

-Treatment-
- of -
Nashua Copper Ore.

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Treatment
of
Nesbitt Copper Ore
Woodstock.

The ore.

The kinds of mineral composition that make up the ore. Chemical analyses of the elements found in the ore.

The Kiln. description. its merits. Methods of working the kiln. the roasting. the amount of fuel, draft, time. and suggestions for future work.

The Furnace. Plan, after being rebuilt and bricks put in.

Plan of the bottom. with overflow arrangement. Substitution of one type. subsequent working of same.

Details of run. I, II, III

Changes .. feed and tap record.
Remarks on the working of the furnaces
during this run.

Anal. of matt. slags and ends in wt.
Chemical analysis of the products of this
run.

Roasting of matt. in Kiln and in
the reverberatory furnace. Comparison of
work done by the two furnaces.

Summary -

Chemical methods used -

Color determination of copper. Tables

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1

The ore - to be treated, was given to me already sorted and reduced to a uniform size - and was quite free from dust.

The average size of the ore - was from the size of an egg - down to pieces of the size of a marble.

It was remarkably free from gangue - it being cobbled for us - of extra richness -

what gangue were to be found, consisted mainly of mica schist - a small amount of quartz and mica.

The ore as treated at the miller - has only six per cent of Cu - but that sent here was cobbled of extra richness - the percentage being double that used at the miller - that is twelve per cent.

The mineral composition of the veinstone is mainly of coarse crystalline Chalcopyrite, Pyrhotite - with small quantities of Blende and fine quartz scattered all through.

The relative proportions of the principal constituent parts are approximately as follows

chalcopyrite----34. per cent
 Pyrrhotite----58. " "
 Quartz-----5. " "
 Blenda-----2. " "

The ore was sampled and treated by chemical processes, wh showed its composition to be as follows-

These analyses are those of Mr. Southworth and Mr. Bautol-

Mr. Southworth.

S	33.46	per cent.
Si-mol	6.40	" "
Li-pol	15	" "
Cu	11.25	" "
Fe	46.49	" "
Zn + Co	1.66	" "

Mr. Bautol.

S	33.97	per cent
Si-mol	5.45	" "
Li-pol	20	" "
Cu	11.29	" "
Fe	46.76	" "
Zn	1.27	" "

in each case slight traces of

Alumina

Magnesia

Alkalies

Calcium

Nickel & Cobalt

As shown by the table of percentages - the amount of S_2 is quite large - it now is necessary to roast off about a half or two thirds of the Sulphur - before smelting for metal.

Until lately - it has been the custom to crush and roast in the reverberatory furnace -

Last year a kiln was built by Mr Southwolfe for roasting - and carried on successfully by him -

Its only advantages are - less amount of fuel needed, and less amount of work - while on the other hand - it is quite uncertain - it being impossible to roast "dead" with it - and needs skill and experience to run it well -

On a larger ^{scale} however, I think it is much more economical than the more certain yet more expensive reverberatory furnace -

I shall endeavor to show by comparison with work in Kiln and the same amt. of work done in Reverberatory furnace - the amt. of

labor and fuel were used, to produce the same results.

The Kiln is made of fire brick - rectangular in form, with an arched top.

The following figures give the interior dimensions - on floor - 23" x 18" wide height - 43.5"

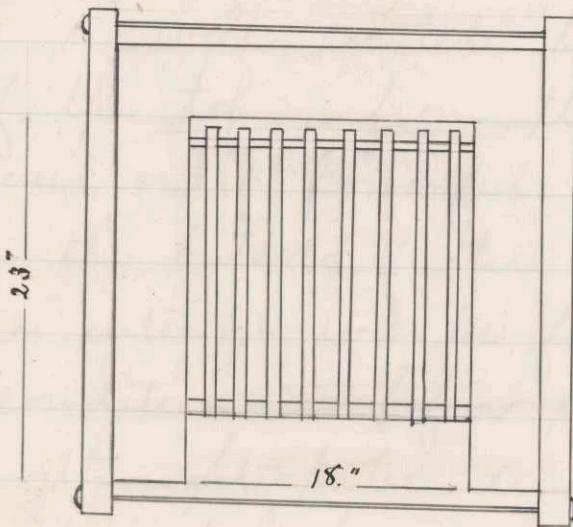
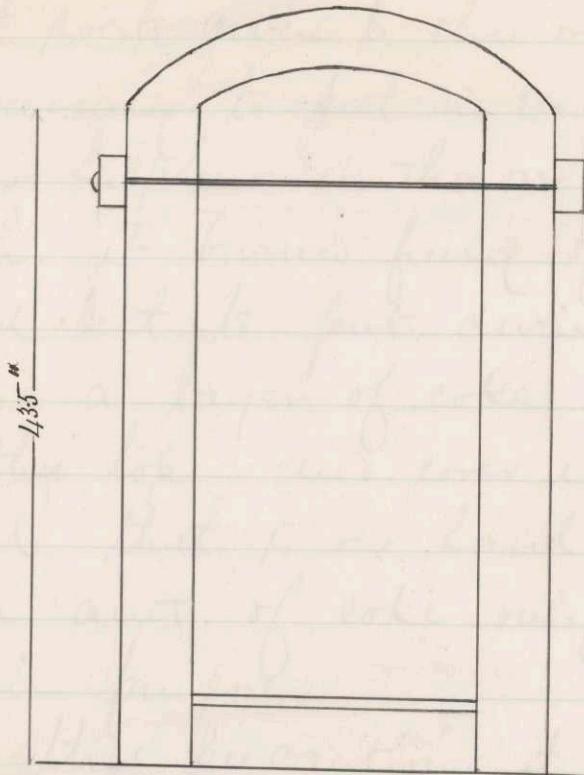
It has a grate about 5" from floor, with movable grate bars -

To start a roast - it was my custom to have a good supply of shavings to start the kindling wood and preclude the possibility of it going out.

Then on the shavings - an amount of charcoal - that varied from 4 - 6 lbs

This charcoal I found necessary to get the coke well under way -

On the charcoal - was put about half a hod of coke - then the ore was put on and it depended on the ore whether any more coke was put in or not.



