

60002481

DISTRICT	Manhattan - see back
DIST_NO	2960 - see back
COUNTY	Nyc - see back
If different from written on document	
TITLE	Hughes Series Employees and Consultant Reports - Skinner, Paul
If not obvious	
AUTHOR	Skinner P; Block, F
DATE OF DOC(S)	1974
MULTI_DIST <input checked="" type="checkbox"/> N?	See back
Additional Dist_Nos:	
QUAD_NAME	Manhattan 7½' - see back
P_M_C_NAME (mine, claim & company names)	Hughes Tool Co.
COMMODITY	Gold silver
If not obvious	
NOTES	Correspondence; handwritten notes; assay procedures; expenses
NOTE: Some pages double sided Scan Folder front	
58p.	

Keep docs at about 250 pages if no oversized maps attached
(for every 1 oversized page (>11x17) with text reduce
the amount of pages by ~25)

Revised: 1/22/08

SS:	DD	12/22/08
Initials	Date	
DB:		
Initials	Date	
SCANNED:	MT	1/12/10
Initials	Date	

District	District No.	ID	Quad	PMC	County
McCoy	3020	60002472	McCoy 15'	Hughes Tool Co.	Lander
Silver Peak	4390	60002484	Silver Peak 7 $\frac{1}{2}$	Hughes Tool Co. Mary Mine	Esmeralda
Tonopah	4840	60002485	Tonopah 7 $\frac{1}{2}$	Hughes Tool Co.	Nye

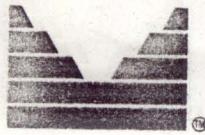
SKINNER, Paul

25

60002471 2960

HUGHES SERIES
EMPLOYEES & CONSULTANT REPORTS
SKINNER, PAUL

25



Land Exploration
and Mining Division

66662471 2960
Post Office Box 1126
Tonopah Nevada 89049

A Division of
Summa Corporation

#3-N

October 29, 1974

Mr. Frank E. Block- Research Director
U.S. DEPARTMENT OF THE INTERIOR
BUREAU OF MINES
1605 Evans Ave.
Reno, Nevada 89505

Dear Mr. Block:

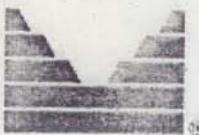
In answer to your letter of October 7, 1974 I would like to commend you on your Open House.

I received some very important information about carbon stripping with ethanol. In fact I believe we are going to use this process in our operation.

I am very pleased to see that the Bureau of Mines is now spending it's time and money on things that are practical. Your rare earth project of past was interesting and informative, but didn't really do anything for mining, where as your more recent work is of a great help to people in the industry. I would be most interested in anymore such demonstrations you have in the future.

Sincerely,

Paul M. Skinner
Chief Metallurgist
Summa Corporation



Land Exploration
and Mining Division

Post Office Box 309
Las Vegas Nevada 89101
702 739 7441

A Division of
Summa Corporation

12-8

November 25, 1974

National Filter Media Corp.
333 S. 3rd East Street
P. O. Box 156
Salt Lake City, Utah

Dear Sirs:

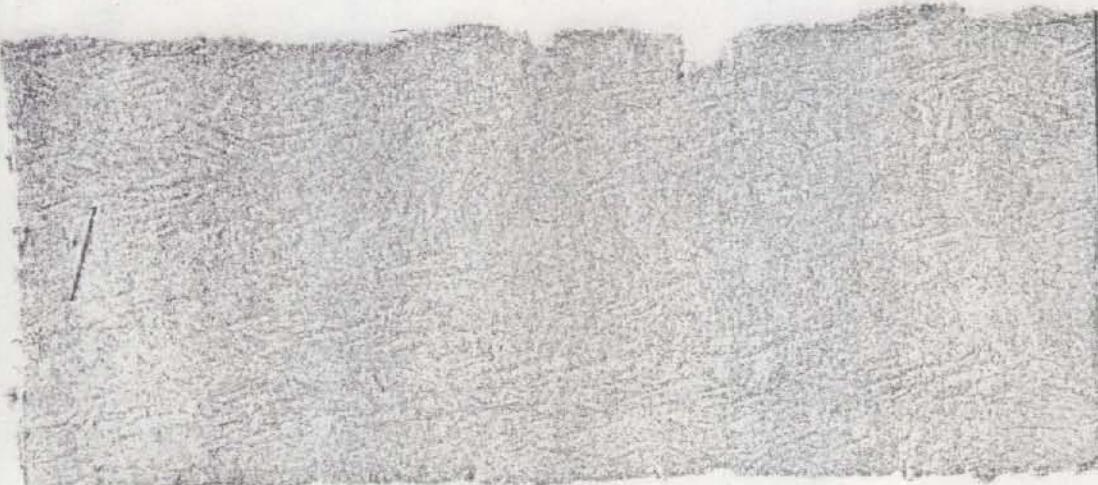
I am interested in purchasing some canvas duck filter cloth and filter paper from you. Enclosed is a sample of the type of paper I would like. The canvas I want is canvas duck suitable for filtering silver and gold precipitate out of cyanide solution. Could you send me prices and a brochure on your products.

Thank you,

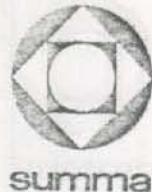
Paul M. Skinner

Paul M. Skinner
Chief Metallurgist

PMS:sfm
Enclosure



Internal Communication



Date: November 26, 1974
To: Dave J. Gribbin
From: Paul M. Skinner, Chief Metallurgist
Subject: Preliminary Report on the Mary Mine Leach No.1

A 500-pound sample of mine run Mary ore was placed in a tub and leaching started. The sodium cyanide was kept at 2.0 pound per ton of solution and a pH of 10.5 was maintained with caustic soda.

After three days the pregnant solution had built up to 7.2 ppm., at which time I put in a carbon column to continuously strip the pregnant solution. After 19 days of leaching the pregnant and the barren solutions were both the same, .008 ppm. I then took the carbon column off and circulated the solution for 2 more days and found that there was no more build-up of gold in the solution. I therefore concluded that the leach was completed.

The sample was then washed with water and a 100-pound sample was split out and assayed.

The assays were as follows:

head .128 oz. Au
tail .031 oz. Au

$$\frac{.031}{.128} \times 100 = 24.2\% = 75.8\% \text{ total gold recovery}$$

head .280 oz. Ag
tail .032 oz. Ag

$$\frac{.032}{.280} \times 100 = 11.4\% = 88.6\% \text{ total recovery}$$

November 26, 1974

Page 2

Conclusions:

The results of this preliminary test show that the Mary ore is amenable to heap leaching. A recovery of 75% on mine run ore is excellent.

I am, however, in the process of running a test on material which has been crushed to -3/8 inch to see if an even better recovery can be gotten.

I am very pleased with these preliminary results and I feel that the Mary ore can be leached at a good profit.

Paul M. Skinner

Paul M. Skinner
Chief Metallurgist

Dist: PMK
WS
Lute
Marymine file
WOM - 7-14-75

BROWNSTONE Mining Co.

P. O. BOX 215 • LONE PINE, CALIFORNIA • PHONE: 875-4640
P. O. BOX 206 • DARWIN, CALIFORNIA • PHONE: 876-3565

January 12, 1975

Statement - Summa Corp, Tonopah, Nevada

The following is my bill for consulting at your Tonopah and Manhatten, Nevada projects.

December 30, January 2 and 3 - 3 days at \$125.00 per day

January 6 through January 10 - 5 days at \$125.00 per day

Total due \$1000.00

Make check payable to Paul M. Skinner

Paul M. Skinner

Paul M. Skinner



Internal Communication

Date: January 8, 1975
To: D. J. Gribbin
From: Paul Skinner
Subject: Amalgamation Run on 75 Pounds of Concentrates

A small amalgamation barrel and amalgamation plate were borrowed from Jim Larson. This barrel has a capacity of about 80 lbs. of concentrates.

The test run went as follows:

75 lbs. of concentrates (assay unknown) were added to the barrel along with 10 ball mill balls, one handful of lime, 2 lbs. of mercury and water to within 2 inches of the top. The lid was put in place and the barrel turned on for 45 minutes. While the barrel was rotating the copper amalgamation plate was dressed down with new mercury and some silver to get it prepared. The barrel's contents were emptied into a bucket and the rock washed away from the mercury with a strong stream of water. This was done over the amalgamation plate so that any escaping amalgam would be caught on the plate. After cleaning, the remaining mercury and amalgam were squeezed through a muslin cloth to obtain the amalgam.

