A practical study of complex sulphide separation using collectorless and nitrogen flotation.

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I. ABSTRACT

This thesis explores the potential for (1) collectorless flotation and (2) the use of nitrogen in complex sulphide flotation. Several ores from the Canadian Shield are used.

Various responses to collectorless flotation are observed. Chalcopyrite can float without collector, particularly in ores from Mattabi Mines. Galena from Mattabi ores also floats though more slowly than chalcopyrite. Galena flotation is attributed to surface oxidation while the mechanism for chalcopyrite collectorless flotation is not fully understood. A process is proposed that includes collectorless flotation and may lead to improved copper, lead separability. No collectorless flotation of pyrite and zinc-bearing minerals is observed.

Nitrogen, used as the carrier gas, promotes pyrite flotation in all ores tested. This is used to selectively remove pyrite after bulk flotation and before zinc flotation. Some zinc-bearing minerals report to the pyrite concentrate. Pyrite floated in nitrogen can, however, be depressed again in air enabling the zinc to be selectively floated after copper activation. The proposed process improves zinc rougher concentrate grades by up to 50% at similar recoveries. This is attributed to an overall reduction in the exposure of pyrite to collector and copper sulphate. The flotatic of pyrite in nitrogen is linked to the removal of dissolved oxygen. It is suggested that this may block a galvanic interaction between pyrite and sphalerite which in an oxygenated environment depresses pyrite.

II. RÉSUMÉ

Cette thèse examine le potentiel de la flottation sans collecteur et de l'utilisation de l'azote pour la flottation des sulfures complexes. On utilise plusieurs minerais du Bouclier Canadien.

La flottation sans collecteur donne des résultats qui dépendent beaucoup du minéral flotté. La chalcopyrite flotte parfois sans collecteur, surtout avec échantillons provenant des mines Mattabi. La galène des mines Mattabi flotte également sans collecteur, quoique plus lentement que la chalcopyrite. On attribue la flottation de la galène à son oxydation en surface; on ne peut toutefois pas proposer d'explication satisfaisante pour la flottation de la chalcopyrite sans collecteur. Nous prosposons un schéma de traitement qui fait appel à la flottation sans collecteur, et qui pourrait améliorer la séparation de la chalcopyrite et de la galène. On n'a pas pu flotter sans collecteur la pyrite ou les minéraux zincifères.

L'azote, lorsqu'utilisé comme gaz transporteur, favorise la flottation de la pyrite pour tous les minérais étudiés. Cette technique permet l'extraction de la pyrite, après la flottation de la chalcopyrite et de la galène, mais avant celle du zinc. Une fraction des minéraux zincifères se retrouve dans le concentré de pyrite. Cette pyrite, flottée avec l'azote, peut toutefois etre déprimeé avec l'air, pour ainsi flotter le zinc sélectivement après activation au sulfate de cuivre.

Ce procédé permet d'obtenir des concentrés de dégrossissage dont la teneur est de 50% plus élevée, à récupération équivalente. On explique cette amélioration par une réduction du contact entre la pyrite, d'une part, et le collecteur et le sulfate de cuivre, d'autre part. La flottation de la pyrite par l'azote est liée a l'élimination de l'oxygène dissout dans la pulpe, ce qui, croyons-nous, empêche l'interaction galvanique (entre la pyrite et la sphalérite) qui déprime la pyrite, en présence d'oxygène.

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1. INTRODUCTION

1.1 BACKGROUND AND THESIS OBJECTIVES.

Froth flotation has been used to concentrate sulphide minerals for 80 years and organic-based collectors such as xanthates have been used since 1925 (26,27,96).

floatability of a sulphide mineral is dependent on the hydrophobicity of its surface. A few sulphide minerals such as molybdenite are naturally hydrophobic because of their specific lattice structures, and will float without any pre-treatment This is usually referred to as natural floatability (26). Other sulphide minerals exhibit self-induced floatability - they can be rendered hydrophobic by some form of pre-treatment (22,25-6,36,96,107-8). Most sulphides, however, need to adsorb a collector onto the mineral surface before they become hydrophobic. Electro-chemical mechanisms are believed responsible for the interaction between collector and mineral, and electrochemical control may be the useful means of controlling mineralcollector interactions.

This thesis assesses the possible application of collectorless flotation towards achieving selective flotation of minerals from some Canadian Shield ores. Using the same ores, the thesis then considers the ability of nitrogen gas to alter the electrochemical properties of the pulp, and thus mineral-collector interactions.

1.1.1 Collectorless (self-induced) flotation

In the last 30 years considerable progress has been made toward developing an understanding of self-induced flotation (102) Researchers, however, have generally used pure minerals and perhaps for this reason it has found little commercial application. One of the objects of this thesis is to assess the potential for collectorless flotation of a number of ores from the Canadian Shield.

1.1.2 Nitrogen flotation

In 1953, Salamy and Nixon(87) proposed an electrochemical theory of flotation and this theory has become increasingly accepted as more is learned about mineral-collector interactions (105). One of the consequences, as described by Woods(105) is the importance of pulp potential in mineral-collector interaction. Recently workers have attempted to use pulp potential to develop new approaches to achieve and control selective flotation (30,32,34,81-2,103). Because of nitrogen's chemical inertness, flotation in nitrogen could be a particularly elegant means of adjusting the pulp potential. Nitrogen has already found significant use on copper/molybdenum separation (13,73) but has not yet found any other commercial use. The effect of nitrogen on the flotation of complex sulphide ores of the Canadian Shield will be assessed in this thesis.

1.2 STRUCTURE OF THE THESIS.

This thesis includes literature reviews of collectorless flotation and the role of pulp potential and dissolved oxygen in flotation. This is followed by a description of the operating practice of the three plants from which ore was sampled for the testwork. A review of the current use of nitrogen and gases other than air in flotation is included, together with a section on the supply o one available for nitrogen production.

The test procedures for all three ores are outlined in section three. The results section, section four, is sub-divided into the testwork on each individual ore. Further sub-divisions describe aspects of collectorless flotation and nitrogen flotation.

In section 5, the results are discussed and processes proposed which include collectorless flotation and pyrite flotation in nitrogen. With numerous mechanisms already available to explain collectorless flotation, no new mechanism is proposed in this thesis. The promotion of pyrite flotation with nitrogen is explained by elimination of a galvanic interaction between pyrite and sphalerite, which in the presence of oxygen serves to depress pyrite. An economic analysis of the use of nitrogen is included.

Sections 6 and 7 summarise the conclusions drawn and suggest directions for future work. Following the list of references, the appendices detail the results from the tests reported in the thesis.

2. GENERAL REVIEW OF BACKGROUND TO THE PROJECT.

2.1 REVIEW OF RESEARCH INTO COLLECTORLESS FLOTATION AND THE USE OF NITROGEN

2.1.1 Collectorless flotation.

Some of the earliest processes developed in flotation used the collectorless hydrophobicity of minerals. Work on Broken Hill ore in Australia, as described by Louis (57), used carbonic acid to affect self-induced flotation of sphalerite and collectorless flotation of galena was occasionally used up to 1920.

With the development of xanthates, collectorless flotation of minerals sulphide became largely forgotten. Effective collectorless flotation was considered limited to orpiment and molybdenite. Gaudin (26) explains these exceptions through their crystal lattice structure . However Sutherland and Wark (96) questioned this in 1955, by noting that galena would occasionally respond to collectorless flotation, and in doing so sparked a controversy that is still gaining momentum 30 years later. They summarized work by numerous authors who had all found that galena could float without collector, but could not agree on the mechanism and particularly whether the surface needed to be clean or oxidised. In 1960, Rey and Formanek (80) found that sphalerite would float without a collector if ground in a ceramic mill . More recently, Kocabag and Smith (49) confirmed this observation, making an in-depth study of the effect of grinding media in flotation .

Since then most of the work on collectorless flotation has concentrated on chalcopyrite. Plaksin (69) used chalcopyrite to show that hydrophobicity of sulphide minerals was possible through exposure to air. Lepetic's work (55), in 1974, showed that chalcopyrite can be floated without collector following dry grinding, achieving superior grades and recoveries than through the conventional route of wet milling and collection. Both explained their observations by suggesting that the mineral surface was adsorbing molecular oxygen, leading to a de-hydrated and hence hydrophobic surface in flotation.

An exhaustive study of collectorless flotation was performed by Finkelstein and co-workers(22) in 1975. They checked the collectorless floatability of a wide range of ores finding some chalcopyrites, pyrites, galenas, stibnites and all copperactivated sphalerites were floatable without collectors.

While that work probably convinced most researchers that effective collectorless flotation was possible, the responsible mechanism has become an object of increased interest and often considerable disagreement over the past ten years. The controversy was intensified when Heyes and Trahar (36) identified in 1977 that collectorless flotation was only possible under oxidising conditions, and Yoon (107) found, 4 years later, that chalcopyrite would float naturally following addition of a reducing agent, sodium sulphide.

While an increasing amount of evidence seems to endorse Heyes and Trahar's approach (102) that oxidising conditions are required

the exact role of oxidation/reduction in collectorlass flotation is still an open question.

Finkelstein and Woods agree with Heyes and Trahar (12,22,25). But while agreeeing that oxidising conditions are necessary for formation of a hydrophobic surface on the chalcopyrite there was disagreement about the specific product of oxidation that imparts this floatability. Finkelstein's work (22) concluded that elemental sulphur played no role in collectorless flotation. Gardner and Woods (25), however, proposed an oxidation reaction to explain the collectorless flotation of chalcopyrite, that included the formation of elemental sulphur:

$$CuFeS_2 + 3H_2O -> CuS + S + Fe^{3} + 3e$$

Later, Woods and Buckley (12), using X-ray Photoelectron Spectroscopy failed to identify elemental sulphur on the chalcopyrite surface concluding that the oxidation product was some form of copper poly-sulphide. Trahar (99) in 1983 remained consistent with the line of thinking that oxidation of the chalcopyrite surface was needed to render the mineral surface naturally floatable, though he made no conclusions as to the actual species responsible for this hydrophobicity.

Yoon's (107-8) observations in 1981 that adding sodium sulphide, a reducing agent, to the grind rendered both chalcopyrite and copper-activated sphalerite floatable, was in contrast to the other investigators. His work, on Kidd Creek ore amongst others, identified that if sodium sulphide was added to the mill, the

resultant grade/recovery relationship from collectorless flotation of chalcopyrite and sphalerite was similar to that using collectors. He suggested that sodium sulphide helped dissolve superficial oxidation products leaving a clean particle surface.

More recent work by Luttrell and Yoon (58-9) illustrated again the advantage of using sodium sulphide in collectorless flotation of chalcopyrite. They concluded that freshly ground chalcopyrite would not float naturally under reducing conditions and that oxidised chalcopyrite would also not float naturally. Chalcopyrite, however, treated with sodium sulphide under oxidising conditions would float naturally.

In 1984, Heyes and Trahar (37) concluded that pyrite could also be floated following oxidation and treatment with sodium sulphide. They concluded that sodium sulphide causes sulphur to form on the pyrite surface and this sulphur is responsible for the minerals hydrophobicity and natural flotation characteristics.

Only a thin sulphur layer can, however, be formed on chalcopyrite because of the relative instability of elemental sulphur at common flotation potentials. This indicates that elemental sulphur formation is not the mechanism behind natural flotation of chalcopyrite. Nevertheless, Walker and co-workers (102) observed that chalcopyrite could be made floatable by electrodeposition of sulphur at potentials well below those permitting the oxidation of the mineral. In short, the mechanism

behind the collectorless flotation of chalcopyrite remains unclear (102).

contrasting theories behind the collectorless flotation of chalcopyrite may illustrate the problem of trying to explain it on a pure mineral basis. Mineralogical texts describe that chalcopyrite can contain any of eleven elements replacing the iron, copper and sulphur (16).Furthermore the copper:iron:sulphur ratio can vary (one single specimen gave copper contents varying from 25.8% to 30.7%), and measurement of X-ray Absorption edges have shown that the copper can exist the +1 or the +2 states, the iron then existing in the +3 and the +2 respectively. Analysis through cryoscopic states electrolytic methods .ave also shown that both metals can be divalent (16,63,74). Such variations in the mineral must lead to different reactions and different reaction mechanisms.

The mechanism behind the collectorless flotation of pyrite and galena seems to be more clearly understood. Rao and Finch (77) following sufficient oxidation, showed that, pyrite from Brunswick Mining ore can float without collector, albeit weakly, and this could be used to improve subsequent zinc-pyrite selectivity. Voltammetric data suggests the initial product from oxidation of the pyrite and galena surfaces, is the polysulphide ion, though as further oxidation occurs elemental sulphur forms on the sumface (37). The actual species present on the mineral surface is probably related to the degree of oxidation and the relative kinetics of the reactions involved, but with sufficient oxidation a relatively thick sulphur layer can be formed on the surfaces of pyrite and galena.

This is particularly relevant as Walker and Richardson (102) observed that the hydrophobicity of the mineral surface is related to the thickness of the elemental sulphur formed on the surface. Such conditions seem to have occurred on a number of the pyrite-rich gold tailings dumps in South Africa which often respond readily to collectorless flotation (67).

While researchers do seem to agree on a mechanism behind the collectorless flotation of pyrite and galena, variations for mineralogical reasons are still possible (galena, for instance can have a eutectic intergrowth of up to 30% iron sulphide(16)).

This thesis will check the effect of collectorless flotation on some Canadian ores, and will highlight the variations in response to collectorless flotation possible in some similar pyritic ores of the Canadian Shield.

2.1.2 ROLE FOR NITROGEN: THE ELECTROCHEMICAL THEORY OF FLOTATION.

The use of nitrogen in flotation was conceived as a logical progression from the electrochemical-based theory developed over the last thirty years to explain mineral-collector interaction.

Only in recent years has significant progress been made in the understanding of collector attachment to minerals. While a basic understanding of the role of collectors was available by the 1950s, little was known about the mechanism of collector-mineral attachment (26-9).

In the 1950s it was largely accepted that collector-mineral reactions largely involved ion adsorption (26) through a relatively simple series of well-defined reactions, although other theories had been suggested as outlined by Gaudin and more recently Leja (26,54).