"Plan of Kilu.

The first roasts given to the ore it was only necessary to put in enough fuel to get the Sulphur in the ore well on fire - after wh. it burned freely of itself.

It is found best to put during these first roasting a layer of coke about a foot from the top and cover it with small material that is on hand -

A small amt. of coke only is needed for this purpose -

Without this precaution it is found that the ore is well roasted to within a foot of the top. - from this point the fire seems only powerful enough to oxidize the outside of the lumps of ore and the interior will be found in the same condition as before roasting -

This is thought to be due to the too free escape of heat - wh. not having a larger body of ore to confine it easily passes off - only touching the outside of the lumps of ore

This remedy was tried and it was found that the space was lessened - that is - from 4"-6" from the top it was not much roasted - It was however an improvement on the former roasts.

After the ore had been roasted twice it became necessary to increase the amt. of coke, as the amt. of Sulphur in the ore was less -

This was done by putting in layers of coke - {it could hardly be called a layer but rather a judicious sprinkling of coke over the surface of the ore,} and then filling up - about 10" of ore - then repeat -

The average time taken for a roast, was about 30 hours - that is - a roast that was put in at 10 A.M. of one day was taken out at 4 P.M. of the following day -

It was found best not to hurry the operation but to let the fuel burn slowly, with little air let in at bottom - the whole front being plastered.

The following table show the amount of ore, coke, charcoal-shavings, that were used for each roast.

I

5% 4 lbs. Cu ore.

coke, 24 lbs - top not good
char - 4 " middle - good
wood, shavings, 6" bottom - caked

At one time the fin must have been too hot, as the lower foot was caked, solid, showing that there was too much fuel at that point and not enough fuel at the top.

II

590.5 lbs Cu. wt. — top part roasted at all
coke 20.5 lbs centre. good
char. 6.5 " bottom. caked.
wood shavings. 6"

The conditions of this road must have been like the first, as the result was much alike.

III

659. lbs Cr. Ore.

Coke. 14 lbs

→ did not roast at all.

char. 3.5 "

made second roast.

wood. shavings. 5"

ore. same.

coke. 25.25 lbs.

very good result.

char. 5.25 "

very little cake at

wood. shavings. 8"

the bottom

The first attempt was a failure owing to the very small amount of fuel used.

The only result of the roast was a complete burning out of the fuel which only gave out enough heat to oxidize the outer coating of the lumps of ore.

The second roast of the same ore. with a different arrangement of fuel. was a success. the ore was but little caked. A layer of coke was put near the top wh. aided the roasting of the top very much.

III

621.25 lbs Cu. ore - very good roast, a
 coke 24 lbs. small amt. of ore
 char. 4.75 " not roasted. found near
 fuel 7: the front, near the cracks.
 very little bottom

distribution of coke same as in last
 roast.

The ore was then taken and sampled
 and a chemical determination made for
 the amount of S it contained.

The result of this analysis showed
 that the ore still contained 18.51%
 of S - which was judged to be too
 much. as it would give us a
 matte of but little more richness
 in Cu. than the ore itself.

Accordingly the whole amount was
 run through the kiln a second time
 with beneficial results - as the amount
 of S was reduced to about 10 per cent.

I.

char. 5 lbs

coke. 35 " ~ my good roast.

fuel. 8 "

II

char. 5.5 lbs.

coke. 32 " ~ fair roast.

wood. 7 "

III

char. 5.5 lbs

coke. 40 " ~ good roast.

wood. 6 "

IV

char. 5.5 lbs

coke. 33 " ~ good

wood. 5 "

This we see picked over and the pieces
 that were not roast enough, put back
 for another roast.

V

char 6. lbs

coke. 28 " not very satisfactory-

wood 6 "

The ore was now sampled and an analysis made for the determination made of the 'S' that remained from the two roasting.

The analyses gave - 6.21% and 6.52 These results were wholly unexpected and for this cause, were discredited and in the calculation for slag, the determination was ignored - and the calculations were made from a basis of 10%.

Subsequent results showed that we had figured too low, even at that percent. The cause of such a low estimate being made will be explained in the chapter on the method used for the chemical determinations.

The Kiln-läken altogether did its work well - and by carefully arranging the fuel - very good roasts can be had. The damper that was put on, I did not use often for the purpose intended, but rather as a "peep hole".

The ore weighed before roasting. 2421.75 lbs.
 " " " after " 2340.5 "

This loss is not due entirely to s' but also to the loss of fine dust, which the best of care cannot prevent. This loss is not great as it would seem, but the ash, of the coke, charcoal and the shavings, make it appear less.

Preparation were now made for a furnace small for metal, containing about 25% of metallic Cu.

Before the run was made however the furnace was torn down and "boles" put in - also one tuyere was substituted for the three that were formerly used -

As this took some time I made some Cu. assays -

The first assay - gave me only a button of metal - and as I thought that the roasting was complete - I put the blame on the Soda used -

I therefore tested the Soda for 18, and found a larger amount - wh. showed clearly where the trouble lay -

The second time, chemically pure Soda was used and a Cu button was obtained wh. gave only 7%.

Again I tried it - carrying along two assays. using the same Soda, and the results were quite good -

In one case the button was free from mala and looked clean, did not break under the hammer and was quite flattened - the percentage obtained was 10.9. wh. comes near the determination by the wet method. 11.25.

The second crucible did not get enough heat - the Cu. being scattered through in fine buttons - not result obtained -

By practicing constantly this method could be made to give very fair results but the time being limited - this method of determination was dropped -

The bottom of the furnace was arranged so as to give an overflow of the slag - at the top - and at the lower tap - the metal was run out, being tapped and plugged - The figure explains itself.

At the slag outlet charcoal was put. wh. by burning, kept the slag from getting cold. One layer was substituted - instead of stone that had been used in the Pb. run - of 1.5" diameter.

A new Roots Blower was substituted for the old Sturtevant-Blower - and it has been found to work much better than the old form. not only a higher pressure being obtained: but more air was sent into the furnace.

A gate was put into the air pipe, so that the pressure could be regulated at will.