All in all the test went well, but we did have a longer than normal mercury loss because of all the heavy black sands. This was strictly a physical loss and not chemical. What happens is that when cleaning the amalgam the heavy black sands make you use a stronger stream of water than should be needed. This in turn causes more mercury than normal to overflow the bucket onto the plate. When the mercury hits the plate it runs along the top of the heavy, dense black sands and is insulated from the plate, keeping it from sticking, and in turn causes the mercury to run into the tailings.

Paul Skinner to D. J. Gribbin

January 8, 1975

Amalgamation Run on 75 Pounds of Concentrates -- page 2

This problem can of course be remedied by doing what I did in the lab experiment -- remove the black sands first by magnetic separation, then amalgamate.

Paul M. Skinner

Paul Skinner



Internal Communication

Date: January 8, 1974
 To: D. J. Gribbin
 From: Paul Skinner
 Subject: Placer Amalgamation (Test)

P. S. 126
 Screen Analysis 254.6 g.

Screen Size	q.	% Dist.	mg.- Au - Ag	Au
+10	12.4	4.8	62.44	= 43.71
-10+20	80.0	31.4	873.77	611.64
-20+30	39.0	15.3	444.96	311.47
-30+48	60.8	23.9		
-48+80	33.2	13.0	-30 through -100	
-80+100	5.4	2.2	928.73 by amalgamation	
-100	23.8	9.4		

The -30 mesh sample was amalgamated for 10 minutes with Hg. The amalgam was then taken down with hot HNO_3 . The metallics were then fired and weighted. The tailings from the amalgamation assayed 6.10 oz. Au and 3.96 oz. Ag per ton.

$$\text{mg.Au in tails} = \frac{252 \text{ q.} \times 6.10 \text{ mg.}}{29.166}$$

$$= 52.7 \text{ mg. Au in the tails}$$

Paul M. Skinner

Paul Skinner to D. J. Gribbin
January 8, 1975
Placer Amalgamation (Test) - page 2

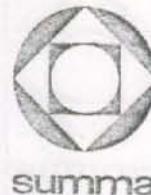
43.71
611.64
311.47
928.73
52.70
1948.25 mg. Au total in the sample
1895.55 mg. Au taken out = 97.3% recovery by amalgamation.
$$x = \frac{454 \times 1.948}{254.6} = 3.47 \text{ g. per lb. of sample}$$
$$= 6953.7 \text{ g. per ton}$$
$$= 223.57 \text{ oz. per ton}$$
(calculated head assay)

Conclusions:

This test shows that by simple amalgamation 97.8% of the gold can be recovered from the Manhattan placer concentrates. I feel that this is the route we should take in cleaning up these concentrates. A small, 2 or 3-foot amalgamation barrel should work real well.

Paul M. Skinner

Internal Communication



Date: January 8, 1975
 To: D. J. Gribbin
 From: Paul Skinner
 Subject: Placer Magnetic Separation Test

P.S. 131

100 g. Manhattan placer concentrate was taken and separated with a magnet with the following results:

		Au oz./t	gm.oz.	%Dist.	%Dist.Au
Head	100 g.	4.015	401.5	100	100
Magnetic	79.5 g.	.048	3.816	79.5	.95
Non-magnetic	<u>20.5</u> g.	<u>19.40</u>	<u>397.7</u>	20.5	99.05
	100.0		401.52		

$$H \times P = \frac{401.52}{100} = 4.015 \text{ oz. per ton}$$

Conclusions:

From this test it can be seen that 99% of the gold is in the non-magnetic portion of the concentrates, and the non-magnetic part constitutes 20% of the total. There was also an upgrading of approximately 5 to 1 when the magnetic fraction was removed. I would highly recommend having the placer concentrates dried and put through a magnetic separator before amalgamation. This would help in two ways: one is it would cut your need to run the amalgamation plant down to a minimum because of the reduction in volume, and a lot of heavy magnetite is a real nuisance when you amalgamate.

A handwritten signature in cursive ink that reads "Paul M. Skinner".

Date: September 17, 1975
To: WOM, DJG, WS
From: Susie Mollison *SJM*
Subject: Spectrographic Analysis "Skinner #1 03/28/75" by
USBM Reno Metallurgy Research Center
(Sample of cracked Summa Bar)

Dr. Howard Heady, Director of the Analytical Group for the above center, telephoned the following to me this afternoon. (I had spoken to his assistant this morning requesting the information.) He believed the gold sample had come to him from Harold Heinen with a request to identify contaminants.

The elements he detected in the sample are in the following percentages:

Aluminum	.01
Copper	.06
Iron	.06
Magnesium	.002
Manganese	.2
Nickel	.01
Lead	.5
Silica	.0005
Tin	.004
Zinc	less than .1
Silver	4.0
Gold	major (the remainder of the sample)
Platinum	.003

PROJECT MANHATTAN AL + Zn DUST PRECIPITATION

Notebook No. _____

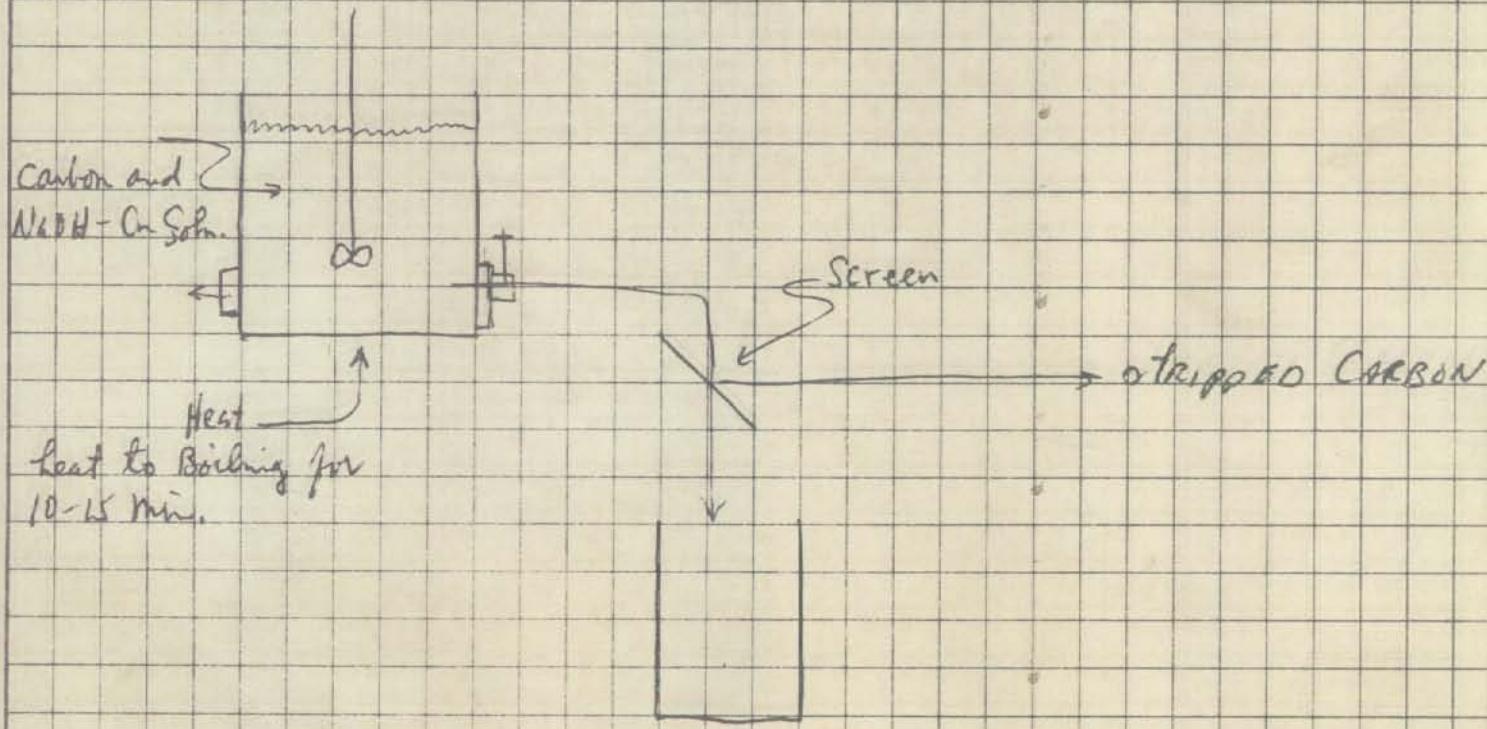
Continued From Page _____

100 ml preg. soln from carbon strip cell, touch of Pb acetate added then zinc dust. Soln. agitated for 5-10 min. Soln. filtered through filter paper. ppt. then treated with HCl to dissolve off excess zinc, then dried with Boro glass, rubic. Soda Ash. filtrate analyzed with A-A.

preg soln = 85 ppm (by A-A) 2.48 g/l for soln.