In 1952, Salamy and Nixon (87) proposed an electrochemical mechanism involving the simultaneous anodic oxidation of collector and cathodic reduction of oxygen, on the mineral surface. This has become increasingly accepted as more is learned about the mechanism of the various mineral sulphide-collector-oxygen systems, and of the variety of surface species produced from the reactions (105).

Sulphide minerals are semi-conductors and can thus act either as a source of electrons or an acceptor of electrons. They act as the former in the reaction with the collector, and the extra electrons are conducted away and react with oxygen (see Figure

2.1). The reaction with the collector (the anodic oxidation reaction resulting in spare electrons) may be of two types:

formation of dixanthogen: $2ROCS_{2}^{2-} \longrightarrow (ROCS_{2})_{2} + 2e-\dots <1>$ or reaction (chemisorption) with the mineral itself: $PbS + 2ROCS_{2}^{2-} \longrightarrow Pb(ROCS_{2})_{2} + S + 2e-\dots <2>$

The surplus electrons would then reduce oxygen, effectively simultaneously, in the following cathodic reduction reaction:

Which of the two anodic reactions predominates depends on the mineral, the collector and the pulp potential. Investigations by Goold and co-workers (32) in the early 1970s led to a correlation between minerals, collectors, and potentials at which the reactions start to occur (rest potentials) and surface products, as shown in Table 2.1. The table shows the effect of the rest potential on the final reaction product. If the rest potential is high (above the reversible potential for the oxidation of xanthate to dixanthogen) then the product is dixanthogen. If it is lower than the reversible potential, then the product is the metal xanthate. The one exception is covellite and xanthate. This is explained in terms of the reaction of dissolved cuprous ions from the mineral surface with the xanthate (32,105).

PbS + 2ROCS₂ - Pb(ROCS₂)₂ + S + 2e
collector-mineral interaction

ROCS - H+

2H₂O

O₂ + 2H+ + 2e- - 2H₂O

oxygen reduction reaction

FIGURE 2.1 - Principle of collector oxidation and oxygen reduction on galena surface.

TABLE 2.1: The reaction product and starting (rest) potential of the anodic reaction of various mineral-collector combinations. (105)

mineral	collector	rest	brogact
		potential	
		vs S.H.E.	
pyrite	KEX	0.22V	dixanthogen
arsenopyrite	KEX	0.22V	dixanthogen
pyrrhotite	KEX	0.217	dixanthogen
chalcopyrite	KEX	0.14V	dixanthogen
rev. potential	for oxidation	to dixanth	ogen - 0.13 V
covellite	KEX	0.05V	dixanthogen
bornite	KEX	0.06V	metal xanthate
galena	KEX	0.06V	metal xanthate
pyrite	DtC	0.475V	disulphide
rev. potential	for oxidation	to disulph	ide - 0.176V
covellite	DtC	0.115V	metal DtC
chalcopyrite	DtC	0.095V	metal DtC
galena	Dtc	-0.035V	metal DtC
bornite	DtC	-0.045V	metal DtC
chalcocite	DtC	-0.155V	metal DtC

note: KEX - Potassium ethyl xanthate (0.000625M at pH 7)

: DtC - Sodium diethyl dithiocarbamate (100.ppm at pH 8)

Generalisations are always dangerous. Gaudin (28) identified metal xanthates as well as dixanthogen on the surface of pyrite and it is likely that in all cases both reactions occur, although one will tend to predominate over the other. Most of the evidence in the past twenty years has suggested that dixanthogen is the active species in pyrite flotation (21,22,61).

Pyrite floats in air most effectively at potentials above that of dixanthogen formation (105). This suggests that by maintaining the pulp potential below the rest potential for dixanthogen formation on pyrite, or preferably below the reversible potential for xanthate oxidation to dixanthogen, pyrite could be depressed. This is an example of electrochemical control of flotation which has commanded much attention at the U.S. Bureau of Mines in recent years (30,81,82,103).

Investigators at the U.S.B.M. have shown that, with single minerals, the flotation of chalcopyrite, pyrite, bornite and chalcocite is strongly dependent on pulp potential (82). When mixed beds are used interactions between the minerals tend to distort the result. For example, in a chalcocite-pyrite mixture, pyrite is floated at different pulp potentials than in the case of pyrite alone possibly through activation by copper ions dissolved from the chalcocite surface (Figure 2.2)(30).

While the idea of selective flotation through control of the pulp potential has possibilities, external control of the potential through contact with electrodes, as being attempted at the U.S.B.M., pose severe practical difficulties on a large scale.

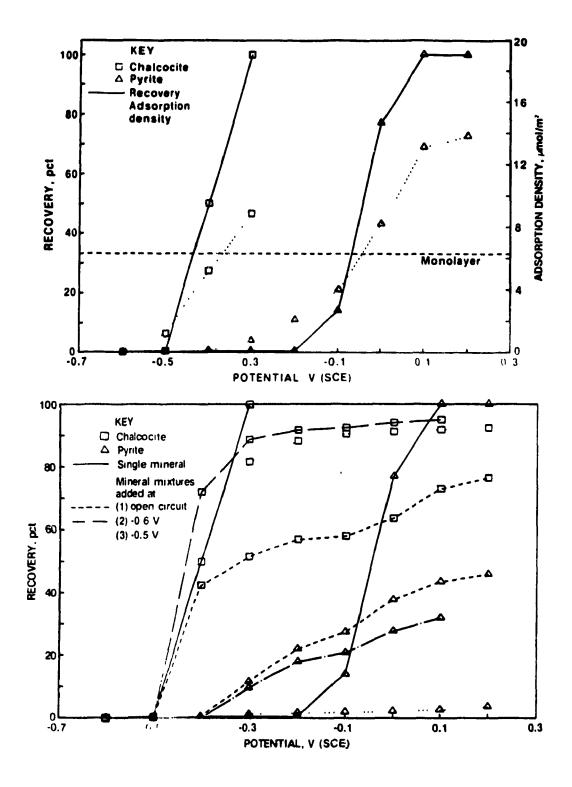


FIGURE 2.2 - Collector adsorption densities and flotation recoveries of pyrite and chalcocite as single minerals and combined.

The first graph shows that if flotation was to take place at -0.2 to -0.3 volts, chalcocite would float and pyrite remain depressed, based on their floatabilities as single minerals. The second graph shows how the flotation recoveries of the minerals become less potential dependent when added together.

Chemical control of potential has possibilities (and may already be practiced albeit unknowingly). Outokumpu have run pilot tests using control of pulp potentials, as against pH. This testwork used sulphuric acid for potential control. It resulted in savings in reagent costs (particularly xanthate and frother where up to 50% savings were reported), and an improvement in economic returns from concentrate production. So far tests have been run on copper/nickel ores and gold ores. Whether the technique can be applied to more complex ores such as the pyritic copper/lead/zinc ores of the Canadian Shield remains to be seen(34).

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Development of improved process control methods is only one consequence of recent progress in flotation electrochemistry. Many reagents used in current circuits may activate or depress by changing the pulp potential. If this is the case then a reagent that adjusts the pulp potential without interfering with any other aspect of flotation chemistry could be extremely useful.

Nitrogen may be such a reagent. Nitrogen gas is currently used in the flotation separation of molybdenite from chalcopyrite (see Section 2.3)(73). Nitrogen gas as the gas phase in flotation would reduce the pulp potential. It is also an inert gas and should thus not interfere directly with any chemical mechanisms. In, for example, pyritic ores containing zinc, zinc-pyrite selectivity could be improved by lowering the potential below the rest potential for pyrite flotation (76-7). One of the objects of this thesic is to extend this idea one step further, and incorporate nitrogen into a complex copper/lead/zinc flowsheet using a currently processed ore.

Nitrogen will also purge dissolved oxygen out of the pulp. This may reduce flotation through blocking Reaction 3. However the role of oxygen in a flotation pulp is not fully understood and its removal could affect other mechanisms which contribute to mineral surface hydrophobicity. One such mechanism is galvanic interaction between different minerals, as described by Kocabag and Smith in 1985 (46). Figure 2.3 illustrates the mechanism they proposed for pyrite, a particularly noble sulphide, and another sulphide mineral. The two results relevant to flotation are:

- the production of elemental sulphur on the mineral acting as the anode (and possible resultant collectorless flotation), and,
- the formation of hydrophilic ferrous hydroxide on the pyrite surface, making it less floatable.

Dissolved oxygen is fundamental to such a mechanism so use of nitrogen could block it by removing the oxygen. This would render the pyrite more floatable, counter-acting the effect of inhibited collector-mineral interaction.

Further consideration will be given to the possible role of galvanic interactions in depressing pyrite in the discussion section (Section 5.2.1).

sphalerite

- less noble

pyrite

- more noble

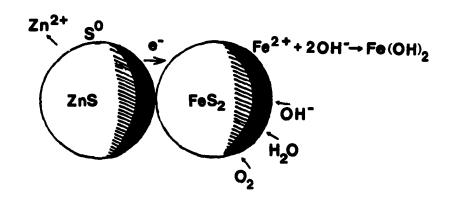


FIGURE 2.3 - Proposed galvanic interaction between pyrite and sphalerite, leading to an oxidised sphalerite surface and a hydrophilic pyrite surface.

2.2 REVIEW OF ORE MINERALOGY AND CURRENT METALLURGY AT THREE CANADIAN OPERATIONS.

2.2.1 Brunswick Mining and Smelting (Figure 2.4).

The Brunswick #12 ore deposit is one of the largest complex sulphide deposits in the Canadian Shield (93). It contains 100 million tonnes of complex sulphide ore, averaging 9.0% zinc, 3.5% lead, 0.3% copper and 98 grams per tonne silver.

The deposit, situated 27 km south west of Bathurst, N.B. of sedimentary and volcanic rocks which metamorphised and deformed during Ordovician and Devonian times (53,86,95). The ore is 80% sulphides, of which almost 60% is pyrite. The principal zinc minerals are sphalerite sulphide, low in iron) and marmatite (zinc sulphide, high in iron), lead is mineralised as galena (lead sulphide), copper as chalcopyrite (copper, iron sulphide) and silver as tetrahedrite (copper, silver, antimony) sulphide and other sulphide minerals. other minerals occurring the are pyrrhotite, arsenopyrite, marcasite, boulangerite, stannite, cassiterite and pyrargyrite.

While high-grade bands contain relatively coarse-grained values $(100-1000\mu\text{m})$, low grade bands are much more finely disseminated. The economic optimum grind size is currently 65%, -37 μ m for primary grinding, though it is significantly finer in the regrind circuits.

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KEY TO PRINCIPAL REAGENT POINTS:

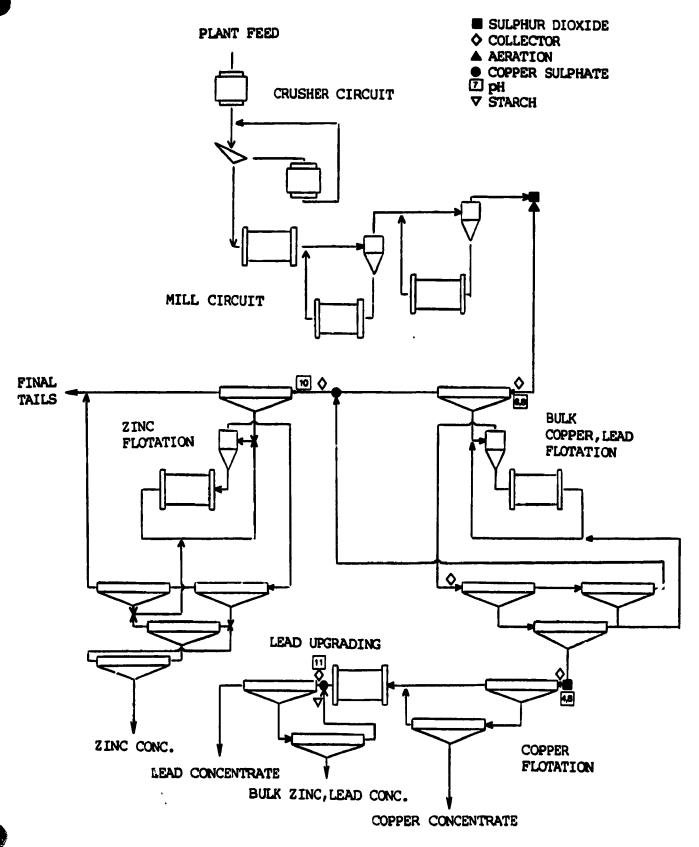


FIGURE 2.4 - Brunswick Mining general plant flowsheet.

Mine production has been entirely from underground since 1980. The mine uses a mechanised cut-and-fill method which replaced open stope mining in 1971. Ore is hoisted using 15 ton skips through numbers 2 and 3 shafts from three primary crushing units, roughly 1000 m underground. The mine currently extracts 10,250 tonnes of run-of-mine ore daily (68).

The ore is crushed to minus 15cm underground using 2 Traylor type-H jaw crushezs and one Allis-Chalmers gyratory crusher. Once on the surface it is crushed further, to minus 1.2cm in each of two plants. A secondary 1.7m Symons standard cone crusher operates in open circuit, the crusher product sized on a 1.2cm vibrating screen. Oversize is crushed in a tertiary 1.7m shorthead cone crusher, in closed circuit with the screen.

Brunswick Mining has three grinding lines (17). Numbers 1 and 2 (2900 tonnes/day each) receive ore from crusher plant number one. They consist of a 3.2m by 4.3m primary rod mill with 3.2m by 4.0m secondary and tertiary ball mills. Number 3 grinding section (4200 tonnes/day) receives ore from number 2 crusher plant and consists of a 3.8m by 4.9m primary rod mill with 3.8m by 4.6m secondary and tertiary ball mills. 76mm rods (and 100mm in line 3) are used for primary rod milling, and 32mm Manmet slugs are used for ball milling.

Two stage cyclones size the mill products. The primary cyclones are 25cm diameter Krebs cyclones. They classify the rod mill discharge and are in closed circuit with the secondary ball mill. The secondary cyclones, 15cm Krebs cyclones, are in closed

circuit with the tertiairy ball mill.