Plan of the

Furnace

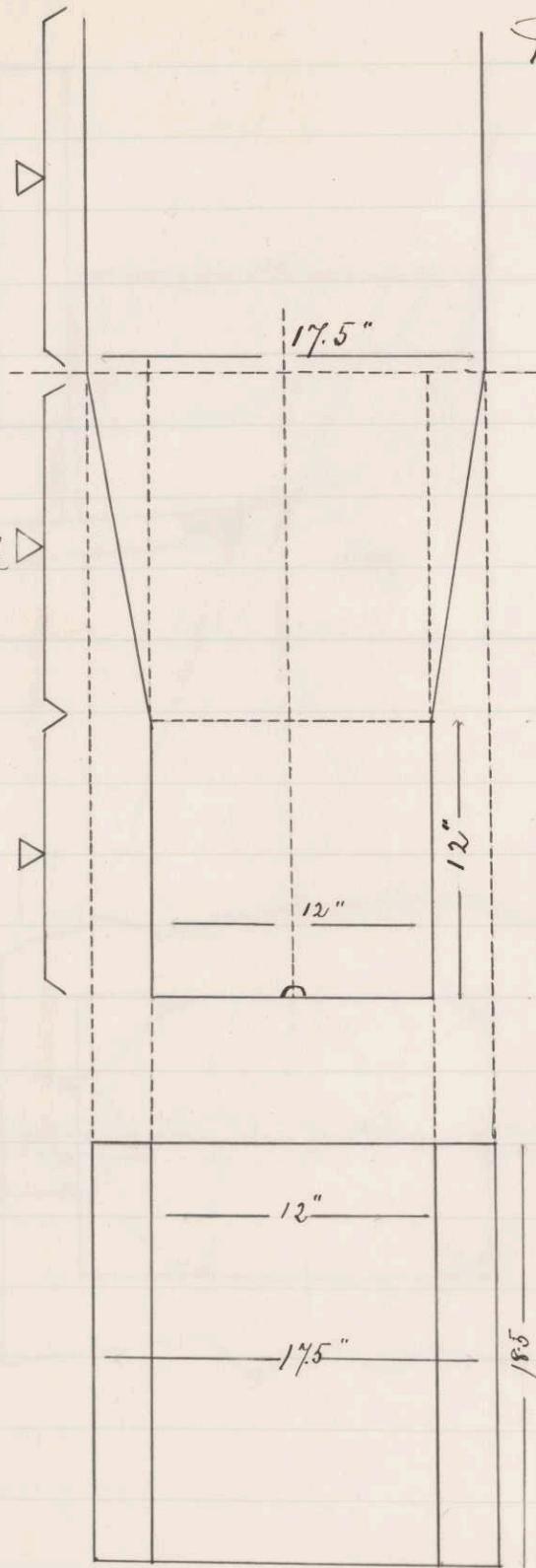
after being
rebuilt.

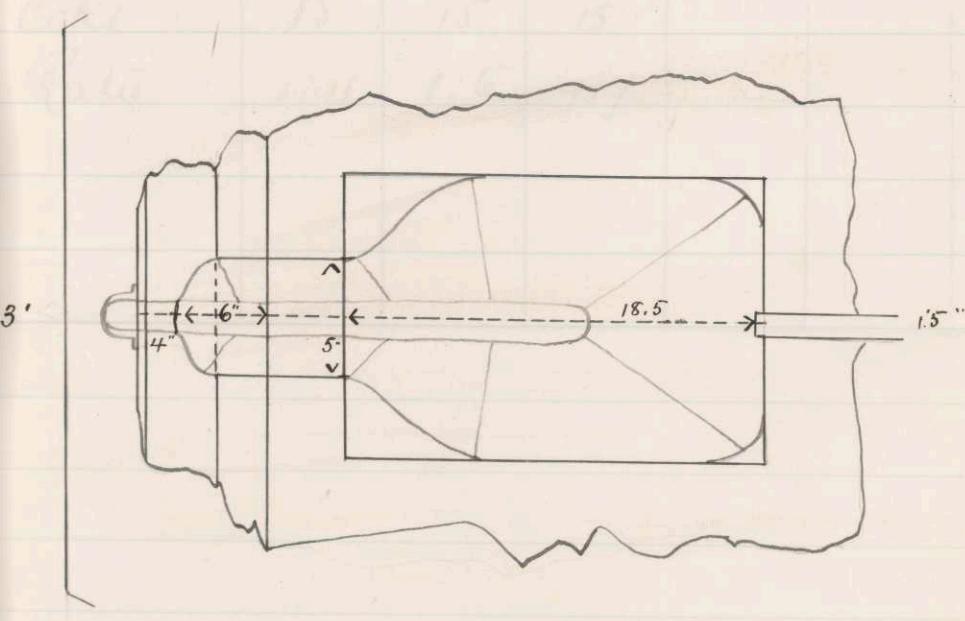
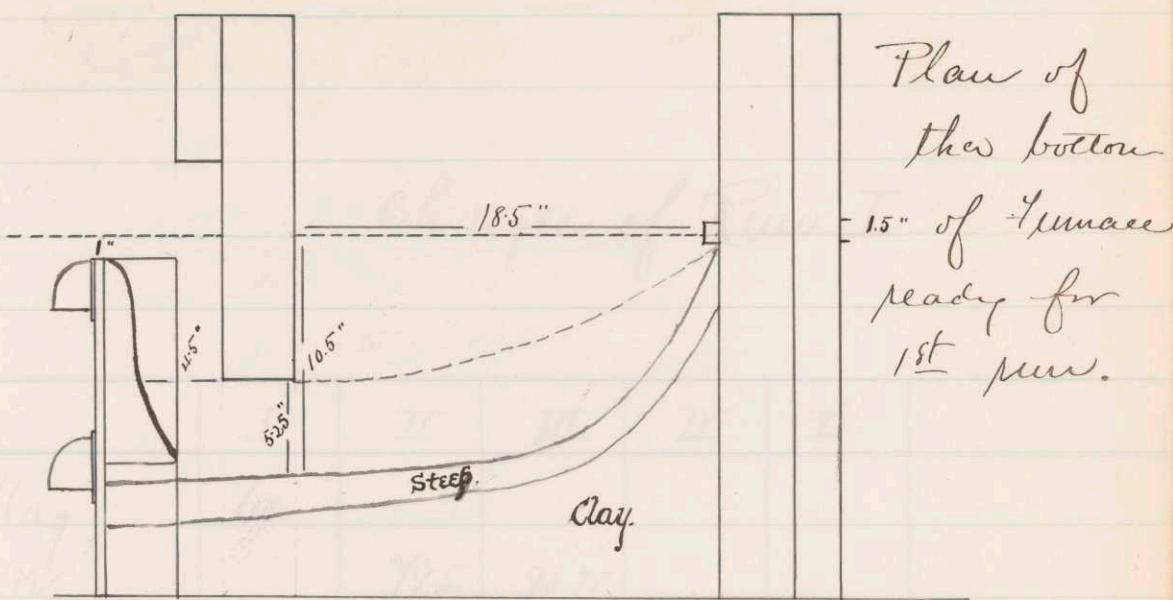
and "bodes"
put in.