Boro soln = 0 ppm (by A-A)

ppr. wt =



Continued on Page _____

Read and Understood By

Signed _____

Date _____

Signed _____

Date _____

2
PROJECT MANHATTEN - CARBON TEST

Notebook No.

Continued From Page

1000 ml - 3% NaCN / Ton of Soil.

500 ml = PREC. CARBON = approx 450 g OF CARBON

AGITATE FOR 20 MIN. AT 200°F

160°F 12 ppm Au

180°F 65 ppm Au

200°F 103 ppm Au

200°F 130 ppm Au after 15 min.

After, precipitated carbon was then filtered. The carbon digested and the solution then analyzed. The gold-tin zinc dust was added to the soln. and allowed to stand with some stirring. The ppt. was filtered and the soln. showed no Au on the A-A machine. The ppt. was then dried and fired to get the Au-Ag.

Continued on Page

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Date

Signed

Date

1000 ml - 10% NaCN Soln.
300 g Carbon (prey.)

Heat soln. to 200°F, then add the prey carbon to the agitated
soln.

Samples

No.	RPM Ag	Time Start	Time Stop
1	5 min	56	8 to 9:00
2	10 min	61	3 to 9:02
3	15 min	—	9:02
4	20 min	—	9:07
5	30 min	62	9:02

After 15 min 10 more g NaCN added - continued boiling for 30'
total.

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Date _____

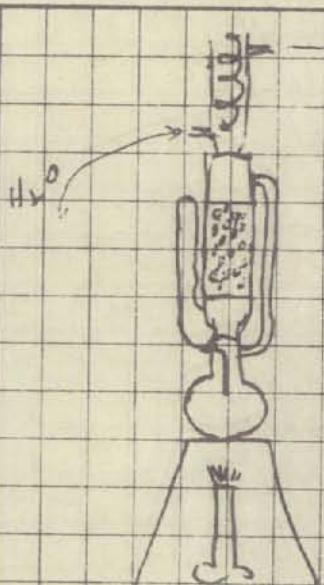
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Date _____

PROJECT SOXHLET REFLUX TEST

Notebook No. P.S. 116

Continued From Page



130 ml Soln in flask. Soln made up of
 .1% NaCl
 1.5% NaOH
 20.0% H₂O
 78.4% METHANOL = 102 ml methanol added

50 g loaded pellet Carbon

turned on at 9:00 A.M. cycle time ± 20 min
 turned off at 5:00 P.M. approx. 20 bed vol thru
 the carbon.
 temp at ± 75°C

Return methanol - drain out 45 ml

45 ml

Carbon Load I 200 oz. Au

Return - 90 ml

Carbon load II 200 oz. Au
 carbon total 8 hrs. 138.8 Au, 30.64 Ag
 " 36 hrs.

The carbon was then drained off its methanol and removed to the large flask where H₂O was added and the carbon and water mixture then heated and the methanol distilled off of the carbon. The temperature was slowly raised until a temperature of 105°C was reached to purge the system of methanol.

Buat No. 1 Soln. after Amm. Sulfide ppt. - 46.37 ppm Au, 0 ppm Ag

Buat No. 2 " zinc dusting - 42.4 Ag

fusior No. 1 ammon. sulf ppt. - .156 mg Ag, 8.134 mg Au

" No. 2 zinc dustt ppt. - .024 mg Ag, .366 mg Au

8.29 mg Soln = 1.88% Ag

.156 mg Ag

= Dore Rollin = 981.2 fine in Au

Continued on Page

Read and Understood By

11-4-74

P.M.S.

Conditions same as that P.S. but extractor run for 24 hrs.
or at 12:00 noon 11-7-74

Solution made up of 200 ml H₂O

1g NaCl

15g NaOH

800 ml Methanol

± 50 g carbon used

150 ml soln. used

Carbon load ± 200 mg Au

Carbon load 117 mg Au, 14.4 mg Ag

prey Soln at 24 hrs - 255.6 ppm Au

" " #36 hrs - 388.4 " Au, 28.76 ppm Ag

Au
P.P.M.

Boat #1	sol before zinc or hydride but after sulfide	cone	631.2
#2	sol after zinc	cone	513.4
#3	" " hydride	cone	544.16
#4	" " prey. Monett	11-13-74	.16
#5	" " Bane Monett	11-13-74	.07
#6	" " prey. Mary	11-13-74	.08
#7	allied extraction Soln. from 50g Carbon (pellet)	cone	48.06 g Au

Silver Sulfide from NH₄S precipitation .290 mg Au, 18.57 mg Ag

Fine coconut carbon to be used for methanol extraction 81.3 g Au, 17.24 g Ag

Continued on Page _____

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Date

Signed

Date

Same as Test 116, 117, 118 only ethanol was used.

5.0 g sample of pellet carbon used.
10 ml of Solvent.

Began at 3:00 P.M. 11-12-74

-111 at 11:00 p.m. 11-12-74

Carbon washed and arranged

all of the ash from the strip distilled down to 25 ml and the put a boat and taken to dryers

pure Carbon 200.0 g

pure Carbon 80.8 g Au, 17.14 g Ag

$$\frac{30.8}{200.0} = 40.4 \quad 100.0 - 40.4 = 59.6\% \text{ pure Au}$$

Continued on F

Read and Understood By

Signed

Date

Signed

Same as last P.S. 118 only pre-cooked with methanol only for 16 hr before

50 carbon

100 ml methanol stirring each.

Start extraction at 8:30 A.M. 11-13-74

1st dec at 8:49

2nd dec at 9:09 - 20 min per dec volume
- 3 per hr - 20 in 3 hrs

pp at 4:30

Carbon washed with water and dried the analyzed
soh, fine added (wt of carb + 25 mg) 1 ml of 10% Na₂S added, this
dropped out what looks to be all of the Ag₂S in nice large pieces.

pp carb = 20.0 g Au

Basis Carb = 21.06 g Au, 7.10 g Ag

$$\frac{21.06 - 10.53}{21.06} \times 100 = 49.47\% \text{ net in 8 hrs}$$

$$\begin{aligned} \text{pp. wt Na}_2\text{S added wt. of ppt.} &= 2.98 \text{ mg Au, } 49.58 \text{ mg Ag} \\ &= 52.56 \text{ total} \end{aligned}$$

$$\frac{49.58}{52.56} = 92.569 \text{ fine Ag}$$

$$\text{predicted ppt} = 52.7 \text{ mg Au, } 5.48 \text{ mg Ag} = 58.18 \text{ mg total}$$

$$\frac{5.48}{58.18} = 907.40 \text{ fine Au}$$

Basis wt after pre-cooking = 43.0 ppm Au, 0 ppm Ag

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Date

PROJECT Soxhlet Extraction - Same as 120 Except Small Carbon Continued From Page

50 g carbon 15:1, 81.3 g Au, 17.24 g Ag

100 ml methanol soln.

Condition 24 hrs

on at 8:00 A.M.

off at 4:00 P.M.

pre carbon assay 81.3 g Au, 17.24 g Ag

carbon Carbon 1.10 g Au, 2.58 g Ag

2 98.6 % extraction of Au

Continued on Pg

Read and Understood By

Signed

Date

Signed

PROJECT SOKHLET EXTRACTION PELLET 12 Hrs.

11-21-74

50g Sample pellet Carbon
 \pm 100 ml methanol extraction Soln.
 24 hr. condition time
 12 hr. run time.