Soda ash is added to maintain a pH of 8.8 in the grinding circuit together with 150 g/tonne sulphur dioxide. The role of sulphur dioxide is to depress sphalerite and pyrite in the bulk flotation, although its exact action is not known. Results from work at Brunswick Mining by Hill appear to confirm this philosophy (38). Theoretical aspects of sulphur dioxide depression will be considered at a later stage (Section 2.3).

The secondary cyclone overflow is fed to an aeration stage with 25 minute retention time. Aeration further depresses pyrite flotation. After flotation two collectors are added consisting of (1) plant xanthate containing sodium isopropyl xanthate (80%) and amyl xanthate (20%) at 30 g/tonne and (2) Cyanamid AEROFLOAT dithiophosphate (241) added at 20 g/tonne.

Bulk flotation is carried out using 8 m3 Outokumpu cells, producing a concentrate containing 18% combined lead and copper. The bulk rougher concentrate is then re-ground to 81% minus 38 um in a 3.2m x 4.0m ball mill.

Two stages of cleaning produce a concentrate containing 29% lead, 2.5% copper and 700g/tonne silver. The bulk cleaner circuit is open, the thickened tails from each stage combining with the bulk scavenger tails to form the zinc flotation feed.

The bulk cleaner concentrate is treated with activated carbon to remove dissolved organics (e.g excess collector), though the carbon may also remove some of the collector coating from the

galena surface (38). It is also treated with starch which helps to depress the galena. Sulphur dioxide is then added to the pulp which further depresses the galena, and maintains a pH of 4.8 during conditioning and copper flotation.

A copper concentrate is floated using 10-20 g/tonne of an alkylalkyl thionocarbamate (Cyanamid AERO 3894) and cleaned three times. The final copper concentrate contains 22% copper, 6% lead and 3000 g/tonne silver.

Significant quantities of marmatite float with the copper/lead concentrate. Marmatite has been found to be very finely disseminated in galena so Brunswick Mining are forced to produce a low-grade zinc concentrate from the lead concentrate (35,72).

The copper flotation circuit tail is reground to 85% minus 38µm in a 3.7m by 4.6m ball mill before being fed to the zinc, lead separation circuit. The ball mill discharge is heated to 70 deg C using steam injected into the pulp, to destroy any residual collector or starch from the copper flotation circuit. The pulp is then cooled to 35 deg C by decantation and repulping with cool water.

Lime is added to pH 10.8 and copper sulphate added to activate the marmatite (1000 g/tonne). A bulk (predominantly zinc) rougher concentrate is then floated with xanthate and cleaned twice, producing a final bulk zinc/lead concentrate containing 34% zinc and 18% lead. The tailings from this circuit assaying 35% lead and 5% zinc is the lead final concentrate.

The zinc circuit is conditioned with 650 g/tonne copper sulphate at pH 10.5 using lime for pH adjustment. Further conditioning with plant xanthate (30 g/tonne) and cyanamid's AERO 3894 (20 g/tonne) leads to the flotation of a zinc rougher concentrate (34% Zn), from 32 Outokumpu 8 m³ cells.

The zinc rougher concentrate is reground in a 3.8m by 4.9m ball mill to 95%, -38µm and floated three times in a closed zinc cleaner circuit. The final concentrate contains 52% zinc.

The zinc rougher tail was formerly directed to a scavenger, or secondary zinc flotation circuit. This used to produce a low-grade zinc concentrate that was up-graded using reverse flotation of pyrite from the zinc. The assay of this secondary zinc concentrate was 48% zinc, and it was combined with the primary zinc circuit concentrate. This circuit is, however, no longer used.

In summary, flotation problems are a combination of difficult chemistry increased by the fine grind and finely disseminated low-grade bands in the ore. Some of the features of the circuit as a result of these factors are:

- * Zinc floats to the bulk circuit, necessitating the production of a zinc concentrate from the lead/copper bulk concentrate.
- * Excessive pyrite floats to the zinc concentrate, resulting in low zinc grade in the rougher concentrate and increased reliance on the zinc cleaner circuit.

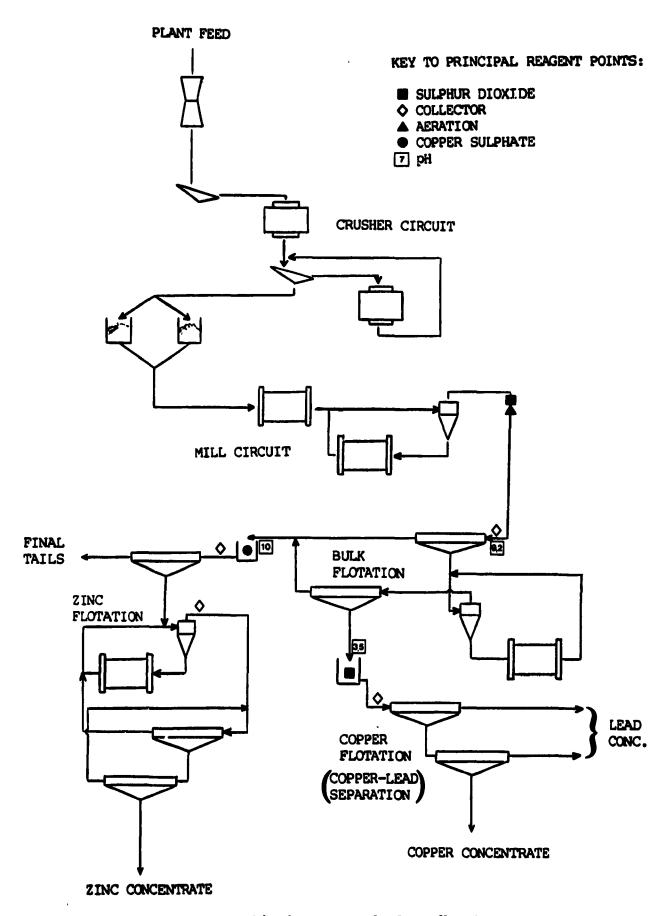


FIGURE 2.5 - Mattabi Mines general plant flowsheet.

2.2.2 Mattabi Mines Ltd, Milling operations (Figure 2.5-4,65).

The mill, set on the shore of Sturgeon Lake 90 km north east of Ignace in Ontario, treats 3000 tonnes per day of pyritic copper, lead, zinc, silver ore mined underground.

The ore treated through the mill when this testwork was performed, was mined from the Mattabi (80%) and Lyon Lake (20%) deposits. Typical head grades are 11% zinc, 0.65% copper, 0.6% lead and 32% iron. The ore also contains approximately 100 g/tonne of silver. The principal valuable minerals are sphalerite, chalcopyrite, galena and tetrahedrite. The most abundant gangue mineral is pyrite, which comprises 70% of the ore. In all, sulphides comprise between 85 and 90% of the ore.

The ore is, by Canadian Shield standards, relatively coarsegrained. The economic optimum grind size being 75% -74µm from primary and secondary milling though intermediate products are ground considerably finer in the bulk copper/lead and zinc regrind circuits. The flowsheet is summarised by Figure 2.5.

The ores from the two mines are batch-crushed and stored separately for later blending. Primary crushing is performed underground by a Bowborough jaw crusher for Lyon Lake ore and through a gyratory crusher on the surface for the Mattabi ore. All ore currently passes through the gyratory crusher, to aid material handling. Both ores are stockpiled separately after primary crushing in an ore storage building with a live capacity of 8000 tonnes.

The mill uses a 33cm \times 2.13m Allis-Chalmers hydrocone crusher for secondary crushing to -1.9cm. A 1.83m \times 4.88m double-deck screen sizes the primary crusher product, the oversize feeding the secondary crusher.

A larger (2.44m x 6.10m) single deck screen sizes the secondary crusher product. This is in closed circuit with an Allis-Chalmers 12.7cm x 2.13m hydrocone tertiary crusher. Product from the crusher plant is stored in two 4000 tonnes fine ore bins. Number 1 bin stores Mattabi ore and number 2 bin stores Lyon Lake ore. Ore fed into the grinding circuit is monitored by a Merrick Model E-315 weightometer.

The current grinding circuit consists of primary and secondary ball milling. The primary mill is a Dominion Engineering 3.05m x 4.88m ball mill, and is charged with 7.62cm forged steel balls. Primary mill discharge comprises a significant amount of coarse rock, predominantly rhyolite. A 1.27cm trommel screens off this coarse material which is barren in base metal sulphides. This is used for road bedding.

The trommel undersize is classified in four 38 cm Krebs primary cyclones. Cyclone underflow feeds a 3.05m x 4.88m Allis-Chalmers secondary ball mill using 2.54cm x 1.27cm "Norcast" slugs. Coarse rhyolite is again screened off using a 1.11cm trommel. Screen undersize is circulated back to the primary cyclone feed. Primary cyclone overflow is directed to the flotation circuit.

The cyclone overflow is aerated and treated with sulphur dioxide to depress the pyrite and the sphalerite. The pH is adjusted to 9-9.5 using soda ash. Collector composed of sodium ethyl xanthate (Cyanamid 325 - 2.5 g/tonne) and aryl dithiophosphoric acid (Cyanamid 241 - 2.5 g/tonne) is added to float the copper, lead and silver.

A bulk concentrate is floated in 16 Agitair 1.4 m³ cells and includes a middlings product, which is circulated back to the conditioner in the bulk float feed. The copper, lead rougher concentrate contains roughly 25% copper and lead combined. This is reground in a Marcy 1.52m x 2.44m ball mill to 95%, -74µm and refloated using eight 1.34 m³ flotation cells to a combined copper, lead grade of 35%. Methyl isobutyl carbinol (MIBC) is used as the frother in the copper/lead circuit.

Starch (wheat dextrine) is added to the copper, lead bulk concentrate together with a small amount of activated carbon to remove residual reagents in solution. Steam is then injected into the pulp, raising the pulp temperature to 50 deg C which strips the adsorbed collector off the mineral surfaces. The pulp is then slowly cooled to 30 deg C and sulphur dioxide added to depress the galena.

A diisoamyl dithiophosphate (Cyanamid 3501) is then added to the pulp and a copper concentrate floated using 8 Denver (1.34 m^3) cells. This is then cleaned using more sulphur dioxide at pH 3.5 and further flotation in 8 Denver 1.34 m^3 cells. The final copper concentrate contains 24% copper and 12% lead. The tails from

copper, lead separation and copper cleaning are combined and directed to the lead concentrate thickener. This final lead concentrate contains 27% lead, 11% zinc and 5% copper.

In contrast to the separation problems encountered in the flotation of copper and lead, the production of a zinc concentrate is relatively straightforward. Copper sulphate (550 g/tonne) is added at pH 10-10.5 (adjusted using hydrated lime), activating the sphalerite. A mix of collectors is added: a thionocarbamate (Minerec 2030) and sodium ethyl xanthate (Cyanamid, aerofloat 325) at a combined rate of 38 g/tonne.

Eighteen Agitair 1.4 m³ cells are used for zinc rougher flotation, producing a concentrate that is directed to a regrind circuit and a middlings that is recirculated to the zinc conditioner. The zinc rougher concentrate contains 43% zinc and is reground in a 2.13m x 3.35m Allis Chalmers ball mill, in closed circuit with 6 Krebs cyclones to 95%,-74µm. This is now cleaned using 16 Agitair 1.4 m³ cells (middlings and tails are redirected to zinc cleaner feed and zinc rougher feed respectively). The cleaner concentrate is re-cleaned using 8 Denver 1.34 m³ cells in closed circuit, producing a final concentrate of 53% zinc.

Overall plant recoveries are 91% of the zinc, 75% of the lead and 50% of the copper. The main metallurgical problem lies with copper/lead separation and zinc depression in the bulk circuit, though the latter problem has only become significant since a greater proportion of the ore has originated from the Lyon Lake

deposit (65).

Zinc flotation is relatively straightforward, rougher grades being roughly 45% zinc. Consequently not much is demanded of the cleaner circuit making it relatively easy to operate.

2.2.3 Kidd Creek Mines "D" Division (Figure 2.6-2,84)

The Kidd Creek Concentrator is located 28 km east of Timmins, Ontario. It treats two types of ore known as A ore (a copper-zinc ore) and C ore (a copper-lead-zinc ore). The total tonnage treated is 12,370 tonnes/day. This thesis is concerned with the C-type ore, which is treated separately from the A-type ore. The throughput of "C" ore is 3100 tonnes/day.

The ore is mined from a massive pyritic deposit, which contains 9% zinc in the form of sphalerite, 0.9% lead as galena, 0.6% copper as chalcopyrite and 230 g/tonne silver, largely as native silver. The chief gangue mineral is pyrite with pyrrhotite occasionally occurring. The chief non-sulphide gangue is rhyolite though graphite may also occur.

Primary crushing at the mine-site reduces the run-of-mine ore to minus 150 mm. This is further crushed in a 1000 tonne/hour fine crushing plant, to minus 16 mm. Primary crusher product is screened on a 50 mm square hole rubber deck screen, the oversize feeding the secondary crusher. T.o Allis-Chalmers hydrocone crushers perform the secondary crushing, these being set to 32 mm. Undersize is screened on a 19 mm steel rod-deck screen. Oversize from the rod-deck screen is combined with oversize from the crusher product screens and recycled to the tertiary crushers, three Allis-Chalmers hydrocone crushers set at 13 mm. Crusher discharge is sized using two 40 mm by 23 mm slotted screens.

FIGURE 2.6: Kidd Creek Mines "D" Division general flowsheet.

The grinding circuit consists of an Allis-Chalmers 3.2m by 4.9m rod mill and 3.7m by 5.5m Allis-Chalmers secondary and tertiary ball mills. The rod mill is lined with Noranda Ni-hard wave liners and Domite chrome-moly end liners and uses forged steel rods. The ball mills are rubber lined and use forged steel balls. The primary mill discharge is classified in 38cm Krebs cyclones, secondary and tertiary mill discharges being classified by 25 cm Krebs cyclones. These cyclones produce a flotation circuit feed at 73 percent minus 45 microns. Sometimes incorporated into the grinding circuit is a silver flotation circuit, which treats a 20 tonnes per hour bleed from the secondary mill discharge. Four Outokumpu 1.5 m³ cells are used for rougher flotation and six Wemco-Fagergren 0.09 m³ cells are used as cleaners. Cyanamids R208 collector is used to collect the silver, MIBC being used as a frother.