Straight
courses.

5 courses bend.

4 courses
straight





Charges of Run I

	I	II	III	IV	V
Slag	60				
ore		79.5	90.95		
Sand		11	12.5		
Total		90.5	103.25		
Coke	15	15	15		
Ratio	1:4	1:6	1:7		

Feed Record - I

Time	Interval	charge	depth		Notes.
5. P.M.		1 hood charcoal.	4 hoods coke.	to start	
8.10	3/4	" "	5 "	" .	Blast on.
8.15		I			
8.50	5	I	-		
8.52	2	I	2.5		
8.55	2	I	3.		
9	5	I	3.		
9.10	10	II	3.		
9.16	6	II	3.2		
9.22	6	II	3.5		
9.30	8	II	3.5		
9.38	8.	II	-		
9.46	8	II	-		
9.54	8	II	-		
10.2	8	II	-		
10.12	10	II	5.		bad stoke
10.25	13	II	5.		bad coke with
10.35	10	II	5.		bad coke with
10.40	5	II	5.		slag flooded
10.45	5	II	5.		
10.55	10	II	5.		

Feed Record. I.

Time	Int	charge	depth.	Notes
11.05	10	II	5.	
11.15	10	II	5.1	
11.25	10	II	5.1	
11.40	15	III	5.	
11.50	10	III	5	
12.6	16	III	5.1	
12.21	15	III	5.	
12.34	16	III	5	
—		II		
—		III		lost top coke
—				when the furnace began to fire
1.50	13	III	4.5	
2	10	III	4.5	
2.10	10	III	5.	
2.20	10	III	4.8	
2.40	20	V	4.5	Last of the one 3 hds coke on top
3.15	35		3.	furnace hot on the back
3.30	15			slag - stopped.

-Tap Record-

I.

Time	Int.	Matte	Slag	Notes
9.10				Slag appeared.
9.16			I	
			II	
9.23			III	
			IV	
			V	
9.30			VI	
9.45		I		
10.1		II		
10.30		III		lost lap hole.
12.05				when the furnace was
12.05		IV		tapped a break
12.05				occurred - lost the
				upper run.. took off
				the blast. and sealed
				the leak with slag
				wh. hardened.
1.38				Blast now again
1.45				going well.
-				
3.35				End -

In the early part of the run, the lower tap hole was lost so that the slag and matte had to overflow together over the top run. Toward the end of the run a break occurred between the iron plate and the brick work letting the contents of the furnace out onto the floor. The blast was taken off immediately and by means of new sleep and with the aid of the slag itself (the slag having in cooling) the leak was stopped and the run proceeded.

After the lower tap hole was lost no coke was kept of the number of the buggies or of their amount. It was on account of the foulness of the run that this was done.

The entire slag was filled with matte so that the matts with slag as soon as cool, were dumped into a heap to be sorted and picked over.

The first few buggies that we got from the lower tap hole were very rich

Matter

Copper	21.40 .. 21.32. %	} 97.467 %
Iron	52.37 .. %	
Sulphur	22.70 % .. (14.4 : 16.64)	
Silica	.997. %	

Slag

Copper	.876 .. .895 %	} 103.33. %.
Iron	42.52. 780. 66.8 %	
Silica	35.65. %	

Foul Slag

Copper	4.83 % .. 4.77 %
--------	------------------

Euds

Copper	4.48 % .. 4.43 %
--------	------------------

Analysis of the products of
1st run.

The results obtained from the matal are very satisfactory - but those from the run slag are very poor.

The trouble was in the solution used for the titration of iron. too much HNO_3 being used to change the iron into the ferric oxide.

This free HNO_3 . in the sol. caused the poor results. I think this may be the cause. but am not certain
And as the time was getting short.

I did not try the analysis over again.

The determination of Copper in the Foul slag - and in the ends. show what a waste took place by the poor working of the overflow arrangement.

It put the whole operation back. this having so much Foul Slag - and matal in the ends.

For it became necessary to roast the matal. to get rid of the Sulphur.
but we see that in the two other products the foul slag and the ends

there was enough Sulphur in combination with Copper. to hinder the operation of enrichment.

For if the mass was roated to any low per cent. the resulting matte of the next fusion or smelt. would be but very little richer.

This turned out to be the case. it being only a gain of 7%.

The matte was run through the crusher. and was roated in the kiln.

The same arrangements were made as in the first place - the ant of S. being quite larger. only enough fuel was put in at first. to light the S. when it would burn of itself. Towards the last of the operation more fuel was put in. and a much higher heat obtained.

Roasting of Matte.

I { coke. 20. lbs good roast. top not much
 char. 5. " touched - a little
 shaving sc. 5. " melted and ran through.

II { coke. 18. lbs fair - top not much
 char. 4 " roasted.
 wood sc. 5. " did not melt any -

Ia. { coke. 28 lbs good roast - but not
 char. 6 " enough fuel.
 wood sc. 6. " more needed.

IIa. { coke 30 lbs bottom not roasted.
 char. 6 " center and top - good.
 wood sc. 7. " - more fuel needed

Ib { coke. 40 lbs very good roast.
 char. 5 "
 wood 6. "

II b. { coke 45 lbs at bottom not much
 char 11.5 " roasted.. but at
 wood 6" center and top - very good

Through the active operation of this
 heat. I find that the bottom of
 the Kiln - just above the grates has
 about a foot - that the ore did
 not roast at all.