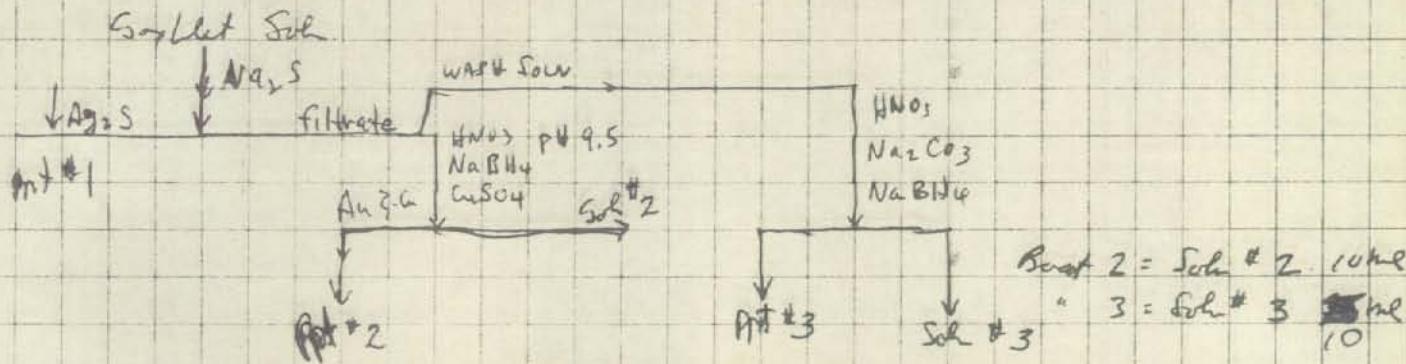
pure carbon - 200 mg Au
 barren Carbon - 14 g Au - 8.56 mg

Soln #1

Soln. from Sokhlet extract add 2 ml of 10% Na_2S to the 25 ml Soln.

part called P.S. 122 Ag2S ppt. - filtrate ^{Soln 2} pH adjusted down to 9.5 (HNO_3), Borohydride added 2 ml and wash of CuSO_4 called 1st gold part.

WASH ml from Ag2S ppt. = Soln #3, taken to acid side brought back with Na_2CO_3 , borohydride added, some lime



Boat #2 Au 13.96 mg

Continued on Page _____

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Date

Signed

Date

PROJECT Soxhlet extraction (24 hr) methanol

Notebook No. P.S. 124

Continued From Page 11-22-74

50 g Carbon

Methanol Soln to soak and cover
Condition 24 hrs.

start 7:00 A.M. 11-22-74
temp = 70°C

Soh 1 - 1/2 full soh. 5 ml Start at 50 ml 3332 pp 986 PI
Soh 1D. 2 - sohn after Na₂S (filtnet) diluted up to 250 ml, 10 ml total
5244 11.6

Boat 1A - Apt on beach

Boat 2A - Sohn after chitidy on whole thing

Hg₂S ppt more 37.8 mg, Au 1.74 mg, Ag 36.156 mg

Bare Carbon 31.65 g Au, 2.94 g Ag

load -
Total -

Continued on Page

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Signed

Date

Signed

Date

62 # of many ore through lab

6200 ml of 50%, add 10 g NaOH to start
2000 " " " more

\$200 more of sales added Total

466v hue of Soln back after saturation

~~3,600~~ me' of ore to saturate 62% of ore

9.78 NaCN added to start, NaCN put in at 5:15 A.M. 11-21-74
Lead assay - .070 Au, .05 Ag

Sample out for array as of 11-25

22-40 ne

23 - 40 me

24-40 mc

21-40 ml

Total mg Au in Sample = 67.49 mg
1st Soln return 3500 ml = 21.56 mg Au
2nd Soln return 5000 ml = 15.60 mg Au
3rd Soln return 3700 ml = 5.40 mg Au
42.56

$$\frac{42,56}{67,49} \times 100 = \underline{\underline{63,06\%}}$$

	#/T NaCl	pH	Am ppm	Bg ppm	Gloweto ml/min	Added NaCN	Added NaCN
11-22-74	1.8	10.8	2.96	4.88		14.2	14.0
11-23	1.8	10.6	10.08	0		—	—
11-24	2.0	10.6	8.96	0			
11-25	2.0	10.6	4.32	24.88	100		
11-25	AFTERNOON (2:30) LANT ASSAY		6.16	0	Complete change of soln.		
	New Soln 4.0#/T NaCN, pH 11.5 (4000 ml)						
11-26-74	2.1	11.5	3.12	5.44	added new soln	2.0	3.0
11-29-74	2.1	11.0	3.12	1.68			
12-1-74	2.2	11.0	2.64	0			
12-2-74	2.1	11.0	3.04	4.96			
12-3-74	n		3.12	12.08	11 at 8:00 A.M. & 5004 ml soln disp.		
12-4-74	New Soln 2.0#/T NaCN, pH 11.0 (4000 ml)				on at 7:00 A.M. 12-4-74		
12-5-74	On for good 3:00 P.M.						
	Soln return	3700 ml	1.44	0			

Continued on Page

Read and Understood By

Report Made

Signed

Date _____

Signed

D

PROJECT Placer Conc. (manhattan)

254.6 g placer conc.

screen analysis

		Mt. %	g. Au
+ 10	12.4	4.8	62.44 = 43.71
-10 + 20	80.0	31.4	1873.77 = 611.64
-20 + 30	39.0	15.3	444.96 = 311.47
-30 + 48	60.8	23.9	
-48 + 80	33.2	13.0	928.73 by amalgamation
-80 + 100	5.4	2.2	
-100	23.8	9.4	

Took sample and amalgamated for + 5 min. Mercury then taken off and dissolved with HNO_3 . The amalgam was then fired with flux and weighed. The sample taken for assay.

Assay 6.10 g/tion 3.96 g/tion = 10.06 Au-Ag alloy

$$\frac{10.06}{6.10} \text{ mg} = \frac{x}{252.9 \text{ (raw)}} \quad x = 52.7 \text{ mg Au in the tons}$$

$$\begin{array}{r}
 62.44 \quad 43.71 \\
 1873.77 \quad 611.64 \\
 444.96 \quad 311.47 \\
 928.73 \quad 928.73 \quad 1895.55 \\
 \hline
 52.70 \quad 52.70
 \end{array}$$

my Total gold : 97.77 % recovery by amalgamation

$$\begin{array}{r}
 1.948 \\
 2.368 - 2 \\
 \hline
 254.6 g \quad 454
 \end{array}
 \quad x = \frac{454 \times 3.96}{254.6} = 1.948 \quad 3.47 \\
 = 4.20 \text{ g/lb.} \\
 = 8.953 \text{ g/tion} \\
 = 270.6 \text{ g/tion Calculated lead} \\
 223.57$$

Continued on Page

Report Made

Read and Understood By

Signed

Date

Signed

Date

50g sample conditioned over night (large pellet carbon) by pumping 70°C water through the carbon sample then put into the extractor for 24 hr at 800 A.M. 12-4-74

Boat #1 left over conditioning soln. (25 ml) 7.05 ppm
Boat #2 soln. after prec. with aluminum shavings (25 ml)

Continued on P

Read and Understood By

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50 g sample conditioned with methanol only for 13 hrs., then run through the extractor for 8 hrs.

Carbon assay (pres.) 2700 g
Carbon assay (stripped)

gold soln. from above precipitated with Na_2S and filtered.
precipitate put into beaker electrolytic cell and power turned on.
after 30 min. A-A showed no Au in soln. 25 ml taken for assay.

Boat

BOOZE -

MASH - 5 GAL H_2O
9# SUGAR
1 CUP YEAST } LET WORK UNTIL SOUR, 7-14 DAYS AT 70°F
IF NOT SOUR ADD H_2O

$$5 \text{ lb soln or 5 ml} \quad \frac{4.0}{200} = \frac{y}{5000}$$

Continued on Page _____

Read and Understood By _____

PROJECT Man Mine - 1/8" alluvial ore (well water)

Continued From Page

Lead - .078

Well H₂O - 6.8 pH

50# ore crushed to finest setting on crusher, sample put into column and 2 gal H₂O run into sample

$$2 \text{ gal} = \times 3.785 = 7.57 \text{ liter H}_2\text{O in } 5 \text{ pts or } .9463 = 4.73 \text{ liter return}$$

$$\therefore 7.57 - 4.73 = 2.84 \text{ liter H}_2\text{O feed by 50# of ore. on 2:00 12-17-74}$$

10g NaOH added, when then pumped through the column until basic soln. returned, 5g more NaOH, 10g more, +5 more +10

added 10g NaCl - titrate after 2 hrs. = .91% T, added 10 mole NaCl

DATE	#/T NaCl	pH	Ag ppm	flow rate	added NaCl	NaOH
12-17					20g	40g
12-18	.45	9.3	1.80	0	10g	10g
12-19	2.35	11.0	2.88	-	-	-
12-23		TR				
12-20		2.72				15g

12-26 Replaced all of the soln. with fresh soda, 5#/ton soln., one gallon - started at 10:00 A.M.