The grinding circuit product is treated with 500 g/tonne sulphur dioxide (without aeration). A copper concentrate is then floated at pH 8.5 (adjusted using lime) with 60 g/tonne Cyanamid R317 and/or R208 collectors. MIBC is used as a frother. Rougher concentrate is cleaned three times, at pH 6.5 (dropped using sulphur acid to aid galena depression). Denver and Wemco cells with a total capacity of 55 m³ float the rougher concentrate, cleaning is performed in Denver cells with a capacity of 16 m³. The final copper concentrate is relatively low-grade, being typically 17% copper.

Following copper flotation, 48 g/tonne of xanthate collectors (R208 and R317) are added to float the lead. The cell configuration is the same as the copper circuit, although Denver cells with 23 m³ capacity are used for cleaners and no Denver cells are used for the roughing stage. Flotation is performed at pH 8.2, yielding a final lead concentrate containing 12% lead and 10% copper.

and is conditioned for approximately 8 minutes at pH 10. Xanthate is added during copper conditioning at approximately 60 g/tonne. Zinc roughing and scavenging uses three banks of Wemco 4.2 m³ cells, each bank with 14 cells. Rougher concentrate is cleaned three times at pH 11 using Wemco 4.2 m³ cells. The tails from each cleaner stage are combined with a scavenger concentrate and reground in a 2.4 m by 3.7 m Allis-Chalmers regrind ball mill. This is in closed circuit with 15 cm Krebs cyclones. Cyclone overflow is retreated with copper sulphate, collector and frother and added mid-way down the rougher circuit. The final concentrate assays 50% zinc.

In recent years the zinc concentrate has been subjected to a pyrite reverse float, whereby significant quantities of sulphur dioxide have been added to the concentrate (roughly 2000 g/tonne of concentrate) dropping the pH to 4.0. This is conditioned for 15 minutes and cyanamid 3418 collector added. A silver-rich pyrite concentrate is floated and cleaned three times. The pyrite circuit tails now assay 55% Zn, the pyrite concentrate is added to the copper concentrate increasing the silver assay of the

latter.

The success of the pyrite float is indicative of the poor zincpyrite selectivity achieved in the zinc circuit. Poor copper-lead
selectivity is also a feature of the circuit, this particularly
affecting lead grade and recovery. The low feed grade of the
lead, together with the low silver content and economic value of
the lead concentrate reduces the economic significance of the
lead circuit. Much of the attention of research into the plant is
therefore focused on the zinc circuit.

2.3 THE USE OF GASES OTHER THAN AIR IN FLOTATION

While air is probably used as the flotation gas in more than 99% of the plants world-wide, gases other than air played a significant role in the early development of flotation. Some of the earliest recorded work in flotation development was performed by the Bessel Brothers in the 1870s and 1880s. They used steam and carbon dioxide as the gaseous phases in flotation. The work by Potter in Australia made use of reactions between acid and carbonate minerals to form carbonic acid, and this became the carrier gas in the flotation process (57).

The application and potential application of some gases is described in this section.

2.3.1 Nitrogen

The industrial use of nitrogen in flotation is currently limited to copper/molybdenum separation. While the use of nitrogen (as patented by Delaney in 1972, (18) is not to control mineral collector interaction a brief description of its use in copper/molybdenum flotation metallurgy is still useful.

A copper/molybdenum bulk concentrate is normally floated using conventional xanthate and dithiophosphate collectors (13). However the subsequent separation of molybdenite by flotation from chalcopyrite takes advantage of molybdenite's natural hydrophobicity. So, for effective copper/molybdenum separation, the collector coating must be removed from the chalcopyrite.

Acidification and conditioning alone fails to remove all this coating so a strong reducing agent is added. In the case of Gibraltar Mines (British Columbia) sodium hydrosulphide (NaHS) is used for this purpose (although Nokes reagent is also commonly used). This, like any other strong reducing agent is readily oxidised by dissolved oxygen during flotation and as a result over 8,000 g/tonne of sodium hydrosulphide is used, considerably more than the 25 g/tonne that, according to stochiometric calculations, is needed (13).

The idea of using nitrogen appears to have derived from work by Soviet investigators on using various gases, including steam in copper/molybdenum reparation. The gas lowers consumption of the reducing reagent and so reduces reagent cost. Nitrogen may find a use in flotation whenever reducing conditions are beneficial; and it could theoretically reduce the consumption of other reagents such as sodium cyanide and sodium hydrosulphite.

Savings upon using nitrogen are often considerable. Podobnik and Shirley reported that the savings in Nokes reagent were over 60% (73). At Gibraltar Mines, the drop in sodium hydrosulphide consumption has been more than 75%, saving over a million dollars per year. In the Soviet Union the use of nitrogen has been reported to have cut reagent costs in half (89).

While Shirley (1980) noted that nitrogen gas could only be used beneficially in copper flotation, the increasing use of flotation columns and the resultant reduction in gas volumes used in flotation may open new economic avenues for the use of nitrogen.

Furthermore, Soviet investigators have recently established that nitrogen effectiveness is related to pulp temperature, temperature control being one way for further optimization in the use of nitrogen (89).

Outside copper/molybdenum separation, the potential value of nitrogen as a carrier gas in flotation has never been assessed. Its role as a pyrite depressant in sphalerite/pyrite separation has been proposed by Rao and Finch (76-7) though the results in this thesis will indicate that the role of nitrogen as a pyrite depressant may be hard to control.

The use of nitrogen in flotation is of particular relevance to this thesis so Section 2.4 is devoted to describing how nitrogen is produced for industrial applications.

2.3.2 Carbon dioxide

Carbon dioxide is produced as a by-product from nitrogen production at plants such as Gibraltar. In this case it is used as a substitute for sulphuric acid to reduce the pH in the molybdenum circuit (13).

Miller and co-workers have recently proposed the use of carbon dioxide to float fine coal. Carbon dioxide was used to produce a super-clean coal (so-called carbon dioxide coal flotation) by improving ash rejection and increasing the rate and extent of flotation for high-volatile bituminous coal (46). The success of the procedure is attributed to the coal's high adsorption potential for carbon dioxide.

Following this use in coal flotation another possible area is bitumen flotation. It is difficult, however, to imagine widespread application of carbon dioxide in froth flotation (98).

2.3.3 Steam

Many complex sulphide plants use steam as a conditioning gas prior to flotation. Two examples are Brunswick Mining and Mattabi Mines, as described in the previous section. Steam's normal role is to destroy residual reagents in solution after bulk flotation (68). However Berger and Evdokimov (10) have experimented with the use of steam in combination with air, as the flotation carrier gas. They tested this novel approach on several ores and observed reduced reagent consumption and flotation time, and improved extraction and selectivity. The effect of steam may be to improve froth cleaning by raising water temperature and aiding film drainage (67).

2.3.4 Sulphur dioxide

Sulphur dioxide is now probably the most commonly used gas beside air in flotation. Its use, however, is purely as a conditioner and it never acts as the carrier gas. In Canada, numerous plants use sulphur dioxide prior to complex sulphide flotation. Examples are given in Table 2.2.

sulphur dioxide is mainly used for the depression of pyrite, sphalerite and galena (2,4,11,20,68,85). While Table 2.2 shows that many of the larger complex sulphide circuits in Canada use sulphur dioxide, other mines continue to use the older established depressants such as cyanide and zinc sulphate (e.g. Curragh Resources, Polaris Mine and Westmin Resources)(42,88). Yamamoto (106) gave two reasons for the replacement of cyanide with sulphur dioxide as a depressant - (1) cyanide is an environmental hazard and (2) cyanide tends to dissolve some of the precious metals in the ore. Gaudin (26), in fact, mentions that cyanide depression of sphalerite is only effective when significant quantities of iron are associated with the zinc. (This perhaps explains the use of cyanide at Curragh resources where the sphalerite contains roughly 15% iron in the matrix.)

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Table 2.2: Some Canadian base metal operations using sulphur dioxide during flotation.

name:	minerals recovered	addition rate g/t
Brunsvick Mining:	Cu/Pb/Zn/Ag	900
Geco:	Cu/Pb/Zn	430
Kidd Creek Mines, "A"	div:Cu/Zn	341
а и и 'иDи	div:Cu/Pb/Zn	381
Matagami Lake:	Cu/Pb/2n	294
Mattabi Mines:	Cu/Pb/Zn/Ag	345

Significant research has recently been conducted into the chemistry of sulphur dioxide action in complex sulphide flotation. A major literature review of the theory behind the use of sulphur dioxide, or more precisely the derived ions such as

the sulphite ion, in complex sulphide flotation is beyond the scope of this thesis. Some of the more important findings will, however, be covered.

At Brunswick Mining, Kidd Creek Mines and Mattabi Mines, sulphur dioxide is used to depress galena, sphalerite and pyrite at various stages in the process. All three plants add sulphur dioxide to the pre-aerator for pyrite and sphalerite depression prior to copper flotation. However, the two Noranda mines use sulphur dioxide to selectively float galena to the bulk concentrate and then to selectively depress galena in copper/lead and zinc/lead separation stages. This apparent contradiction of uses illustrates the complexity of sulphite ion / metal sulphide interactions. Peres (71) conducted an extensive study to evaluate the effectiveness of sulphur dioxide on the flotation of copper, lead and zinc. The depressing action of sulphur dioxide on different minerals is compared in Table 2.3.

TABLE 2.3: Depressing action of sulphur dioxide in order of effectiveness.(71)

Sphalerite most effectively depressed Marmatite
Pyrite
Galena
Covellite
Chalcocite
Chalcopyrite least effectively depressed

Several mechanisms have been proposed for the depressing effect of sulphite ions on sphalerite flotation.

1. Sulphite ions may depress sphalerite through the formation of calcium or zinc sulphite films on the sphalerite surface (96,106). So, for effective sphalerite depression zinc or calcium ions must be available in solution. Table 2.4 shows that while some operations appear to coincide with this approach, others contradict it (2,4,64,68):

TABLE 2.4: A list of operations using sulphur dioxide for sphalerite depression, and the regulators used with it.

Operation	Amount SO ₂ added	Other regulators
Mount Isa	500-1000	zinc sulphate sodium cyanide
Ainai, Japan	70-1000	zinc sulphate sodium cyanide
Hanava, Japan	70-1000	zinc sulphate
Brunsvick Mining	150	none
Mattabi Mines	150-500	none
Kidd Creek Mines 'D' Division	350	none

2. Gaudin et al, taking an approach based on solid/solution equilibria (27), proposed that the depression of sphalerite was the result of lowering the cupric ion concentration in solution through reduction of the solution potential. Gaudin calculated that the addition of 600 g/tonne sulphur dioxide would effectively remove copper ions from solution (26).

3. Misra (64) found that the depression of sphalerite from chalcopyrite also stems from the decomposition of xanthate through action of sulphur dioxide. He found that the rate of decomposition is faster than that of xanthate adsorption onto the sphalerite surface, thus keeping the surface clear of collector. It is slower than the adsorption of xanthate onto chalcopyrite, however, and so cannot prevent a multi-layer coating of collector from forming on the chalcopyrite surface. The respective rates are dependent on sulphur dioxide and xanthate concentrations and the pH. Consequently all these factors influence the effectiveness of sulphur dioxide in promoting selective copper flotation from a copper/zinc mineral association (64).

Galena depression by sulphite ions seems to be the result of formation of lead sulphite on the galena surface. A pre-requisite to the formation of such a film is an oxidised galena surface, indicating that sulphur dioxide will only depress galena if it is heavily oxidised (91). Interestingly, Shimocisaka (91) found a similar mechanism for the depressing action of chromate ions on galena. This, however, was not restricted to oxidised galena, suggesting that chromate ions may be a more effective depressant. In fact Brunswick Mining originally adopted the use of chromate depression of sphalerite and pyrite, only to later reject the approach (66).

Sulphur dioxide also depresses pyrite. As sulphur dioxide is often a reducing agent (depending on the other ions in solution), pyrite depression is often considered electrochemical in nature. Dixanthogen is the most important hydrophobic species on the surface of pyrite (8,105). This can only form at pulp potentials higher than +0.22v vs S.H.E. By lowering the pulp potential below 0.22v vs S.H.E. the mechanism for pyrite collection using xanthates is blocked. However, Hill (38) found that sulphur dioxide does not reduce the potential in streams at Brunswick unless the pH is below 2. Further, sulphur dioxide is used to improve the selectivity of chalcopyrite flotation, and the rest potentials for xanthate adsorption on chalcopyrite pyrite differ by only 80 mV. A more likely explanation is given by Yamamoto (106) and is similar to that proposed by Misra for sphalerite depression. While the adsorption of xanthate onto the pyrite surface is not blocked by sulphite ions, they do desorb the xanthate effectively - the xanthate being decomposed to ethyl alcohol and carbon dioxide through the action of sulphite ions and oxygen. The applicable pH range is limited (pH 6-8) which would help to explain why sulphur dioxide can be used for selective pyrite depression (at pH 8) and flotation (at low pH as formerly at Brunswick Mining).

Recently, Hoyack and Raghavan (43) showed that the depression of pyrite at higher pH levels (as at Kidd Creek Mines) may be due to electrochemical reactions leading to the formation of a hydrophilic surface product, $Fe_2(SO_4)_2$. $Fe(OH)_3$.

The potential value of sulphur dioxide in copper/nickel flotation has been assessed from time to time. Extensive pilot plant—work on the use of sulphur dioxide in floating severely altered "pothole" copper/nickel/platinum group metal-bearing ores in South Africa resulted in improved pentlandite and pyrrhotite grades (45) but this has not been adopted commercially. Results from work by Peres (71), on pentlandite and chalcopyrite flotation are concordant with those noted in South Africa.

2.3.5 Other gases in flotation.

Hydrogen and oxygen are used as carrier gases in electroflotation. The basic principal of electroflotation is to produce hydrogen or oxygen through electrolysis, and to use the gas produced as the carrier gas in flotation. Electroflotation has two possible advantages:

- 1) The chemical nature of the gas can aid the flotation process. An example is the use of hydrogen to float cassiterite, where the gas reduces the oxide to metallic tin, aiding particle hydrophobicity. (41,104)
- 2) Finer bubbles can be produced, and the bubble size better controlled. This could aid the flotation of fine particles. The bubble sizes produced are typically 10-60µm in diameter, there being normally a compromise between bubble size and the stability of the froth produced.(9,31,50,104).