The fuel seemed to burn out. with
 out any effect on the charge at
 that point.

But from about a foot above the
 grates. very good results were
 obtained. the ore being converted
 into a porous condition.

I tried an increase of fuel -
 but the result was not marked
 with any degree of success.

The reason of such action. has
 not been satisfactorily explained

An analysis of the matts was now made for the S. it contained.

The analysis showed that it still contained

13.17%

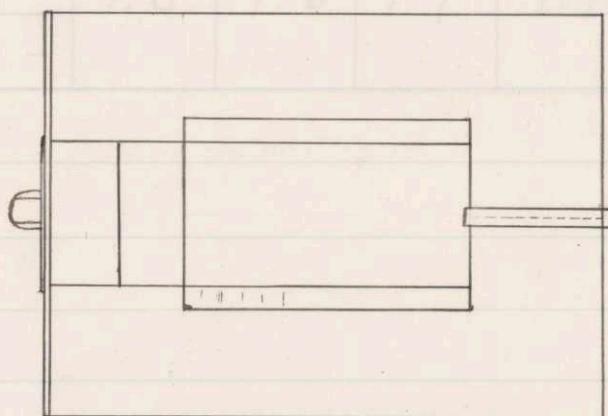
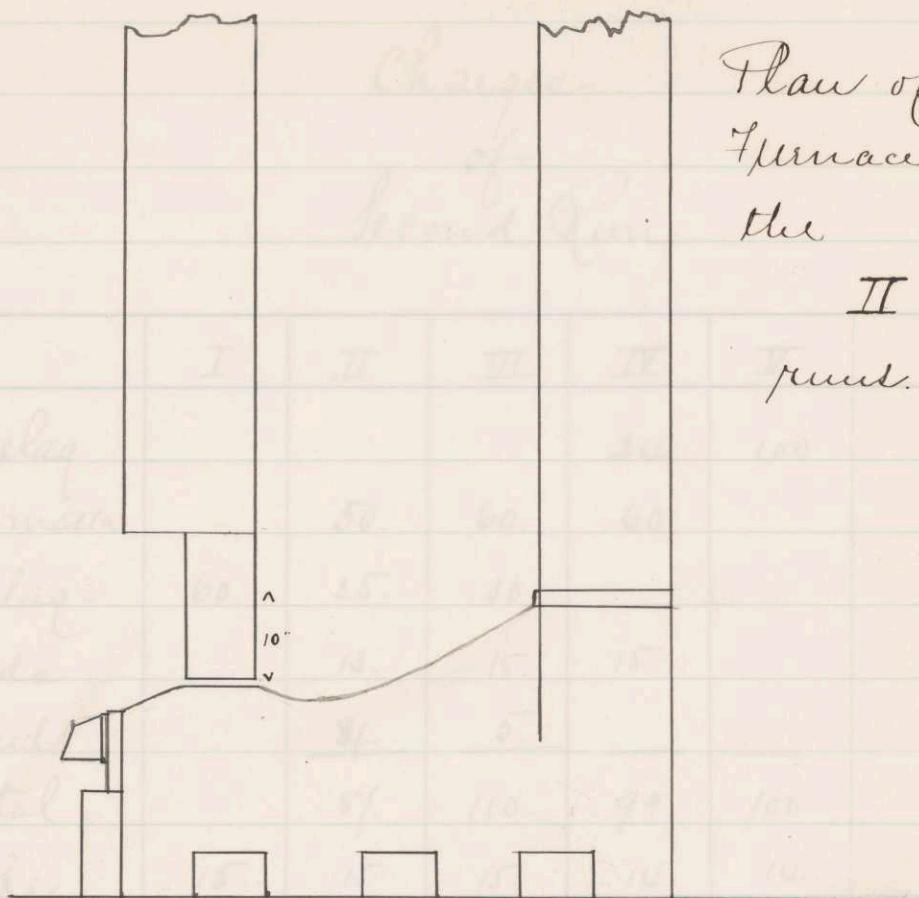
which was quite a reduction from the percentage obtained before coal wh. was

22.7%

showing that it had lost nearly one half of its S.

As the amount of foul slag and coke was so large, it was thought best to run with this percentage of S. only for a richer matte. The amount of S in the slag and coke, being so large as to preclude the hope of seeing Black Cu.

Plan of the
Furnace. for
the
II and III
period.



Tall R Run No. II
Charges
of
Second Run.

	I	II	III	IV	V	
Fr. slag				24.	100	pounds
road matter		50.	60.	60	"	
Fr. slag -	60.	25.	30	"	"	
Euds		13.	15.	15.	"	
Sand		<u>84.</u>	<u>5.</u>	—	—	"
Total		87.	110.	99.	100	
Coke	15.	15.	15.	14.	14.	"
Ratio	1:4.	1:6.	1:7.	1:4.	1:4.	

Tap Record II

Time	int.	buggy amt.			Note
Apr. 26.	4 P.M.	4 hods coke.	15 hour char.		
Apr. 27	4 P.M.	2			
9.40					blast on.
10.15					slag coming.
10.40					p.
10.45					t.
10.47		1.25			p.
10.54					t.
10.58		2			p.
11.10					t.
11.12		2.25			p.
11.23					t.
11.24		2			p.
11.35					t.
11.37		2.5			p.
11.48					t.
11.49		2.5			p.
12					t.
12.2		2.75			p.
12.13					t.
12.14		2.5			p.

Tap record II.

time	met.	buoy amt.			Notes.
12.23					t.
12.24		2.5			p.
12.23					t.
12.34		2			p.
12.46					t.
12.48		2			p.
12.59					t.
1.1		3			p.
1.13					t.
1.15		2.75			p.
1.26					t.
1.28		1.5			p.
1.38					t.
1.39		1.25			p.
1.48					t.
1.49		2.25			p.
1.58					t.
1.59		1.45			p.
2.8					t.
2.9		145			p.