12-26 3.52

Lead .078

total .012

$$\frac{.078}{.012} \times 100 =$$

$$\frac{.012}{.078} = 15.4 = \underline{\underline{84.6}}$$

1 ton ore will hold $\frac{2000 \times 2.84}{50} = 113.6 \text{ l/ton} = 113.6 \times 2642 = 30 \text{ gal soda}$

Report of
Information given to Bob Sutzy about mid June

Continued on Page

Read and Understood By

Signed

Date

Signed

Date

Screen Analysis of Head Sample

	Grams	%	Cumulative %	Au	$\frac{\text{g}}{\text{gm}}$
+ 1/2"	573	1.88	1.88	.030	1.71
- 1/2 + 3/8"	432	14.27	16.15	.018	7.78
- 3/8 + 1/4"	842	27.81	43.96	.065	54.73
- 1/4 + 1/2"	522	17.24	61.20	.042	21.92
- 6 + 10	468	15.46	76.66	.046	21.53
- 10 + 16	177	5.84	82.50	.060	10.62
- 16 + 28	161	5.32	87.82	.072	11.59
- 28 + 48	157	5.18	93.00	.094	14.76
- 48 + 65	54	1.78	94.78	.108	5.83
- 65 + 100	40	1.32	96.10	.132	5.28
- 100	168	3.90	100	.162	<u>19.12</u>
					174.87 gm-oz

81.97% - 3/8"

To B66 fix 3

$$H \times p = \frac{174.87 \times 100}{3028} = \underline{\underline{.0578}}$$

RS. 130-A

Regular Soxhlet extraction, 24 hr conditioning time, 29 hr extracts H₂O soln. properly left with some about 1/2, th alcohol. 3 ml 10% Na added, filtered and put into electrolytic cell.

boat # 130-A - preg. soln. before elect. 10 ml - 65.2 ppm
 130-B same soln. after elect. 25 ml - 1.04 ppm

Carbon

Bo. ready assay - 10g ore, 6g flour (2 spms), = 52g button

Continued on Page

Read and Understood By

Signed

Date

Signed

D

PROJECT P.S. 131 (Manhattan placer conc.)

Notebook No. P.S. 131

21

Continued From Page

100 g placer, sample separated with the magnet.

Kernets	Au B/T 4.015	gm/B 401.5	% DUST & % BRK FL
Lead 100 g		401.5	100.0 100.0
Magnetic 79.5 g	.048	3.816	79.5 .950
Non-Magnetic 20.5 g	19.40	<u>397.7</u> 401.52	20.5 99.05

99.05 g of the gold is in the non-magnetic portion.

Report Made

Continued on Page

Read and Understood By

Signed

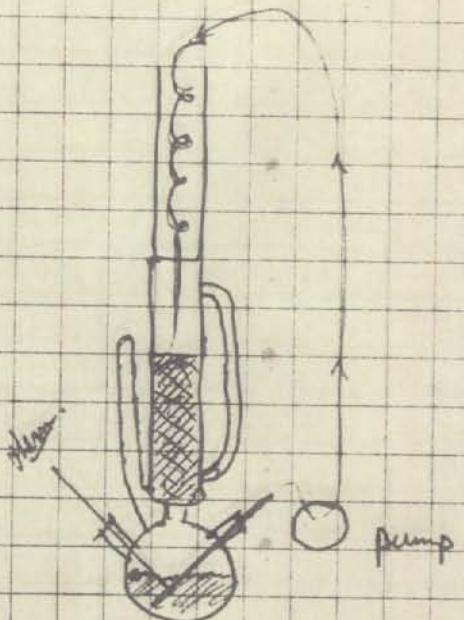
Date

Signed

Date

50 g small pellet field loaded carbon were added to the Sox Extractor. The carbon was washed saturated and then 100 ml more soln. Sol - Methanol strip soln. The complete system was circulated for 3 hrs., then the ash was changed.

Temperature of ash in flask 70°C, approx in the extractor.



heat and pump on at 10:15
1st ash removed at 1:15, all new ash
added, 107 ml.
flowrate 38 ml/min.

$$\text{Soln} = 87.98 \text{ g / 100 ml}$$

$$.8798 \times 38 = 33.4 \text{ g / min}$$

assays -

Soln. #1 - done 1252 ppm

Soln. #2 -

baren Carbon -

Continued on P.

Read and Understood By

Signed

Date

Signed

Sample from hole No. 32-51-75, 100 lbs cutout, 50 lb. of sample was put into the leach column and the leach started. The other 50 lbs. was split down and arranged for assay.

1 gal. Soln. Containing 10 g NaOH added = 3.785

No good
cyanide
percolate
will not
soil

Continued on

Read and Understood By

Signed

Date

Signed

PROJECT Wet Screen Analysis of McCoy drill cuttings

Notebook No. P.S. 135

Continued From Page

Sample of 32-51-75

765.3

~~1000~~ g as came from drill hole

		Our % Au	Ag Sp - g.	17.83 NaCl wt. %
+ 8 mesh -	225	218.2 29.1 .038	8.292	8.24
- 8 + 16	-	144.0 19.2 .092	13.248	13.16
- 16 + 35	-	113.3 15.1 .108	12.236	12.16
- 35 + 65	-	66.2 8.8 .146	9.665	9.60
- 65 + 270	-	82.5 11.0 .254	20.955	20.82
- 270	-	<u>125.0</u> <u>16.7</u> .290	<u>36.250</u>	<u>36.02</u>
		749.2 100	100.646	100.00

HXP .134 rising 749.2

HXP .132 rising 765.3

166 .061
167
168
169
170

A
Information Sheet
to R.R. futs

Continued on Page

Read and Understood By

1000 g ore

3000 ml solution at 8.0% NaCN, 5.0% lime? avoid

Roll of 72 lbs.

Ttate 4.02 * NaCN / Ton ore
.11#/T AuO40.2 ml AuO, 1lb = .1# NaCN/T
.55 lb ore acidic 1lb = .2 CaO

problem - 16.16 ppm

$$\frac{1}{2} = \frac{.11}{x} \quad \frac{x}{.11} = \frac{2}{1}$$

$$x = .2 \times .11$$

$$= .11$$

25 me = .404 kg Au

$$\frac{.404}{25} = \frac{x}{3000}$$

$$x = \frac{3000 \times .404}{25} = 48.48 \text{ mg Au out}$$

Tailings 1.03 Au/T

.015 g Au

~~1000 g @ .03 = $\frac{3000}{1000} \times .03 = .09 \text{ g Au/T} = .015 \text{ g Au/T}$~~

~~3000 ml @ 16.16 ppm = $\frac{48.48 \text{ mg Au}}{3000 \text{ ml}} = \frac{16.16}{34.3} = .48 \text{ g/T}$~~

~~Sb = 72.32 g/T $\frac{.48}{2000} = \frac{x}{3000} = .72 \text{ g Au/Ton}$~~

Total Au .015 g/Ton/1000 lb.

~~.720 g/Ton $\frac{.72}{3000} = .00024 \text{ g Au/Ton}$~~
~~.735 g Au total/Ton $\frac{.72}{.735} \times 1000 = 98.0 \% \text{ recov}$~~

~~HKP = $\frac{.735 \text{ g}}{1000 \text{ lb}} = 1.47 \text{ g/Ton}$ But (I think)~~

Information

Continued on Page _____

Read and Understood By

This test is to check test no. 136

1000 g ore from hole no. 1

3000 ml Soln.

$$\frac{7.5 \text{ g NaCN}}{20 \text{ g CaO @}} = \frac{5.0 \text{ #/ton Soln}}{=}$$

titrated 16.8 ml
25.0 ml

$$41.8 \text{ ml NaCN} = 4.18 \text{ #/ton soln}$$

titrated 5.8 ml Oxalic

3000 ml @ 3.2 ppm

$$\frac{3.2 \text{ ppm}}{34.3} = .0933 \text{ g/T}$$

$$\frac{.0933}{2000} = \frac{x}{3000} \quad \sim .13995 \\ = .1400 \text{ g Au/ton}$$

Tails 1000 g @ .052

$$.052 \text{ mg/2000 g} = .026 \text{ mg/1000 g}$$

.1400 mg / 3000 ml soln

.0260 mg/1000 g tails

.1660 mg Total Au in 1000 g head.