Electroflotation has not been adopted on an industrial scale yet, though testwork such as that on tin ores from Wheal Jane in Cornwall, England has suggested that it holds promise (41).

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2.4 MEANS AND ECONOMICS OF PRODUCING MITROGEN (15).

2.4.1 Nitrogen supply options

There are six ways to supply nitrogen to a flotation plant:

- * Combustion plant on-site.
- * Membrane system on-site.
- Pressure Swing Adsorption system on-site.
- * Cryogenic system on-site.
- * Shipping-in liquid nitrogen.
- * Closing flotation cells and recirculating the air, the oxygen will be consumed by the sulphide minerals.

The relative merits of each system will now be considered.

Combustion plant:

Combustion plants have been built on-site at a number of Canadian copper-molybdenum operations (e.g. Gibraltar Mines and Lornex Mining). They operate by using natural gas to combust the oxygen in the air by the following reaction:

$$20_2 + CH_4 \rightarrow 2H_2O + CO_2$$

The quality of the nitrogen produced depends largely on the level of automatic control applied, being of the range of 95% - 99% N_2 . Plant capacities are variable, though in the copper-molybdenum circuits they tend to be 20 - 30 tonnes per day.

Membrane system:

The membrane system effectively filters compressed air, taking advantage of the fast permeation rates of oxygen, carbon dioxide and water vapour to separate them from the more slowly permeating

Membranes employ the principle of selective permeation to separate gases. Oxygen permeates quickly through a semi-permeable membrane. Nitrogen permeates through at a much slower rate. The driving force which forces the oxygen through the membrane, is the difference between the two partial pressures of the feed and the oxygen discharge.

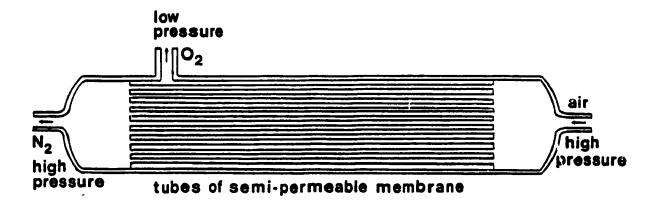


FIGURE 2.7 - Membrane system for producing nitrogen.

nitrogen. Compressed air enters a chamber filled with bundles of semi-permeable membranes formed into tiny, hollow fibres. The faster permeating gases, driven by a higher partial pressure inside the tubes than outside, permeate through the membrane and are collected at a relatively low pressure. Nitrogen remains inside the tubes and discharges the other end of the cylinder (see Figure 2.7). Such a plant produces nitrogen at roughly 95% purity, and has a relatively low capacity of 2.5 tonnes per day.

Pressure Swing Adsorption:

Pressure Swing Adsorption is a commonly used means of providing nitrogen of 99% purity. Compressed air is supplied at the bottom of the first bed where the proprietary sieve adsorbs the oxygen and other impurities (Figure 2.8). When this bed approaches saturation it switches to a regenerative phase while the second bed automatically begins the separation process. A nitrogen reservoir provides surge capacity and ensures a steady output. Plant capacities are generally up to 50 tonnes per day.

Cryogenic plants (101):

Cryogenic plants are the best means for providing large tonnages of nitrogen (more than 50 tonnes per'day). They produce a very clean product (>99.9% N₂) by lowering the temperature of a supply of air to below the boiling point of oxygen (-183 deg C) and separating liquid oxygen from gaseous nitrogen. One stage produces a 50% oxygen containing liquid so two stages are needed to produce a high grade oxygen product. The Linde double column illustrated in Figure 2.9 is an example of a cryogenic system.

A Pressure Swing Adsorption (PSA) system is composed of two identical multi-layered beds which perform the separation process. Compressed air enters the bottom of the first bed where a proprietary sieve adsorbs oxygen and other impurities, allowing the nitrogen to pass through. When this bed approaches saturation it switches to a regenerative phase while the second bed automatically begins the separation process. The cycle is completed when the second bed reaches saturation and starts to regenerate and the first bed, once again, begins production.

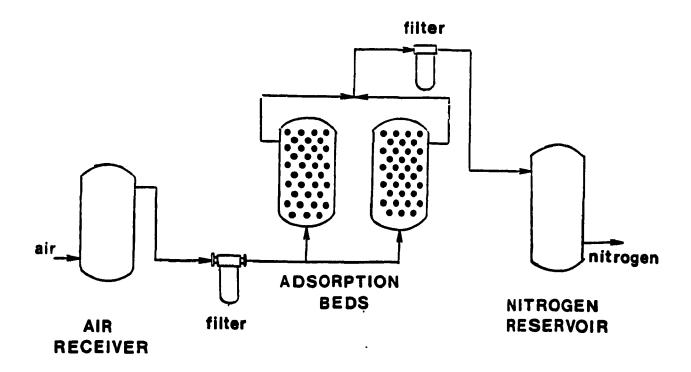


FIGURE 2.8 - Diagram of a Pressure Swing Adsorption system used for the production of nitrogen.

If nitrogen is required at any tonnage above 10-20 tonnes per day, shipping it in on trucks in liquid state is not likely to be viable. If nitrogen is only required occasionally, however, or if a low requirement is expected this may be a viable alternative. This is the only option if the consumer does not want any significant capital outlay.

Probably the most economical application of nitrogen in flotation in Canada is that of Island Copper at Port Hardy, B.C. Plant engineers closed the cells in the copper/molybdenum circuit and recirculated the gas (15). Some oxygen does leak into the system but this is consumed by reagent and the sulphides in the ore. The extra cost of consumed reagent is considerably less than the cost of producing nitrogen which itself would not be entirely clean of oxygen. Brenda Mines have also enclosed the flotation cells in the copper/molybdenum separation circuit (15).

2.4.2 Economic Comparison of the different options.

Table 2.5 compares the economics of the different sources of nitrogen. The capital cost of a nitrogen plant depends on the amount and quality of nitrogen to be produced. A typical plant producing 20 tonnes per day at 95% purity would cost roughly \$750,000. A plant producing 400 tonnes per day of 99% nitrogen would cost roughly \$8 million.

Air goes through tubes cooled in compartments which both cold oxygen and nitrogen flow. It is liquified in the bottom of the lower column where mainly nitrogen rises in gaseous form. It is also greatly cooled in the tubes surrounded by liquid oxygen from the upper column.

The oxygen-rich liquid is transferred to the middle of the upper column where condensation of mainly the oxygen occurs, the more volatile nitrogen rising to the compartment that cools the incoming air. The arrows show the route followed by the liquid and gases.

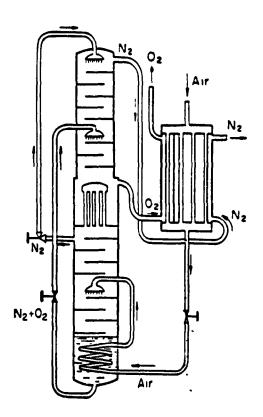


FIGURE 2.9 - Diagram of a double column for separating the oxygen and nitrogen in an air supply (from reference 15).

TABLE 2.5: Comparing the costs of the different means of producing nitrogen.

cost per tonne N,

2.4.3 Other aspects of the use of nitrogen.

A plant that uses large tonnages of nitrogen will probably need to ensure the gas is disposed of for safety reasons. Thus some form of closed cells are going to be needed, which suggests that it would be worthwhile enclosing them completely.

Simply recirculating the air may not ensure consistent quality of nitrogen and will cause problems during start-up so some form of nitrogen producing capacity will probably be necessary. This would also permit a slight positive pressure to be maintained in the flotation cell, reducing the chance of air leaking into the system. While combustion plants have been the most popular they do have the disadvantage of not producing by-product oxygen. The oxygen, produced by membrane, P.S.A., and cryogenic plants can be used either elsewhere in the circuit (e.g. to improve a preaeration stage) or in plant boilers (e.g. for the concentrate driers). This would help off-set the cost of producing the nitrogen. The production of nitrogen will be considered further during the discussion stage of this thesis.

3. EXPERIMENTAL

3.1 ORE PREPARATION

Ore was obtained from the rod mill feed of three plants, Brunswick Mining, Mattabi Mines and Kidd Creek Mines ("D" division). The ore from Brunswick Mining was sampled in 1985 and 1987. The Mattabi plant was sampled in the spring of 1987 and ore from Kidd Creek Mines "D" division was sampled in the fall of 1987.

Plant rod mill feed material is generally too coarse to be ground in the McGill laboratory rod mill without being crushed finer. So each ore was dried (slowly in a cool oven, to avoid oxidation of the ore) and crushed using a laboratory gyratory crusher (Peacock Brothers) powered by a 5.6kW synchronous motor.

The crusher product was split into 1 kg samples using a 12-way rotary splitter, model M12 "prosplitter" (Desert Laboratories, Tucson AZ). The samples were stored dry at room temperature.

The ore was ground wet, at 64% solids by weight, in a stainless steel laboratory rod mill. The mill was 235mm long by 178mm diameter. A combined charge of mild and stainless steel rods was used, consisting of 11, 16mm by 230mm stainless steel rods, 2, 22mm by 194mm mild steel rods and 1, 35mm by 194mm mild steel rod. The total charge weight was 7750 g. Some tests were run using an entirely stainless steel charge but no change in flotation performance was observed.

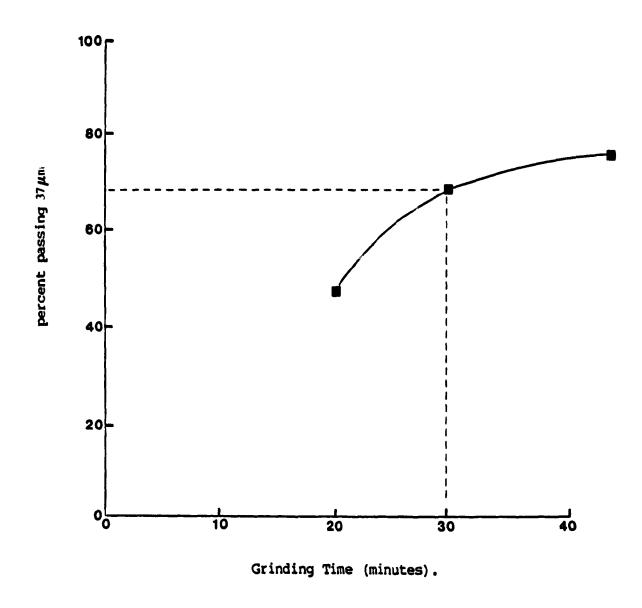


FIGURE 3.1 - Grinding calibration curve for tests on Brunswick Mining ore. The required grind was 65 % passing $37\mu\mathrm{m}$.

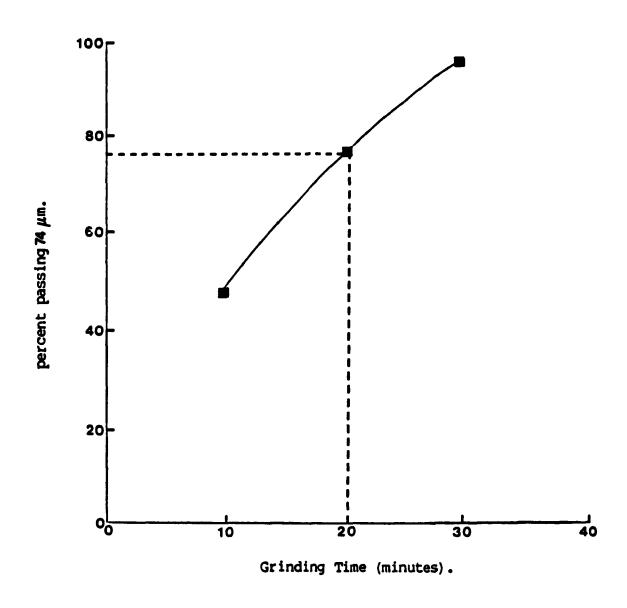


FIGURE 3.2 - Grinding calibration curve for tests on Mattabi ore. The required grind was 75 % passing 74 μm .

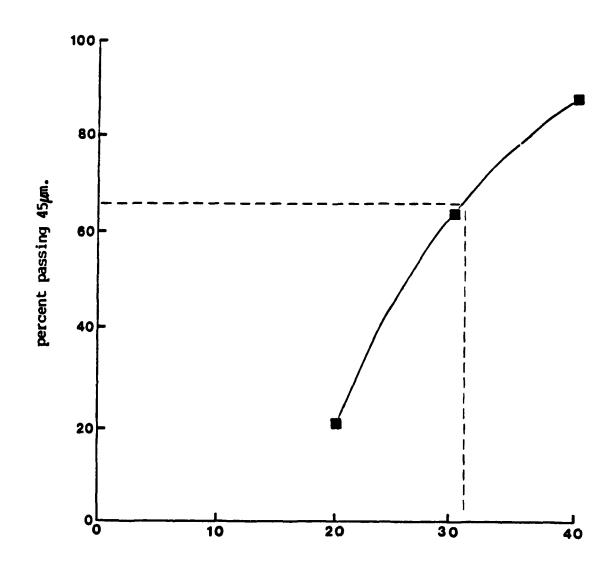


FIGURE 3.3 - Grinding calibration curve for tests on Kidd Creek ore. The required grind was 65 % passing $45\mu\text{m}$.

Grinding Time (minutes).

Grinding time varied from ore to ore, and was determined using grinding tests and preparing a grind time vs size calibration curve. In each case the resultant particle size distribution was as similar as possible to that of the new flotation feed at the respective plant. The grinding test data is presented in Appendix 1, the calibration curves being Figures 3.1 to 3.3. The ground material was transferred from the mill to a 2.5 L Leeds-type flotation cell (Figure 3.4).

