20mCA 2nd
TEST BIT DATA
DOE G.J.C.

INITIAL RUN
TO 406
TO 400
NATURAL GAMMA RANGE
25 CPS/IN
RANGE
2
RATE
15 FPM
SPRANGE
NA
RESRANGE
NA
OTHER
BOTTOM OF PROBE @ 405

LOG
HANDLE
88 DENSITY
RANGE
200 CPS/IN
TC
2
FROM
400'
TO
0'
TOTAL
400'
TAPE
01-25-04A
TOTAL FOOTAGE LOGGED
400

LOG
HANDLE
RANGE
TC
FROM
TO
TOTAL
TAPE

MAJOR DIVISIONS			GRAPH SYMBOL	LETTER SYMBOL	TYPICAL DESCRIPTIONS	
COARSE GRAINED SOILS	GRAVEL AND GRAVELLY SOILS	CLEAN GRAVELS (LITTLE OR NO FINES)		GW	WELL-GRADED GRAVELS, GRAVEL-SAND MIXTURES, LITTLE OR NO FINES	
				GP	POORLY-GRADED GRAVELS, GRAVEL-SAND MIXTURES, LITTLE OR NO FINES	
		MORE THAN 50% OF COARSE FRACTION RETAINED ON NO. 4 SIEVE	GRAVELS WITH FINES (APPRECIABLE AMOUNT OF FINES)		GM	SILTY GRAVELS, GRAVEL-SAND-SILT MIXTURES
					GC	CLAYEY GRAVELS, GRAVEL-SAND-CLAY MIXTURES
	SAND AND SANDY SOILS	CLEAN SAND (LITTLE OR NO FINES)		SW	WELL-GRADED SANDS, GRAVELLY SANDS, LITTLE OR NO FINES	
				SP	POORLY-GRADED SANDS, GRAVELLY SANDS, LITTLE OR NO FINES	
		MORE THAN 50% OF COARSE FRACTION PASSING NO. 4 SIEVE	SANDS WITH FINES (APPRECIABLE AMOUNT OF FINES)		SM	SILTY SANDS, SAND-SILT MIXTURES
					SC	CLAYEY SANDS, SAND-CLAY MIXTURES
FINE GRAINED SOILS	SILTS AND CLAYS	LIQUID LIMIT LESS THAN 50		ML	INORGANIC SILTS AND VERY FINE SANDS, ROCK FLOUR, SILTY OR CLAYEY FINE SANDS OR CLAYEY SILTS WITH SLIGHT PLASTICITY	
				CL	INORGANIC CLAYS OF LOW TO MEDIUM PLASTICITY, GRAVELLY CLAYS, SANDY CLAYS, SILTY CLAYS, LEAN CLAYS	
				OL	ORGANIC SILTS AND ORGANIC SILTY CLAYS OF LOW PLASTICITY	
	SILTS AND CLAYS	LIQUID LIMIT GREATER THAN 50		MH	INORGANIC SILTS, MICACEOUS OR DIATOMACEOUS FINE SAND OR SILTY SOILS	
				CH	INORGANIC CLAYS OF HIGH PLASTICITY, FAT CLAYS	
				OH	ORGANIC CLAYS OF MEDIUM TO HIGH PLASTICITY, ORGANIC SILTS	
HIGHLY ORGANIC SOILS				PT	PEAT, HUMUS, SWAMP SOILS WITH HIGH ORGANIC CONTENTS	

NOTE: DUAL SYMBOLS ARE USED TO INDICATE BORDERLINE SOIL CLASSIFICATIONS.

SOIL CLASSIFICATION CHART

UNIFIED SOIL CLASSIFICATION SYSTEM

13 L C