$$\frac{.026}{.1660} = 84.3\%$$

33.2 g

Calculated Head = .332 g/Ton

Information to
(I think)
further

Some non-magnetic plac conc was added to a beaker along with with 1:1 HNO₃ and boiled, then filtered. The filtrate was assayed for Au-Ag, as was the residue, a fusion was run on the residue.

test p.s. 138 25 mc

222.6 mg Au-Ag from plac test sample of 10 Kilo (+10 m)

Continued on Page _____

Read and Understood By

Signed _____

Date _____

Signed _____

Date _____

4 - 10 ml samples of pres rock. soln. put onto 250 ml beakers

1. The 10% Na₂S added along with lime
2. The " "
3. 10 ml " "
4. 20 ml "

soln. digested, and ppt. formed and washed

		Ag	As
boat 139-1	5 ml	8.354	1.386
"	-2	8.526	0
"	-3	8.432	.178
"	-4	7.740	0

Port 139-1		.196	.204
"	-2	.116	.024
"	-3	.140	.270
"	-4	.038	.102

1st Rocket soln. - Sodium Sulfide added - approx. 500 g / 40 gal so
Sample then taken for electrolysis. wool boiled away with HCl
Sample of As taken for frieze assay

As - Ag wire	.08389	g
Ag - added	.25025	g
	.33414	g

Std.	.24986
	.24430 : 2.25% loss
	.00556

$\frac{112}{4}$
 $\frac{4}{2}$
 17

Reports to get

- R.I. 7464 - Malmonite Extraction of gold from ore
- R.I. 7660 - Recovery of Ag & Hg from mine tailings by Electrooxidation
- R.I. 7395 - Leaching Au - Bearing Mine tailings from Nevada U.S.A.
- R.I. 7597 Extraction of Au from Carbonaceous Ores: pilot plant studies

Summer mileage

start Summer Sat.	34555 = 151
To Round Mountain, Marlboro by to Tonopah.	34706
Tonopah to Lone Pine	34709 = 172
Lone Pine to Darwin	80 miles 172
Lone Pine back to Tonopah	34164 = 112
total for week 547 @ .15 = <u>82.05</u>	284

Circulating pump.

Cole - Palmer
 1 No. 7545 Variable speed drive
 with Solid State Controller 147.50
 1 No. 7014 Std pump head 35.00

Call collect to 312-647-0272

Cole - Palmer Instrument Co.
 7425 N. Oak Park, Ave.
 Chicago, Ill. 60648

$\frac{172}{2}$	172
$\frac{151}{2}$	151
$\frac{172}{2}$	172
$\frac{172}{2}$	172
$\frac{112}{2}$	112
$\frac{60}{2}$	60
$\frac{80}{2}$	80
$\frac{424}{2}$	424
$\frac{667}{2}$	667

50' No. 6408-43 Tygon tubing @ .22/ft.

Copy and P.U. file
 8-15-75

~ 150
.04
/ 0
6.0

To do

1. Cost report a heap leaching #1 priority
2. See about tunnel sizes at bogie mills
- ✓ 3. Start dump tonnage at also pick sampling and extraction procedure.
- ✓ 4. See what the other minerals are in Tonapah dumps.
5. Lab work, lab float Cell, filter, set of assays
(and about costs)
6. ~~Cost At Monte Tom~~
7. Get tunnel heaters at Monticello (2)

For Capital City equity Co. 150.00/day plus ~~expenses~~

- ✓ 1. How important this clay, Boronate, is to scheme of things.
- ✓ 2. Get cost of mill from Cyprus, also see what replacement cost would be
- ✓ 3. find out amount of land there also.
- ✓ 4. Ask Mulvey about haul, from Reeler to Dunn - Pid set in, how new equipment can do best job.

Friday or Saturday

- ✓ go to Monte mill to get samples from non working there to run metallurgy. \$75.00 plus assays.

Summarize An leach cost analysis.
Manhattan project

- A. Mining - 40 - 50,000 tons
- 1) ore deposit preparation, stripping -
 - 2) mining costs -
 - a) ripping and picking up with cars. (Rides common in)
 - b) ripping and picking up with loader and trucks)

B. Pad -

- 1) Excavation of pad area
 - 2) pad preparation, material
 - a) material for bottom of the pad
 - b) sand and gravel for pad base
- cost known approx

C. Pond -

1. pond excavation] cost known approx.
2. pond lining]

D. Soh stripping plant -]

1. Cost to owned plant] cost known approx.
2. Cost to run plant]

E. An-Ag stripping plant (in Tonapetl)

A. Mining

Striping, all over burden can be push to the east, toward town, this shovel took only 2 ~~days~~ days with the D-8 cat. Cost - 34 hrs at \$40.00/hr. cost if hired.

\$1060.00

An Method for Solution by A-A

1. pipet 10 ml of Sample into a large screwtop test tube.
2. add a crystal of KMnO_4 to each ~~cold~~ test tube.
allow to dissolve, heat in water bath if necessary.
3. add (with a burette) 2 ml HCl
4. let stand 10-15 min. until color is gone, or mostly gone.
5. add 5 ml M.T.B.R to each sample
6. Shake 2 min.
7. Run wash layer on A.A.
8. prepare std. same as samples
prepare std. to be extracted fresh daily
1, 2, 5, 10 ppm std.

P.P.M.

9.

10.

2.0^v

Mineral Eng. Monte-Verde
Angels Camp.

Ball mills - Denver Ball-Rod #24
need special clamps to match

1.20 yd² - 20 mil

U.S. Machinery Co. - 786-0300

Tony Selig -

Jen Wilson - 14"-16" about 10' long 359-2669

Minerals Equip. Salt Lake M.E.C. 801-262-5471

See Salt Lake phone book

Machinery Center Salt Lake 801-484-7601

Denver Equip. Denver - 803-831-7311

- Ark about -

Zinc dust precipitator, filter
screen clarifier

Jaw Crusher 18 x 36 or there about

Merill-Crowl. - S.A.

9" X 7' = about \$1000.00

Koruktake - thinks he has Merill-Crowl unit] minerals

- has 9" X 7' classifier] Equip.

Mont Verde - ²⁰⁹ ~~236-2713~~ | 209-736-2626

430

740
110
510

Las Vegas Truckers -

Espy Bros. Cons. - 648-0030

My Trucking - 385-4151 or 642-0440

Wheeler Trucking - 878-9636

Mileage

1. To Tonopah for work 112
2. To mine and back plus trip to Round Mtn. 151
3. Tonopah to Darwin and back to pick up
Leathers 284

$$547 @ .15 = 82.05$$

$$40,000 \text{ Ton } @ .07 = 2800 \text{ q/Au}$$

$$2800 @ 150.00 = \$420,000 \times .75$$

$$= \$315,000 \text{ total # avail. @ 75% use}$$

$$\begin{array}{r} .07 \\ \times 40000 \\ \hline 2800.00 \end{array}$$

$$\begin{array}{r} 4 \\ \times 150 \\ \hline 600 \\ + 2800 \\ \hline 140000 \\ \hline 140000 \\ \hline 1420000 \end{array}$$

$$\begin{array}{r} 1 \\ \times .75 \\ \hline 75 \\ \hline 1420000 \\ \hline 1065000 \\ \hline 2940000 \\ \hline 3150000 \end{array}$$

Summa P.E. 107 A-A.

Pb Std.