3.2 PLOTATION

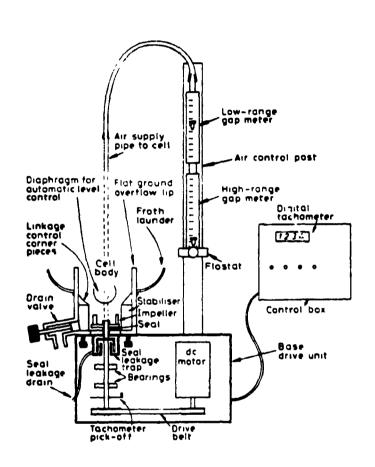
While flotation test reproducibility is always limited by errors in sampling and variations in the operation of a test (6,19), the Leeds cell considerably reduces the amount of error. The cell, designed by C.C.Dell in 1981 has been analysed extensively by researchers in the United Kingdom and South Africa (7,19,67). They concluded that:

- The open top nature of the cell allows excellent access for filling and froth removal, thus maintaining a consistent froth depth.
- * Good control of the impellor speed (through digital readout) and pulp level (controlled automatically or manually) enhances reproducibility.
- * Excellent air dispersion characteristics through use of baffles leading to good flotation performance and stable, deep froths.

The author used a Leeds cell in South Africa, for roughly 150 tests on a magnetic concentrate from Merensky ore (62). Based on that experience, and the experience from the testwork reported in this thesis the author can confirm these observations.

The standard flotation procedures for the three ores are described in Tables 3.1 to 3.4. Most tests, however, were run under specific conditions. These conditions are listed with the respective results sheet in the appendix.

Conditions in the laboratory tests were designed to be similar to standard plant conditions, with one notable exception. Tests at McGill University used zinc sulphate as the principal sphalerite depressant during copper, lead bulk flotation. The main reason for this is the difficulty associated with handling and storing sulphur dioxide at the University, owing to its hazardous nature. Tests at Brunswick Mining compared the use of zinc sulphate with sulphur dioxide, and found that the metallurgical results achieved using the two approaches are very similar. In some tests on Mattabi ore sodium sulphite was used as a substitute for sulphur dioxide. Results from this testwork are descibed in Section 4.2.



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FIGURE 3.4 - Diagrammatic representation of the Leeds flotation cell used in the tests at McGill University.

TABLE 3.1: The standard flotation procedure for tests on Brunswick Mining ore at McGill University:

GRIND:

sample size 1 kilogram

mill steel rod mill (see text)

charge mild/stainless steel rods (see text)

reagents zinc sulphate, 1200 g/t*

pH natural

pulp density 2.1 Kg/litre (64% solids)

note: mill pre-cleaned by grinding with silica for 5-10 minutes.

FLOTATION CELL: Leeds open-top, 2.5 litre

impellor speed 1100 rpm

froth depth 1-2 centimetres

pulp density 1.3 kg/litre (31% solids)

PRE-ABRATION:

Time 10 minutes

Gas flowrate 3-5 litres/minute pH 8.5, modified by lime

BULK PLOTATION: Total flotation

time 4 minutes

collectors Sodium isopropyl xanthate (total 80 g/t)*

frother MIBC

Gas flowrate 3-5 litres/minute

ZINC FLOTATION: Copper Activation

- activator Copper sulphate, 1200 g/t*

- conditioning 5 minutes.

- pH 10.0, adjusted using lime

flotation

- collector sodium isopropyl xanthate (total 80 g/t)*

- time 7 to 11 minutes

- frother MIBC

* prepared each day.

note: Collector prepared as 1% solution.

Copper sulphate prepared as 10% solution. Zinc sulphate prepared as 10% solution.

TABLE 3.2: The standard flotation procedure for tests on Mattabi Mines ore at McGill University:

GRIND:

sample size 1 kilogram

mill steel rod mill (see text)

charge mild/stainless steel rods (see text)

reagents zinc sulphate, 1100 g/t* or sodium sulphite.

pH 9.5, modified by soda ash pulp density 2.1 Kg/litre (64% solids)

note: mill pre-cleaned by grinding with silica for 5-10 minutes.

PLOTATION CELL: Leeds open-top, 2.5 litre

impellor speed 1100 rpm

froth depth 1-2 centimetres

pulp density 1.3 kg/litre (31% solids)

PRE-ABRATION:

Time 10 minutes

Gas flowrate 3-5 litres/minute

pH 9.5, modified by soda ash

BULK FLOTATION: Total flotation

time 4 minutes

collectors Sodium ethyl xanthate, 2.5 g/t and

dithiophosphate, 2.5 g/t*

frother Dowfroth 250

Gas flowrate 3-5 litres/minute

pH 9.5, modified by soda ash

ZINC FLOTATION: Copper Activation

activator Copper sulphate, 1200 g/t*

conditioning 5 minutes.

- pH 10.0, adjusted using lime

flotation

collector sodium ethyl xanthate (total 80 g/t)*

- time 7 minutes

- frother MIBC

prepared each day.

note: Collector prepared as 1% solution.

Copper sulphate prepared as 10% solution.

Zinc sulphate or sodium sulphite prepared as 10%

solution.

TABLE 3.3: The standard flotation procedure for tests on Kidd Creek Mines ore, "D" division at McGill University (5):

GRIND:

sample size 1 kilogram

mill steel rod mill (see text)

charge mild/stainless steel rods (see text)

reagents zinc sulphate, 1000 g/t*
pH 9.0, modified by lime
pulp density 2.1 Kg/litre (64% solids)

note: mill pre-cleaned by grinding with silica for 5-10 minutes.

FLOTATION CELL: Leeds open-top, 2.5 litre

impellor speed 1100 rpm

froth depth 1-2 centimetres

pulp density 1.3 kg/litre (31% solids)

PRE-AERATION:

Time 10 minutes

Gas flowrate 3-5 litres/minute

pH 10.5, modified by lime

BULK FLOTATION: Total flotation

TOTAL TIOCACION

time 4 minutes

collectors Sodium isopropyl xanthate (total 20 g/t)*

frother MIBC

Gas flowrate 3-5 litres/minute

pH 10.5, modified by lime

ZINC FLOTATION: Copper Activation

- activator Copper sulphate, 1000 g/t*

- conditioning 5 minutes.

- pH 11.0, adjusted using lime

flotation

- collector sodium isopropyl xanthate (total 80 g/t)*

- time 7 minutes

- frother MIBC

* prepared each day.

note: Collector prepared as 1% solution.

Copper sulphate prepared as 10% solution.

Zinc sulphate prepared as 10% solution.

TABLE 3.4: The standard flotation procedure for tests on Brunswick Hining ore at the plant laboratory (3).

GRIND:

sample size 2 kilogram

mill steel rod mill (Denver mill)

charge stainless steel rods

reagents sulphur dioxide, 150 g/t* pH 8.0, modified by soda ash pulp density 2.2 Kg/litre (66% solids)

FLOTATION CELL: Venco, 3 litre

impellor speed 1500 rpm

froth depth 1-2 centimetres

pulp density 1.9 kg/litre (55% solids)

PRE-ABRATION:

Time 20 minutes

Gas flowrate valve fully open

pH 9.2, modified by soda ash

BULK FLOTATION: Total flotation

time 7 minutes

collectors Plant xanthate (total 50 g/t)*

frother MIBC

Gas flowrate valve fully open

pH 9.2, modified by soda ash

ZINC FLOTATION: Copper Activation

- activator Copper sulphate, 700 g/t*

- conditioning 10 minutes.

- pH 10.1, adjusted using lime

flotation

- collector plant xanthate (total 50 g/t)*

- time 7 minutes

- frother MIBC

* prepared each day.

note: Collector prepared as 1% solution.
Frother prepared as 1% solution.

Copper sulphate prepared as 10% solution. Zinc sulphate prepared as 10% solution.

A Fisher Scientific "Accumet" pH meter (Model 815 P) with associated electrode, was used for pH measurement. Dissolved oxygen measurements were made using an Orion Research dissolved oxygen probe, model 97-08-00, linked to the above meter.

Pulp potentials were monitored using a system described by Labonte (52). A gold coil electrode was used as a sensing electrode and was measured against a standard calomel electrode. Other factors (such as pH and temperature) affect measurements of pulp potential and were accounted for using the system illustrated by Figure 3.5. Measurements were processed on-line using the program "FLDATAC" on an IBM-computer, giving read-outs at 6 second intervals.

3.3 SAMPLE PREPARATION, ASSAY AND METALLURGICAL BALANCING

Samples from the flotation tests were dried in electric ovens at below 100 deg C, again to avoid oxidation. The dried samples were weighed and, if necessary, split before being sent for assay. Most samples were assayed by Le Centre de Recherches Minerales in Sainte Foy (Quebec), or at the respective mine assay laboratories. A mass balance was generated from the results using the IBM-Lotus 1-2-3 spreadsheet program.

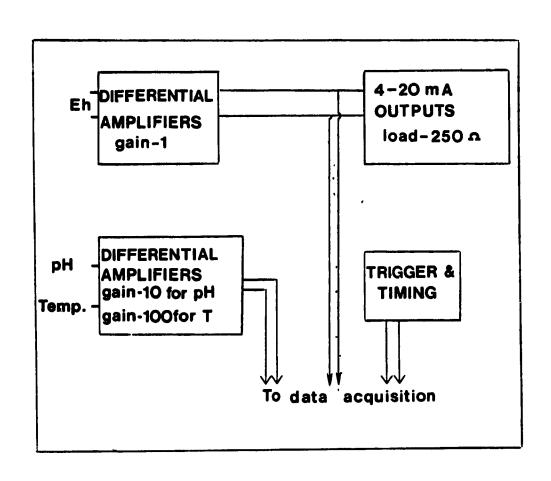


FIGURE 3.5 - Block diagram of the readings used and how they are weighted (amplified) in Labonté's potential measuring system. (from 52)

4. RESULTS

4.1 PLOTATION TESTWORK ON BRUNSWICK MINING ORE

Perhaps the principal problem facing metallurgists at Brunswick Mining is the flotation separation of sphalerite and pyrite. The current flowsheet only allows a 30% zinc rougher concentrate grade to be produced, because of significant mis-direction of pyrite. As a result operators have to rely greatly on the cleaner circuit to make grade, resulting in a relatively complex cleaner circuit and a relatively low grade of zinc (51%) in the final concentrate.

Recently, McGill University has been involved in research to find ways of improving metallurgy in the zinc circuit at Brunswick Mining. Hill (39) concentrated on the mineralogy of the circuit; this thesis considers the chemistry behind pyrite flotation.

This thesis applies some novel concepts, largely derived from the increased understanding of flotation chemistry, to improve selectivity in zinc flotation. As discussed in Section 2.1, of particular interest is the possibility of collectorless flotation of pyrite and the use of nitrogen to improve zinc/pyrite selectivity.

The results described in this section show that collectorless flotation of Brunswick Mining ore is largely ineffective. The tests did, however, identify that pyrite flotation can be

strongly promoted in the presence of a nitrogenated, or more correctly, a de-oxygenated environment. A procedure was developed by which a high-grade pyrite concentrate could be produced prior to zinc flotation.

4.1.1 Standard test and statistical comparison with plant results

In such a program of laboratory tests, it is important to develop a standard procedure which produces similar metallurgical results to the respective production plant. As a consequence, the first part of this section is devoted to determining standard conditions. As the chief area of interest is zinc, pyrite separation all tests generally start with a standard copper, lead bulk flotation stage. (This repeated bulk flotation stage acts as a continual check on test reproducibility.)

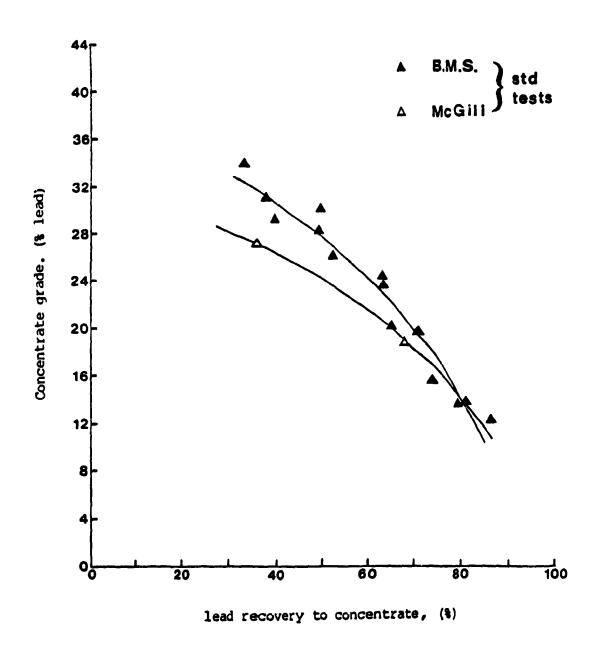


FIGURE 4.1 - Grade vs recovery of lead to the bulk concentrate, from tests at Brunswick Mining employing the Brunswick Mining and McGill University standard procedures.

4.1.1.(a) Results from standard tests employing the McGill procedure.

The tests on Brunswick Mining ore in the McGill University laboratory led to slightly lower combined copper, lead grades in the bulk rougher concentrate than those reported at the plant, at equal recoveries (see Table 4.1).

When the standard Brunswick Mining laboratory test procedure was compared to the McGill procedure, using the same mill and flotation cell the respective results are similar (particularly at typical plant recoveries of 70%, see Table 4.2 and Figure 4.1). This suggests that the improved copper / lead flotation selectivity is a result of a specific plant condition rather than the technique applied. A notable feature from this testwork is the equivalent effect of using sulphur dioxide and zinc sulphate on the flotation depression of pyrite and sphalerite.

At McGill University, the ensuing zinc flotation stage led to a 25% zinc rougher concentrate grade at 72% recovery (or 91.6% recovery from the zinc flotation feed). Again, pyrite rejection in the plant was better than in the laboratory testwork, leading to a 34% concentrate grade at 71% recovery. This may be explained by the addition of the zinc scavenger concentrate to the feed of the plant zinc circuit, raising the zinc assay of the rougher feed.