Tap record II

time	int.	Beeq ₉₉ count			
2.19				t.	
2.20		1.45		p.	
2.30				t.	
2.31		2		p.	
2.41				t.	
2.42		1.5		p.	
2.51				t.	
2.52		1.75		p.	
3.				t.	
3.1		1.75		p.	
3.11				t.	
3.13		2		p.	
3.21				t.	
3.22		2.25		p.	
3.30				t.	
3.31		1.75		p.	
3.39				t.	
3.40		1.75		p.	
3.48				t.	
3.49		1.5		p.	

Tap record II.

time	int	boggy out.			
3.57				t	
3.59	7	-		p	
4.7				t	
4.9		1.5		p	
4.14				t	
4.18		1.5		p	
4.26				t	now flag. coming
4.31		1.5 -		p	
4.39				t	
4.41		2		p	
4.51				t	
4.57		1		p	
5.3				t	
					Furnace torn down at the end of run.

Fired record. II

time	set	charge	depth	ft.	Motes.
9.40					blast on.
10.8	7	I		2.5	
10.15	7	I		2.8	
10.20	5	I		3.1	
10.25	5	I		3.5	
10.30	5	I		3.5	
10.33	3	I		4.2	
10.49	16	II		4.5	
10.59	10	II		4.8	
11.9	10	III		5	flame gone
11.22	13	IV		4.75	
11.35	13	IV		5	
11.46	11.	IV		5.1	
12.2	16	IV		5	during high depth
12.14	12.	IV		5	pressure went up
12.24	10	IV		5	to 2.25 inches.
1-3	39	IV		4.4	flame after charging
1.16	13	IV		4.1	no "
1.36	20	IV		4.1	
1.51	15	IV		4.2	
2.14	23	IV		4.	

Feed Record II

time	inet	charge	depth	Notes.
2.26	12	IV	4.	
2.47	21	IV	4.	
2.58	11	IV	4.1	
3.21	23	IV	3.6	
3.23	2	IV	4.2	
3.39	16	IV	4	
4.3	24	V	3.7	
4.12	9		4.1	2 hds coke
4.20	8		3.5	
4.28	8		3.2	
4.44	16		2.8	
4.54	16		2.2	Almost all iron slag.
5.5	11.		9.	
5.10	5.		1.8	The furnace now only held coke. the slag being about all out. the front wall too out.
				Large "sow" found nestly fr.

This second run was done very successfully
very little foul slag being made.
The plan of having a small tap hole
wh. was plugged with a piece of
charcoal so cut as to fit the hole
quite snugly worked very nicely -
the tapping was done with a small
bar - the hole being made in the
charcoal. the result was, a
nice clean hole, through wh. the
material came in a quiet stream,
not any spattering to speak of.

The melt was picked over and
broken quite small. The slag of two
kinds - viz. I & II.

I slag was that obtained from the buggies
containing melt. and supposed to be a
little foul -

II slag. from the overflow of I and con-
sisted all slag - in some buggies a
very small cake of melt big found
in the bottom. — was thrown away -

The determinations from this run
II. in the various products are
 given as follows.

Matter.

$Cu.$ = in per cents. 30.58

$Fe.$ = " " 30.46

$S.$ = " " 23.05 and 22.7

SiO_2 = " " 1.80

Slag

$Cu.$ = in per cents. 1.2

$Fe.$ " 28.8

SiO_2 " 34.11

The amount of $Ca.$ in the Eude
 was not calculated.

Duplicates not made, except in case
 of SiO_2 . reason - shortness of time.

The product of run II being picked over and sorted, the weights of each were found to be as follows

Matt.	- in pounds	687.7
Seeds	- " "	225.5
Slag	I	648.3
	II	685.25
"Saw"		34.
Total amt of product.		<u>2280.75</u>

The time being so short, it was thought best to roast this matt in the reverberatory furnace - acc. the matt was crushed and passed through the roller, and reduced so that it would pass the $\frac{1}{16}$ th sieve - and was then ready to roast.

The fire was started the night before in the furnace -

The following calculation was given through
to ascertain the amt. per charge.

662.7 lbs metal to be heated.

Furnace I $4 \times 5 = 20$ sq ft. of surface

" II $3.5 \times 3.5 = 12.25$ " "

2 charges. $200 \frac{1}{2}$ I = 10 lbs per sq ft.

" " 131 " II = 13 " " "

3 charges 140 lbs I = 7 lbs per sq ft.

" " 81 " II 7 " " "

4 charges in large furnace

165.5 lbs per charge or 8 lbs per sq ft.

This last was decided upon and the
following arrangement made for shifts
Thursday May 2.

7-10 Am. 4 lbs made

10-4 I charge. -- Adams

4-10 II " -- Richards

10-4 III " -- Fred

4-10 IIII " { 4-7 Fred } { 7-9 Richards } { 9-10 }
Adams

This programme was carried out just as calculated. with the exception that two of the charges were one pound heavier. Some of the matt had been overlooked and was added in the second and third charges -

The amount of coal used for the roasting was.

Starting ~ 104.25 lbs.

roasting ~ 301.75 "

Total amt - 406.00

This was sampled and an analysis made for the amount of S^{18} that it contained -

It was not so low as we hoped but still the time was wanting to roast again

The determination for S in the roasted meat gave:

$$- \text{S} = 7.283 \% -$$

From these percentages were calculated the charges for the third run.

This run was expected to give us black co.

The charges are as follows for
Run III

	I	II	III	IV	V	VI
roast meat		30	60	60	60	13. pounds
sand		7.25	14.5	14.5		"
cinder		10	20	20	15.5	"
slag	60	10	20	20		80 " foul.
coke	15	14	28	26	28	28 "
iron	left out - too much iron					"
iron slag					40	"

Tape Record III

time	T+P	act.		Notes
10.15				slag coming.
10.25	p			fast plug-
10.30	t			
10.38	p	1.25		coming hot
10.44	t			cost 5 minutes to
10.49	p	1.5		start slag -
10.51	p			
10.56	t			hot and clean. layer bar.
11	p	1		
11.8	t			very hot
11.10	p	1.25		
11.19	p			
11.21	p	1.25		hot
11.29	t			
11.31	p	1		Some metallic Cu.
11.39	t			
11.41	p	1.25		" " "
11.49	t			
11.51	p	1.25		
11.59	t			
12.2	p	1.		