4g of 50.1% Pb up to 1000 ml

1 ml = 2 mg Pb

	g Ag	ABS. READING	$\% \text{ error is up}$ to 250 ml
1.	1 ml = 2 mg Pb = .338	.003	.2
2.	10 ml = 20 mg Pb = <u>.338</u>	.032	2.0
3.	20 ml = 40 mg Pb = 6.75	.053	4.0
4.	30 ml = 60 mg Pb =	.076	6.0
5.	40 ml = 80 mg Pb =	.100	<u>+0.0%</u>
6.	50 ml = 100 mg Pb = <u>16.88</u>	.126	10.0%
7.	75 ml = 150 mg Pb =	.180	15.0
8.	100 ml = 200 mg Pb = <u>33.76</u>	.230	20.0%

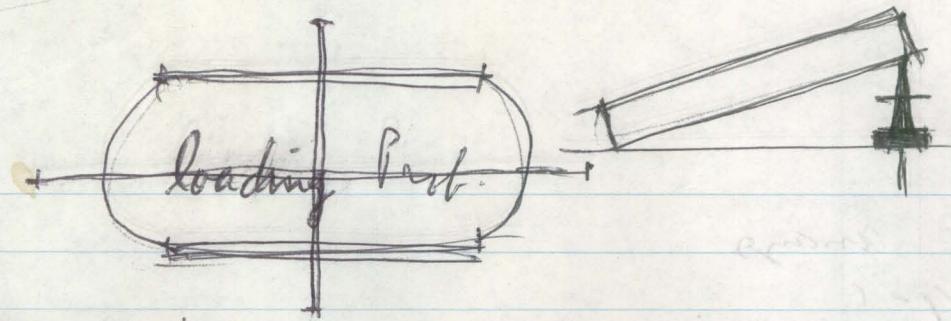
Do

- Call Dad about - Having Women do complete license for Much faster, can have complete by end of the week.
- Tell about end dump in Texas

17.0
129.0
13.3

15.0

Ilo



Call Earl Dringman - operator 6

702-786-0300

Call Orion research 11 Blackstone, Cambridge Mass
617-864-5400

procedure manual for
cyanide
filter

when are you going to get a gold electrode

minimizing cyanide exposure.

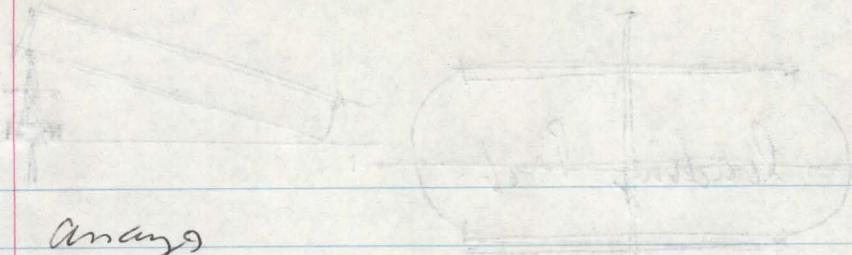
2 atm. 1%, 1 to 10 K

Temp to 20 or

slope 12%

electrode in

good has 95 to 100% slope



Anaays

NP-1

NP-2

NP-2A lost some of it Corel no low

NP-3

NP-4

Beagle Sam ore

5.5# rock

start with 6 l rh.

to Satmatc 4 l

Takes 15 minutes to flow through the 7 ft. column.

pH 6 after running H₂O through the ore

added 2 gm NaOH to bring pH up to 11

put in 24 g of NaCN to start with trying to keep

3.5#/ton ore.

1st sample taken 5:30 June 21

2 sample .. 9:00 June 22 conc. NaCN #5/ton ore.

June 23 NaCN #5/ton ore.

afternoon .. NaCN #34/ton ore.

Sample 1 1.95 g/t extra 6 hrs. pH 11 conc.
3.8

Sample 2 4.5 g/t after 20 hrs.

Sample 3 3.914 g/t

Sample 4 3.7

Sample 5 3.72

Gold in sample showed .145 g/ton in Au

\$8
1/0

charge cost



Beall - Lopez -

$$ft^3 = 40 \# \quad 227 \\ 40 \\ \hline 1080$$

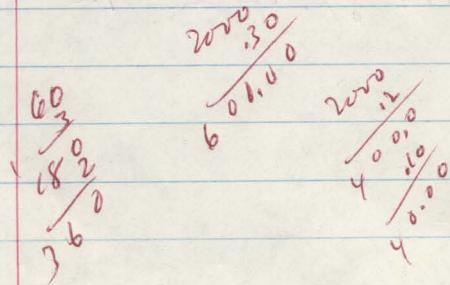
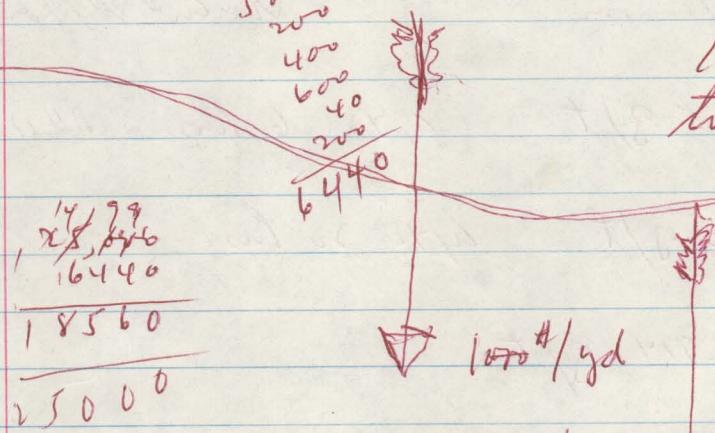
Ron Beers
685-8500

Gis. doing.

Jim Groendyk
213-353-7287

thaw - 26' aluminum

Cook = 13,500 with (P.T. & pump - tank) = 1000 ft wet
kit & cover tire.



5:15

Sodium Bor-Anhydride

will ppt. the following elements

Antimony
Arsenic
Bismuth
Cadmium
Copper
Germanium
Gold
Indium
lead
mercury
Osmiridium
platinum
renium
rodium
 Ruthenium
Silver
Thallium
tin

Ethanol Extraction

Activates with -

But for like not as good for yield

approx - 2-4 bed volumes -

1% - NaOH - up to 3-4% will keep acetate down

.1%, NaCl

up to 25% H₂O

Ethanol.

Activant formed due to electr.

Tom Cernahan - red sample of carbon up
to U.S.B.M. Reno

3 Volts

1000

.01

1000
.001

1.000

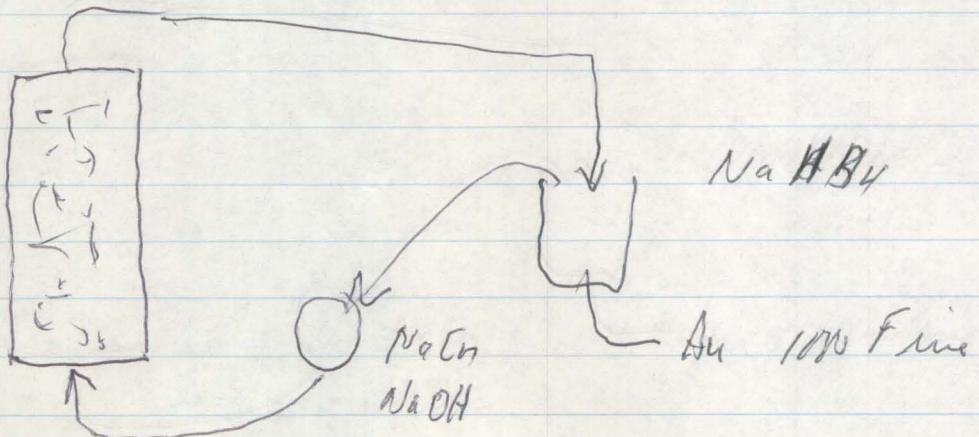
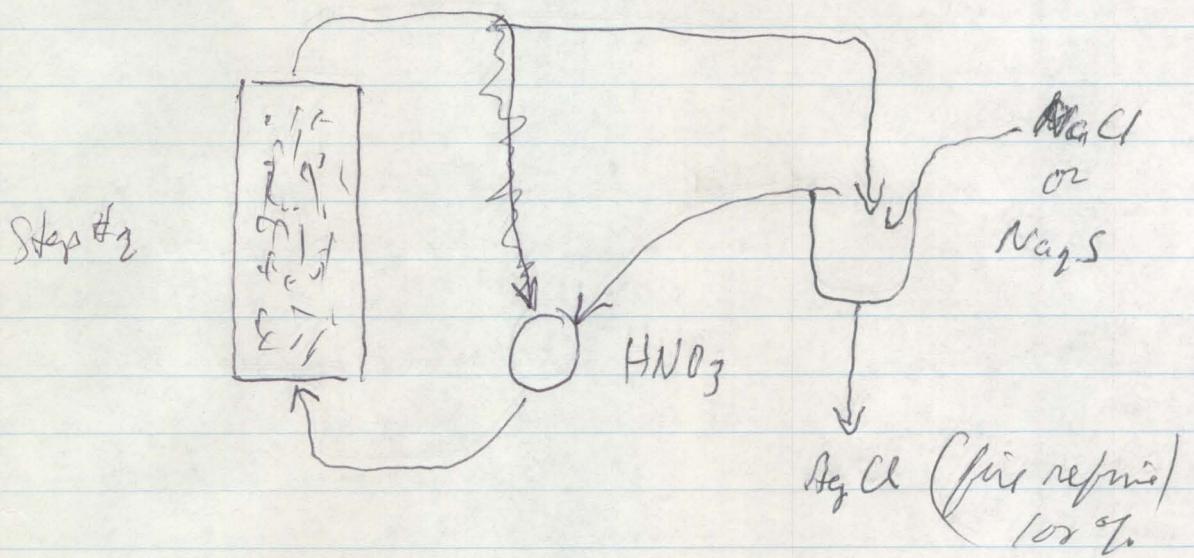
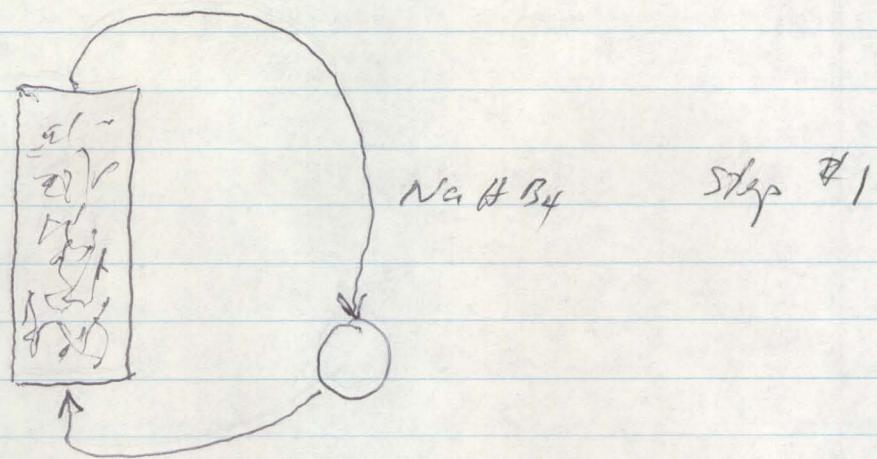
750 ml Ethanol