The laboratory testwork performed on-site at Brunswick Mining included, in each case, some form of pyrite flotation stage prior

TABLE 4.1: Copper, lead, zinc, and iron grades and recoveries in the bulk flotation concentrates from laboratory tests at McGill University and Brunswick Mining #12 plant

test number 10-301

plant data

bulk copper/lead rougher

	grade /	recovery	grade / recovery
	(%)	(%)	(%) (%)
copper	1.14 /	76.1	1.50 / 76.4
lead	10.60 /	75.3	16.70 / 76.7
zinc	7.54 /	21.7	10.50 / 19.3
iron	33.85 /	27.8	28.27 / 17.8

TABLE 4.2: Copper, lead, zinc, and iron grades and recoveries in the bulk flotation concentrates, from laboratory tests using McGill University and Brunswick Mining laboratory procedures. Conditions were otherwise identical. Tests performed at Brunswick Mining.

McGill procedure

BMS procedure

bulk copper/lead rougher

	grade / recovery		grade /	/ recovery
	(%)	(%)	(%)	(%)
copper	1.63 /	57.8	1.58 /	69.4
lead	18.72 /	67.6	19.89 /	72.3
zinc	8.81 /	14.4	9.00 /	14.4
iron	22.99 /	13.4	27.53 /	14.8

TABLE 4.3: Comparison of metal recoveries from bulk copper, lead rougher flotation during laboratory tests and on the plant.

	standard tests	recover other te	plant d ata	
	mean	mean	s.d.	mean
copper	76.1	74.8	3.7	76.4
lead	75.3	73.0	8.5	76.7
zinc	21.7	18.8	5.2	19.3
iron	27.8	17.7	3.2	20.2

to zinc roughing. The zinc flotation feed, therefore, was different in constitution in these cases and the zinc flotation stages cannot be compared.

In most tests a standard bulk copper, lead flotation stage was adopted prior to pyrite and/or zinc flotation. Table 4.3 lists metal recoveries from bulk flotation in the standard test, on the plant, and the mean and standard deviation of all the other tests in the project.

The table shows that deviations from the mean were relatively small. As laboratory flotation tests are always subject to such variable conditions as ore quality, water quality and grind, results have an inherent inconsistency, these relatively low standard deviations are judged acceptable. Table 4.3 also shows that laboratory test results deviate only slightly from the metallurgy achieved on the plant. At the 95% confidence level, the deviations from the plant results are insignificant.

4.1.2 Collectorless flotation of Brunswick Mining ore.

In this set of tests the potential for collectorless flotation of Brunswick Mining ore was assessed. Tests in the section followed the sequence:

collectorless flotation ∇ bulk flotation (tests 10-101,10-102) ∇ zinc flotation

Both air and nitrogen were used as the carrier gases.

Generally, no appreciable collectorless flotation of Brunswick Mining ore in air was observed. However, in one test a longer pre-aeration period resulted a 100% improvement in the collectorless recovery of chalcopyrite. The overall recovery of chalcopyrite, however, was still less than 40% (see Table 4.4).

When a bulk concentrate was floated with collector after the collectorless float, the resulting copper and lead grades were low, reflecting significant flotation of zinc and iron sulphides to the bulk concentrate (see Table 4.5). This in turn led to zinc recovery of only 60% to the zinc concentrate, at a grade of 30% zinc. Zinc flotation without prior bulk flotation led to a low grade zinc concentrate due to the flotation of lead and copper sulphides which had been depressed during the previous collectorless stage. There appears, therefore, to be no advantage in incorporating collectorless flotation using air into the Brunswick Mining flowsheet.

One test was run to check the collectorless flotation of Brunswick material in nitrogen. The results were similar to those in air, showing generally poor flotation of all minerals (see Table 4.6).

Following flotation of the collectorless concentrate, bulk flotation in nitrogen using xanthate was attempted. These results are also outlined in Table 4.6, and indicate strong flotation of pyrite and fairly strong flotation of galena. Other minerals floated poorly.

The remainder of the pyrite floated in the zinc flotation stage.

As a result, only a low grade zinc concentrate could be produced.

TABLE 4.6: Metal grades and recoveries from collectorless flotation in nitrogen of ore from Brunswick Mining, after grinding and nitrogen conditioning.

	Collectorless	flotation concentrate.
	grade (%)	recovery (%)
copper	0.37	11.9
lead	4.81	15.8
zinc	8.68	10.6
iron	26.30	11.5
11011	20.30	11.7
	copper, lead b	ulk flotation concentrate
	grade (%)	recovery (%)
copper	0.48	30.7
lead	4.91	53.5
zinc	2.57	10.3
iron	39.09	56.8
***	33.03	30.0
	zinc flota	tion concentrate
	grade (%)	recovery (%)
copper	0.40	31.7
lead	3.28	26.2
zinc	24.48	74.8
iron	22.51	24.0
411	72172	6 T I V

TABLE 4.4: Hetal grades and recoveries from collectorless flotation in air of ore from Brunswick Mining, after grinding and different levels of pre-aeration.

tests	10-103,10-104	10-102
pre-aera		
time (mi	n) 10	25
	grade/recovery	grade/recovery
	(%) (%)	(%) (%)
copper	0.62/ 16.7	1.07/ 36.8
lead	5.91/ 16.5	5.60/ 20.0
zinc	7.62/ 9.4	7.10/ 9.5
iron	33.51/ 10.5	28.10/ 11.8

TABLE 4.5: The metal grade/recovery relationships from flotation of the copper, lead bulk concentrate in test 10-102.

	copper, lead bulk	flotation concentrate
	grade (%)	recovery (%)
copper	0.70	43.6
lead	9.57	61.1
zinc	9.92	24.0
iron	30.24	24.7

4.1.3 Assessing the effects of aeration and nitrogen on bulk flotation.

The standard pre-treatment method used before bulk flotation involves up to 30 minutes aeration. This is believed to depress pyrite and improve copper and lead grades in the bulk concentrate. (48)

The results described in this section confirm the role of preaeration in pyrite depression and extend this to show that nitrogen can promote pyrite flotation.

The first tests omitted the aeration stage used at Brunswick Mining (and commonly elsewhere). Short periods of xanthate conditioning were used in a nitrogen environment before each bulk flotation step. This resulted in strong flotation of copper, lead and iron-bearing minerals, as indicated by the respective high recoveries in Table 4.7. The amount of zinc mis-directed to the bulk concentrate, possibly by entrainment, limited the grade/recovery characteristics of the subsequent zinc float. Table 4.7 shows that an improvement in zinc grade can be made by removing the pyrite prior to zinc flotation.

The results from these tests show the value of pre-aeration in attaining good pyrite depression. Flushing the oxygen out before xanthate conditioning apparently has no effect on the xanthate collection mechanism. As flotation was in air, it can be argued that the flotation gas provided the necessary oxygen for successful collector-mineral interaction. This is checked in the next section.

TABLE 4.7: Sequential Flotation of bulk copper, lead and zinc concentrates in air following grinding, nitrogen conditioning and xanthate collection.

Bulk copper, lead rougher concentrate

tests 10-209,10-210	grade (%)	recovery	
copper	0.55	82.5	
lead	5.15	79.4	
zinc	5.01	34.4	
iron	38.72	68.3	

zinc rougher concentrate

	grade	recovery
	(%)	(%)
copper	0.19	9.8
lead	1.99	8.7
zinc	32.96	62.0
iron	18.11	9.1

So far, the removal of oxygen during conditioning of the bulk flotation feed has strongly promoted the flotation of pyrite. To further assess the effect of deoxygenated environments on pyrite flotation, the next tests avoided the use of air at any stage after grinding. The results are summarised by Table 4.8.

The overall recoveries of all four elements in two of the three tests were over 80%. As nitrogen was used throughout the tests this suggests that the use of nitrogen does not impede collector-mineral interaction. Whether pyrite is being promoted by nitrogen is unclear from this testwork. It could be argued that air flotation without pre-aeration would produce the same result. This is checked in the next section.

TABLE 4.8: Flotation of (1) a bulk copper, lead concentrate and (2) a zinc concentrate with xanthate, in nitrogen following grinding and nitrogen conditioning.

(1) ref	10-2		k copper 10-2			
141		-	grade/r			
	(%)	(%)	(%)			(%)
copper	1.15/	64.4	0.53/	84.7	0.48/	84.8
lead	3.58/	18.0	4.89/	81.4	5.15/	85.2
zinc	4.74/	12.4	4.22/	28.8	5.06/	38.1
iron	37.41/	23.7	38.84/	75.7	38.66/	80.0
(2)		zin	c conc	entrate		
ref	10-2	08	10-2	05	10-2	06
	grade/r	ecovery	grade/r	ecovery	grade/r	ecovery
	(%)	(%)		(%)	(%)	(%)
copper	0.33/	11.5	0.20/	8.8	0.19/	6.4
lead	8.02/	25.4	1.76/	8.1	1.41/	4.7
zinc	27.44/	45.0	35.24/	66.6	39.71/	55.5
iron	18.37/	31.0	17.21/	9.3	13.44/	5.2

Each individual result is included in Table 4.8 to show that the action of nitrogen in promoting pyrite flotation directly after grinding and nitrogen conditioning, can be unreliable (compare Tests 10-208 and 10-206). While the occasional depression of pyrite can be explained using our current understanding of flotation electrochemistry, through a lower pulp potential, this would not explain the depression of galena.

Zinc flotation is highly sensitive to the metallurgy achieved in the previous stage (bulk flotation). Where substantial flotation of copper, lead and iron sulphides has been achieved as in 10-205 and 10-206 zinc grades and recoveries are higher. The object of the next set of tests was to assess whether the promotion of pyrite could be attributed to the use of nitrogen or the omission of a pre-aeration stage. Strong flotation of copperand lead-bearing minerals was consistently observed. Pyrite also floated strongly, and this is reflected in the poor copper, lead grades.

Eliminating pre-aeration had little effect on zinc recovery to the copper, lead bulk concentrate. Zinc recovery increased from 18% to 21% indicating, perhaps, more entrainment in the bulkier concentrates. The ensuing zinc flotation produced low grade zinc concentrates as illustrated by Table 4.9.

Once again the value of pre-aeration in depression of pyrite has been shown. The results here suggest that pyrite will be activated by any de-oxygenating system (such as conditioners and thickeners), but nitrogen conditioning and nitrogen flotation still lead to the strongest pyrite flotation, as illustrated by Figure 4.2.

FIGURE 4.2: Summary of the effect of different gases and pretreatment steps on the recovery of pyrite in bulk flotation.

<pre>pre-treatment (conditioning)</pre>	flotation gas		pyri	te	recover	γ (%)
•	_		0	20	40	60 80
nitrogen 3 min	nitrogen	80%	11111	111	11111111	
nitrogen 3 min	air	68%	11111	111		1111
none	air	45%	11111	111	11111	
air 10 minutes	air	20%	11111	11		

TABLE 4.9: Flotation of (1) a bulk copper, lead concentrate and (2) a zinc concentrate in air with xanthate, after grinding, no pre-conditioning.

(1)	Cu,Pb concentrate			
		ean		
	grade	recovery		
	(%)	(%)		
copper	0.8	81.4		
lead	9.8	82.6		
zinc	5.9	22.4		
iron	32.4	45.4		
(2)	zinc f	lotation		
	nean			
	grade	recovery		
	(%)	(%)		
copper	0.2	13.8		
lead	1.6	10.4		
zinc	21.3	62.2		
iron	23.6	26.3		

Repeated tests have shown that the use of nitrogen can promote the flotation of pyrite. They have also shown that nitrogen has little effect on the flotation of unactivated sphalerite. So the use of nitrogen can promote pyrite flotation selectively from sphalerite. Such removal of pyrite can improve the ensuing zinc flotation concentrates grade. For this effect to be useful, pyrite must also be extracted separately from the copper and lead-bearing minerals. To this end, a bulk float is now produced in the conventional way, followed by pyrite flotation.

4.1.4 Flotation of a selective pyrite concentrate after standard copper, lead bulk flotation.

In this section, the use of nitrogen for the selective removal of pyrite after standard copper, lead bulk flotation is considered. Results show that air flotation of the pyrite recovers only 23% of the iron from the bulk flotation tails. The use of nitrogen, however, leads to considerably higher recoveries of pyrite.

Standard copper, lead bulk flotation stage results from these tests follow those established in Section 4.1.1 (see Table 4.10). Table 4.10 also shows that further flotation in air yields a pyrite concentrate with both low grade and recovery of pyrite.

TABLE 4.10: Standard flotation of a copper, lead bulk concentrate, followed by the flotation of a pyrite rougher concentrate in air. (Test 10-202)

metal	bulk f	lotation	pyrite	flotation
	gr ade	recovery	grade	recovery
	(%)	(%)	. (%)	(%)
copper	1.45	68.4	0.24	9.3
lead	15.16	74.0	3.05	12.0
zinc	9.13	16.4	8.4	12.2
iron	29.15	18.5	36.0	18.4

Pyrite scavenger flotation was next attempted using nitrogen and air in different tests. The results are compared in Table 4.11 and Figure 4.3, which shows that while flotation in air had reached its recovery limit, the use of nitrogen activated the pyrite, flotation yielding much improved grades and recoveries of iron.

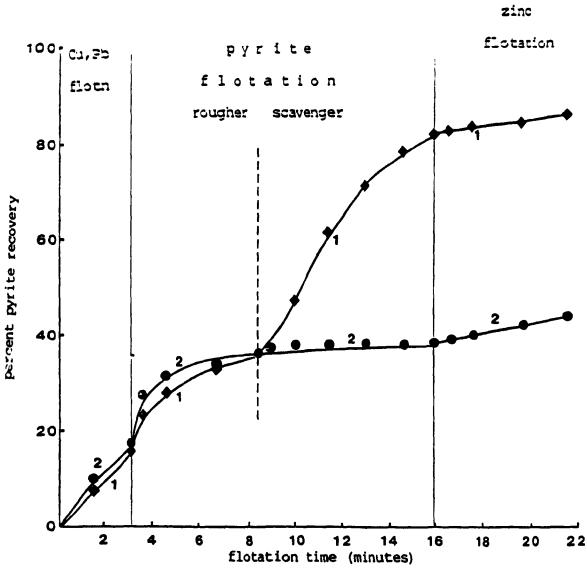


FIGURE 4.3 - Recovery of pyrite as a function of flotation time, showing the effect of nitrogen on pyrite flotation.