Tap Record III

Lines	T and P	aut.			Notes
12.10	t				
12.11	p	1			
12.19	t				
12.21	p	1.25			
12.30	t				
12.32	p	1			
12.44	t	"			
12.46	p	1			
12.57	t				
1.	p	1.25			
1.10	t				
1.11	p	1.5			
1.20	t				
1.25	p	1.5			
1.30	t				
1.32	p	1			
1.38	t				
1.39	p	1			
1.44	t				
1.45	p	1			

Tap Record III

time	T + P	aud.			Notes
1.51	t				
1.53	p	1.			
1.58	t				
1.59	p	1.			
2.8	t				
2.9	p	1.			
2.14	t				
2.15	p	.75			
2.20	t				
2.21	p	.75			
2.26	t				
2.27	p	.75			
2.35	t				
2.36	p	1.			
2.44	t				
2.45	p	1.			
2.52	t				
2.53	p	1.			
3.1	t				
3.2	p	1			

Tap Record III

time	TandP	Amt.		Notes
3.8	t			
3.9	p	1.		
3.18	t			
3.19	p	1.		
3.27	t			
3.28	p	1.		
3.37	t			
3.39	p	1		
3.47	t			
3.48	p	1.		
3.58	t			
3.59	p	1.		
4.8	t			
4.9	p	1		
4.16	t			
4.18	p	.75		
4.26	t			
4.27	p	1.		
4.34	t			
4.35	p	1.		

Tap Record III

Time	Tan ^o P	out.			Notes.
4.43	t				
4.44	b	1.			
4.53	t				
4.54	b	1.			
5.3	t				
5.4	b	1.			
5.16	t				
5.17	b	1.25			
5.27	t				
5.28	b	1			
5.36	t				
5.37	b	1			
5.45	t				
5.46	b	1.25			
5.54	t				
5.55	b	15			
6.3	t				Last tap at 6.55
6.4	b				Furnace torn down
6.12	t				
6.39	t				
6.40	b	1.			

Feed record III.

time	int	charge	depth	pressure 8 $\frac{1}{2}$ ft m.	Notes.
9		4 hoods coke		depth	1.5
9.40	40		1.5		blast on
9.55	15	1 hood coke	1.7		
10.5	10	I	1.9	4	flame strong.
10.12	7	I	2.4	4	
10.40	28	I	2.	4	
10.59	19	I	2.6	4.5	flame not free
11.20	21	II	2.4	4	
11.44	23	II	2.1		
11.54	10	II	2.1		flame after charge
12.1	7	II	2.1		" " "
12.14	13	II	2.5	12	no " " "
12.22	8			4	flame in slight liece flame
1.	38	III	2.3		
1.40	40	III	2.2	4	height kept and regulated throughout
2.14	34	III	2.4		
2.58	44	III	2.	flame	this run so as to just check the flame
3.40	42	III	2.2		use shovel of or put on at time
4.2	22	III	2.2		
4.35	33	IV	2.3		
5.14	39	III	2.3		

Feed Record III

time	wt	charge	depth	pres. 8 ^{10.07} in.	Notes
5.35	21			8	blast changed 4-8
5.41	20	V	2.7		to saw time
5.46	5			11	full blast
6.3	17	VI	2.7	1	all in -
6.9	6		2.9		2 hds coke about
6.21	15		2.4		30 lb
6.44	23		1.6		Furnace down

The same method of tapping and plugging was used, as in the second run, and gave great satisfaction.

Very little foul slag was obtained at the end, as all that had accumulated during the run was set in toward the close of the run.

Several times in drawing out the tapping bar, it was found to be covered with metallic Cu. showing that some might be expected in the buggies.

The products were sorted over and three products were obtained viz. Slag - Matt and black Cu

The amounts of each are given below

Matt - in pounds - 305.25

Slag - " " - 1202.75

Black Cu - " " 30.25

Total amount - 1538.25

owing to the shortness of the time,
the amount of Cu. and the various
constituents of the matt and slag
could not be determined.

But from the appearance of the
matt - it must be very much richer
than before. and acc. to the report
of the Professor in charge - probably
between 65 and 75 per cent of Cu. was
present in the matt.

The slag was very free from matt
but few bubbles being found that
were foul.

A very low pressure was kept
up during the whole of the
run. it being regulated by
a gate made for the purpose

- Summary -

The roasting in the kiln, has been explained in the first part of this paper, and it need not be mentioned again.

The first melt for matt. was not at all satisfactory. the form of the furnace did not give good results. after the lower tap hole had been lost, the slag and matt. overflowed together into the buggies.

This gave us a large amount of foul slag - which interfered with the next melt. the amount of Cu. in shape of sulphides in this slag - which could not be very well roasted over so great that even if the matt were roasted nearly dead. this sulphur, would hinder very much the formation of black Cu.

So the matt was roasted and the whole run down for a rich

mass. great pains being taken to prevent the accumulation of ends and fine slag -

Accordingly in the second run. a new form of furnace - that of tap and plug - using charcoal for plug - and by working carefully very little fine slag was made.

This form of tapping - with charcoal gave such general satisfaction that it is to be recommended for future use - its great recommendation is. the "clean" tap hole it makes - the bar being driven into the charcoal and solv. drawn out. leaves a nice clean opening for the melted material - very little wear of the steep is noticed when using this form of plug - that is, the steep of the run is not worn and cracked, leaving the run in a bad condition for the flow of the mass and slag -

Accordingly in the third run, the same form of furnace was used. The ore in this case, had been run the rolls preparatory to the roasting in the reverberating furnace, and was in a finely divided state, with considerable dust.

This run was very successful indeed the furnace worked well.

A low pressure was kept up, not over half an inch, all the time until the red color it was put on in full force, to clean out the furnace.

The effect was - there was no loss up burning of dust, and the reduction of copper took place with less danger of the reduction of Fe, wh. would enter into the black Cu. and spoil the metal. The little black Cu obtained - showed the effect of the care and precaution.