1000 ml -

200 ml H₂O

15 g NaOH

1 g NaCl



Carbohydrate Count

~~Thurs.~~ Sept. 3

Wed. Sept 4

No breakfast

7

Lunch - corn beef, cabbage (one wedge)

3

- Salad, cheese dressing,

2 5

- $\frac{1}{2}$ cup peas and carrots mixed

17

- small portion of Jello.

0

Supper - T-Bone

3

- Salad, veg. pt

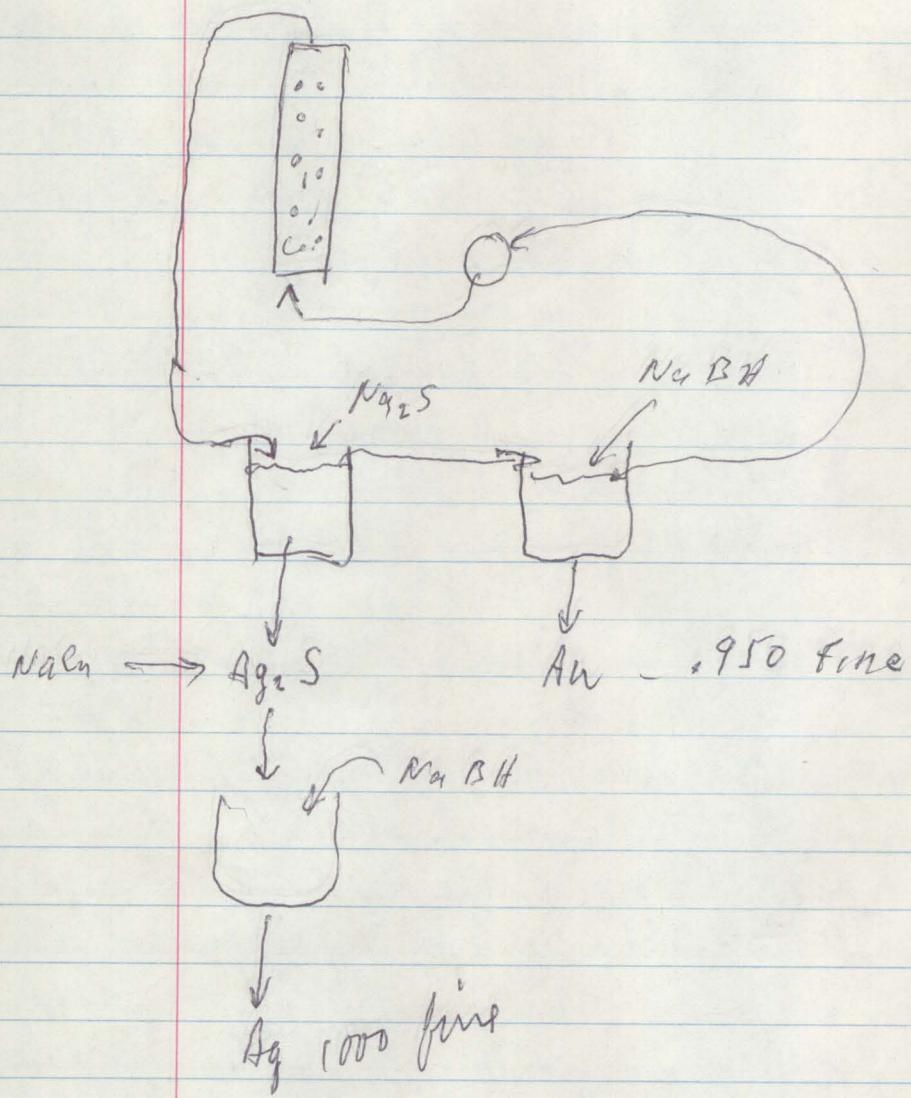
12
47

- 1 piece of toast

1. Song. 55.00
2. Sandy 55.00
(3. " 594.00
4. " 36.00
5. 36.00
6. 280.00

Agr S

Ethanol



1. Leach test on Sawyer ore
2. " " " Merritt Conc.
3. " " " Merritt ore.
4. Make float reagent
5. Float test on Hard Cash ore.
6. Test ethanol method
7. Check nitric acid method

Call Pacific Foundry

Rolli Brown

Ethanol - Commercial Solvents

55 gal barrels San Jose 408-262-2474

formula 1	50A	1.68/gal + freight
"	3A	1.67/gal + "
"	3B	"

Phellacol 1.71/gal + freight

Anahorse " + "

Guixol " + "

Ask about → Fotocol form A2 1.43/gal F.O.B. Calif.
3-9 barrel lots

Dore

Au-Ag

P.S. #	Description	Value
P.S. 5	good looking Poly Ag ore dark colored.	3177.02
P.S. 6	yellow Bi? rock	28.9
P.S. 7	1/2 sample (parent 1/2) of first damp rock	^{1584.16} 1584.16
P.S. 8	west end sample.	43.04
P.S. 9	little (1/2 rock with native Ag)	300.82
P.S. 10	I.A.T. frag. rock.	105.22 270.44
P.S. 11	Cerargite sample	19,851.8

714- 8764102

Skinner

Luson Sonopac files

RL-7-75-70

MEMO

TEST RESULTS OF
MARY MINE ORE

ORE: from: Mary Mine Open Pit

type: Quartz + Alaskite with FeO's and Sulfide

head assay: = .078 oz. Au/ton

PROCEDURE: 50 lbs. of ore was crushed to minus 3/8" diameter and placed in a column with well water from a well in the Mary Mine area. Water had a pH of 6.8. This test had a 9 day leach using 2 lbs/ton NaCN. The pH was maintained at 10.5, test used 65 gms caustic, and 30 gms NaCN.

RESULTS: Tail Assay = .012

1. 84.6% total recovery
2. 5 to 9 days peak of leach time
3. 57% of gold is from (+)1/4", (-)3/8" material
4. One ton ore @ (-)3/8" holds 30 gals. solution in retention.

NOTE

The above information taken from
Paul Skinner notes and dictated to
Robert Lutz on 6-30-75.

In

cc: R. Lutz
D. Gribbin
W. Mollison
W. Simmons

Skinner's Expense Reports

October	240.90	
	137.10	
	114.08	
		492.08
Sept	341.88	
	231.46	
	107.15	680.49
Aug	133.40	
	91.38	224.78
July	117.97	
	178.15	296.12
June	259.99	
	94.21	354.20
		2046.

400/mo
4800/yr