See through the stem runs. no Ca.
was used as a flux.

If had customary in past year to
use Ca. but from observations
made it was concluded best "not"
to use any now.

The effect of Ca. in the flux is
to blow out the Fe. wh reduces
and finds its way into and with
the Cu.

As has been mentioned, the black
Cu obtained, showed clean with
"little" evidence of Fe.
Fe. is present but not in such
great quantities as in the runs
of previous years.

I regret exceedingly that I was
not able to make an analysis
of the black Cu. obtained.

Methode used for the determination of
of the constituent parts of the
products of the stone pens. and of the
ova.

Cu - by means of battery. HNO_3 . sol

Fe - . filtration with bichromate

S - heat with fuming HNO_3 . evaporate
to dryness. to get rid of free HNO_3 .
precipitate the Fe and other things by
means of Na_2CO_3 . make acid and
precipitate with $BaCl_2$ - care must be
taken to get rid of the free HNO_3
as it interferes with the precipitation
of the Barium sulphate - let stand
over night -

Si. treat with acids to decompose as
much as possible. free with Na_2CO_3 .
evaporate the "whole." { acid. dol. and the
carbonate sol. } after being made acid -
heat to make Si insol. add acid to
dissolve out the other constituents.
and weigh as SiO_2 .

without roasting the ore, the first time as there was plenty of spare time the color method of determination for Cu - in small amount, was tried. This method is only applicable to slags or such materials as do not contain more than 2 per cent of Cu.

It depends on the blue color produced when ammonia is added in excess to an acid sol. of copper. The intensity of the blue color depending on the amount of Cu. present.

Two ways that are easy of manipulation are given - descriptions of wh. follow.

Take a gram of pure Cu and dissolve in HNO_3 - } a gram not necessary but I took an even lot. as the calculation would be much simpler. } add an excess of Ammonia - a deep blue color results - now make up to 1000 c.c.

First method.

In this solution of Cu - wh. has been made up to 1000 cc. - 10 cc is equivalent to .1 of a per cent., 20 cc " "

" .2 " " 30 cc " "

" .3 " " 40 cc and so up.

I took a number of larger test tubes and commencing with 40 cc or .4 % arranged a series of tubes containing

40. 50. 60. 70. 80. 90. 100. 120. 130. 140. 150. 160. c.c's

I now filled them with distilled water, all up to the same height, and corked them up tight.

I now had a series of bottles containing a solution of Cu - colored blue, of different intensities.

These were the standard solutions.

I now took a slag containing a small amt. of Cu. .8 of a per cent. treated with acid - and then with Ammonia made up to the same height with water. as the standard and then compared with the standards.

At first. I could not do much at it
but in a short time - it was quite
an easy matter to locate the test tube
containing the sol. at the standard bottle.
I now tried some unknowns and
succeeded very well.

The whole operation was an
experiment, and though I tried to get
perfectly cylindrical test tubes. probably
they were a little out.

But even with that trouble the experiment
was a grand success.

It would have been better to have
had larger test tube or something that
would hold much more solution
as - the whole bulk of the solutions
that I had did not probably exceed
200 cc and the colors were too
intense - it being hard to distinguish
them.

The second method was that of using bottles. The per cent being determined by the amount of water that has to be added to make the unknown solution of the same color as the standard. The standard solution may be taken from that already made. Take .50 cc. then make up in "futab" vessel to 200cc by adding 150 cc distilled water.

The unknown solution is put into a like vessel - an *Pestamonia* added - then add water till the colors of the two bottles are alike.

Did not obtain good results - it being very hard to tell any difference of color even if two cubic centimeters of water over and above the amt. desired was added.

Did not try any unknowns for the above reason.

The first method much the best and is capable of being ^{put} to good use. Practice would make this method very valuable.

Analysis of Ore.

| Elements. | Southworth. | Baile. |
|------------------|-------------|---------------|
| Sulphur | 33.46% | 33.97% |
| Silica - resid. | 6.440. | 5.45.. |
| " sol. | 1.50. | 2. " |
| Copper. | 11.25. | 11.29.. |
| Iron. | 46.47. | 46.76.. |
| Nickel & Cobalt. | 1.66" | 1.27
trace |
| Alumina | | " |
| Magnesia | | " |
| Alkalies | | " |
| Calcium | | " |
| Nickel & Cobalt | | " |

Analyses of Roasted Ore
 and
 Malls for Sulfur.

| | |
|---------------------------------|-----------------|
| 1 st roasted in kiln | 18.51% |
| 2 nd " " " | 6.21% " 6.52% ? |
| 1 st " of mott. | 13.17% |
| 2 nd " " " | 7.28% in R.F. |

Raw, I.

Matt

| | |
|---------|--------------------|
| Copper. | 21.440% .. 21.321% |
| Iron | 53.37% |
| Sulphur | 22.70% |
| Silica | .997% |

Slag.

| | |
|----------|-------------|
| Copper. | .876% .895% |
| Iron. Fe | 52.% |
| Silica | 35.65% |

Foul-slag-

| | |
|---------|----------------|
| Copper. | 4.83% .. 4.77% |
|---------|----------------|

Euds

| | |
|--------|----------------|
| Copper | 4.48% .. 4.43% |
|--------|----------------|

Run II.

Mall.

| | |
|---------|----------------|
| Copper | 30.58 % |
| Iron | 30.46 " |
| Sulphur | 23.05 - 22.7 " |
| Silica | 1.80 " |

Slag

| | |
|--------|---------|
| Copper | 1.20 % |
| Iron | 28.80 " |
| Silica | 34 11 " |

-Fuel used-

| Two roasters in Kile,
used | Coke | char. |
|--|---------|--------|
| | 450.75 | 78.75 |
| "Coke used in the
1 st smelt" | 645. | 16 |
| "Coke used in the
2 nd smelt." | 587. | 10. |
| "Coke used in the
3 rd smelt" | 563. | 10. |
| Total lbs = | 2245.75 | 114.75 |

"Coal used in the R.F.
for starting"
"for roasting"
Total pounds = _____

| | |
|--------|--|
| 104.25 | |
| 301.75 | |
| 406.00 | |