Carrier gas used:		
	1	2
bulk flotation	air	air
pyrite rougher	air	air
pyrite scavenger	nitrogen	air
zinc flotation	air	air

TABLE 4.11: The flotation of a pyrite scavenger concentrate following bulk copper, lead flotation and pyrite rougher flotation. Results show the effect of nitrogen on the promotion of pyrite. (tests 10-202,10-204)

	pyrite scavenger flota air nitro			
	air grade recovery		grade	recovery
	(%)	(%)	(%)	(%)
copper	0.18	0.56	0.08	6.7
lead	2.95	0.35	1.07	8.6
zinc	11.80	0.51	4.47	13.7
iron	26.60	0.41	41.04	45.7

So far, nitrogen has only been used to scavenge the pyrite not floated by a prior rougher flotation stage using air. Both rougher and scavenger pyrite concentrates were next floated solely in nitrogen.

Table 4.12 shows that the bulk copper, lead flotation results correspond to the results from standard bulk flotation tests. Again nitrogen promoted pyrite flotation leading to good grades and recoveries of iron. Some collector was needed, between 10 and 20 g/tonne of sodium isopropyl xanthate. Significant quantities of zinc again floated with the pyrite concentrate.

TABLE 4.12: Standard flotation of a copper, lead concentrate in air, then pyrite flotation in nitrogen, mean result from tests 10-204 and 19-213.

metal	<pre>bulk flotation grade recovery (%) (%)</pre>		<pre>pyrite flotation grade recovery (%) (%)</pre>	
copper	1.28	74.8	0.11	13.4
lead	11.90	66.5	1.88	24.3
zinc	7.98	17.6	5.95	29.3
iron	29.24	20.7	39.38	64.3

The effect of removing the pyrite concentrate on zinc flotation selectivity is now considered. Figure 4.4 shows the grade of zinc concentrate floated as a function of zinc recovery, based on the zinc flotation feed. The zinc flotation characteristics of, first, the test including prior pyrite removal using nitrogen and, second, the standard test are highlighted. It indicates that by selectively removing the pyrite, the selectivity of the subsequent zinc flotation stage is improved.

The recovery of zinc is, however, limited by the prior misdirection of zinc to the bulk copper, lead and pyrite
concentrates, as shown by the zinc recovery vs time curve
illustrated by Figure 4.5. The 40% zinc concentrate was floated
at a recovery of 54% based on plant feed, indicating an
improvement of 15% in the zinc grade but at a cost of 17% in zinc
recovery to the pyrite concentrate. The grade of iron in the zinc
concentrate dropped from 20% to 12%. The next section addresses
the problem of zinc loss to the pyrite concentrate.

4.1.5 Reducing the amount of zinc lost to the pyrite concentrate.

Results from the previous tests have shown that high grade pyrite concentrates can be floated from Brunswick Mining ore with minimal amounts of collector added. Copper activation and xanthate flotation of the pyrite flotation tails then yields a zinc concentrate of higher grade than is achieved through the conventional route.

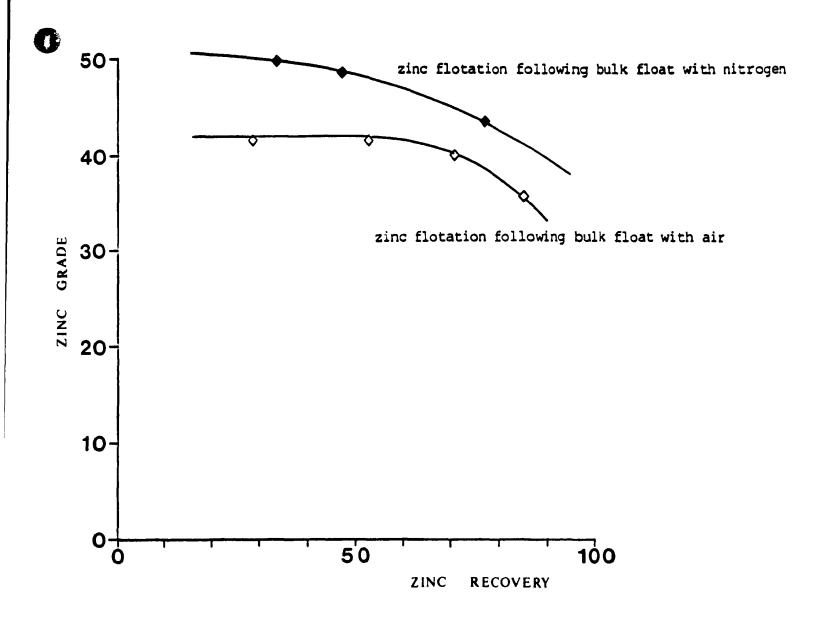


FIGURE 4.4 - Effect of removing the pyrite through nitrogen flotation on ensuing zinc flotation.

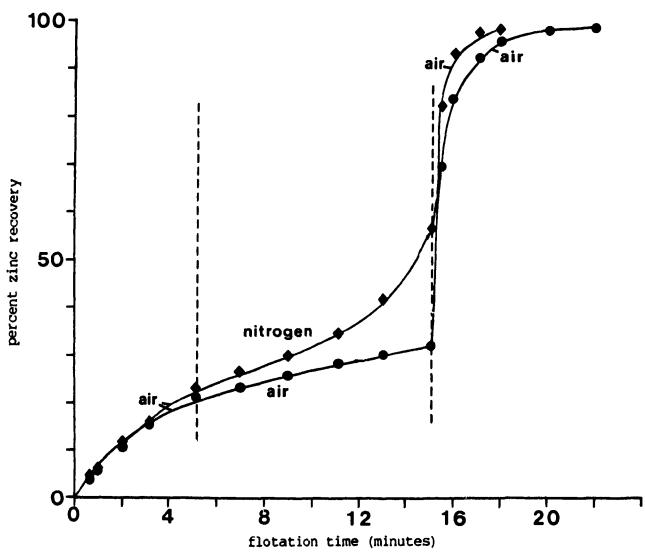


FIGURE 4.5 - Plot of zinc recovery against flotation time showing the effect of nitrogen flotation on zinc recovery to the pyrite concentrate, B.M.S. ore.

The chief metallurgical problem with the process is the loss of zinc that is associated with the pyrite concentrate. Limiting the loss of zinc to the pyrite concentrate and/or recovering it from the pyrite concentrate is fundamental to the economic viability of the proposed treatment route using nitrogen recovery of pyrite.

The work identified two ways to minimise this loss of zinc:

- * Minimising collector addition in nitrogen flotation of pyrite.
- * Floating the zinc from the pyrite concentrate through copper activation and reverse flotation in air.

Table 4.13 correlates the amount of collector added during nitrogen flotation, with the recovery of zinc and iron to the pyrite concentrate. Zinc flotation to the pyrite concentrate can be limited by using as little collector as possible during pyrite flotation. It also shows that at less than 10-20 g/tonne collection addition, the amount of pyrite floated is greatly reduced. Therefore 10-20 g/tonne is the optimum addition rate.

TABLE 4.13: The effect of collector dosage rate on the amount of zinc and iron reporting to the pyrite concentrate during nitrogen flotation.

test	collector dosage	iron	zinc
number	rate	recovery	recovery
	(g/t)	(%)	(%)
10-201	nil	46	13
10-203	10	37	17
10-204	10	51	17
10-501	20	61	23
10-213	20	65	32
10-209	80	80	34
10-210	80	76	39
10-211	100	80	38

Following flotation in nitrogen, the pyrite concentrate is repulped and the pH adjusted to 10.5. Copper ions are added to activate the sphalerite and after 5 minutes conditioning with air xanthate collector is added. A reverse zinc concentrate is then floated in air, the air serving to depress the pyrite. The recovery of zinc to the reverse concentrate is variable, probably being sensitive to conditions not fully understood as yet. In tests at McGill University, the amount of zinc recovered from the pyrite concentrate has been limited (see Table 4.14). Tests onsite at Brunswick Mining achieved significantly better recoveries of zinc as described in the next section.

Pyrite remains depressed during reverse zinc flotation, the recovery of iron being only 3.0%. However starvation quantities of xanthate (10-20 g/tonne) must be used in the pyrite flotation stage to ensure pyrite depression at this stage. While no results are reported here, it was observed that pyrite floated in nitrogen using higher collector addition could not be depressed in the reverse flotation stage.

TABLE 4.14: Metallurgical balance from zinc reverse flotation in air (test at McGill University).

(Sequence - Standard bulk copper, lead flotation; pyrite flotation in nitrogen; pyrite tails to zinc rougher, pyrite concentrate to zinc reverse flotation)

test	10-213			
product	<pre>pyrite final conc. grade/recovery*</pre>		zinc reverse conc. grady/recovery*	
	(%)	(%)	(%)	(%)
copper	0.11	87.8	0.21	12.2
lead	1.48	92.1	1.73	7.9
zinc	4.86	63.5	39.26	36.5
iron	40.50	97.0	17.33	3.0

^{*} note: recoveries based on reverse flotation feed.

4.1.6 Tests performed on-site at the #12 plant, Brunswick Mining & Smelting.

This section was designed to check for any effects on nitrogen flotation of differences in procedure, water type etc. between the standard McGill flotation procedure and the standard Brunswick procedure.

The fundamental difference between the two procedures is the use of zinc sulphate in the grind at McGill University, as against sulphur dioxide. The testwork repeated some of the work already performed, notably comparing the floatability of pyrite in air and in nitrogen, and then checked the effectiveness of the reverse flotation step on the pyrite concentrate.

Bulk flotation results are similar to those using the standard McGill procedure. 81% of the lead floated at a grade of 14.7% and 74% of the copper floated at a grade of 1.0%. Once again pyrite flotation in air was ineffective, recovering 12% of the iron based on the mill feed, at a grade of 39%.

Pyrite flotation in nitrogen was successful. The nitrogen activated the pyrite, leading to an iron recovery of 53%. The froth was, however, visibly more brittle using the Wemco cell at Brunswick Mining than was the case with the tests using the Leeds cell at McGill University. As a result the froths were shallower and permitted more zinc to be carried over with the pyrite. The zinc grade of the pyrite concentrate was 6-8%.

Loss of zinc to the pyrite concentrate is reflected in its low recovery to the zinc concentrate. At first sight the zinc grade (32%) is also low, but closer inspection of the results reveals this is because recovery is high (97% of the zinc in the zinc flotation feed is recovered to the concentrate). Figure 4.6 shows the concentrate grade achieved as a function of zinc recovery, based on zinc flotation feed. This is compared with the same relationship from a standard test at McGill, and shows that the results correspond with those described earlier.

Zinc reporting to the pyrite concentrate was readily recovered through reverse flotation under the conditions prevailing at Brunswick Mining. A pyrite concentrate, floated in nitrogen following standard copper, lead bulk flotation, was cleaned through reverse flotation of the zinc in air. A zinc concentrate containing 32% zinc was floated, recovering more than 70% of the zinc from the pyrite concentrate as outlined in Table 4.16.

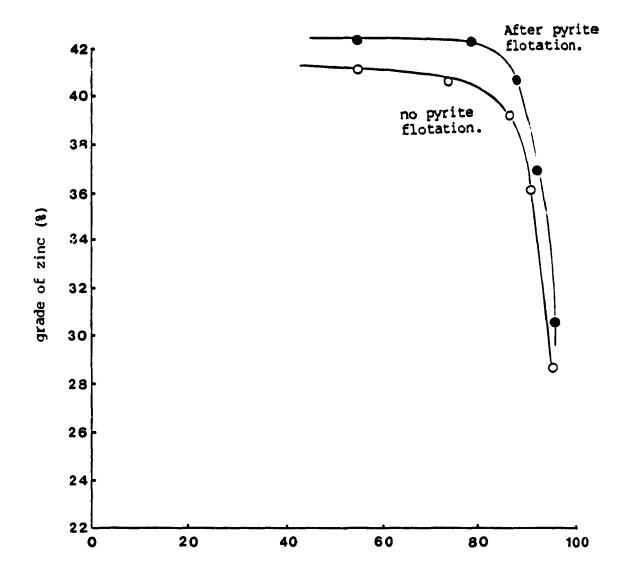
TABLE 4.16: Metallurgical balance from the reverse flotation of zinc from the pyrite concentrate.

(Sequence - Standard bulk copper, lead flotation; pyrite flotation in nitrogen; pyrite tails to zinc rougher, pyrite concentrate to zinc reverse flotation,)

test 10-403

product	<pre>pyrite final conc. grade/recovery</pre>	zinc reverse conc. grade/recovery	
	(%) (%)	(%) (%)	
copper	0.12 84.0	0.14 16.0	
lead	1.20 79.2	1.93 20.5	
zinc	2.14 28.8	32.37 71.2	
iron	42.72 92.2	22.05 7.8	

^{*} note: recoveries based on reverse flotation feed.



recovery of zinc, based on zinc flotation feed (%)

FIGURE 4.6 - Effect of prior removal of pyrite on the grade/recovery relationship from zinc flotation.

(Brunswick Mining ore)

The metallurgical balance for this test is summarised by Figure 4.7, which in turn is compared with the conventional route in the next section.

4.1.7 Summary of the metallurgy achieved through pyrite flotation with nitrogen and zinc reverse flotation using air.

Figure 4.7 summarises the results achieved in these tests from the use of the proposed process on Brunswick Mining ore. It shows that zinc rougher flotation achieves a grade of 38% at 67% recovery. This is considerably better than the grade of 25% at 72% recovery, achieved through the conventional route. An overall comparison of results is illustrated in Table 4.17:

TABLE 4.17: Iron and zinc metallurgy data from the proposed treatment route using flotation of the pyrite concentrate and zinc reverse flotation, compared with the conventional route.

	proposed route		conventional reute	
	Pe	Zn	Fe	Zn
Bulk flotation				
grade(%)	29.7	9.5	33.8	7.6
recovery(%)	24.1	23.2	27.8	21.7
Pyrite flotation				
grade(%)	42.7	2.1	-	-
recovery(%)	40.4	6.1	••	-
Zinc flotation				
grade(%)	15.8	38.4	20.5	25.1
recovery(%)	9.2	67.2	16.7	71.8
Final tails				
grade(%)	19.7	0.9	24.9	0.8
recovery(%)	26.3	3.5	55.5	6.5

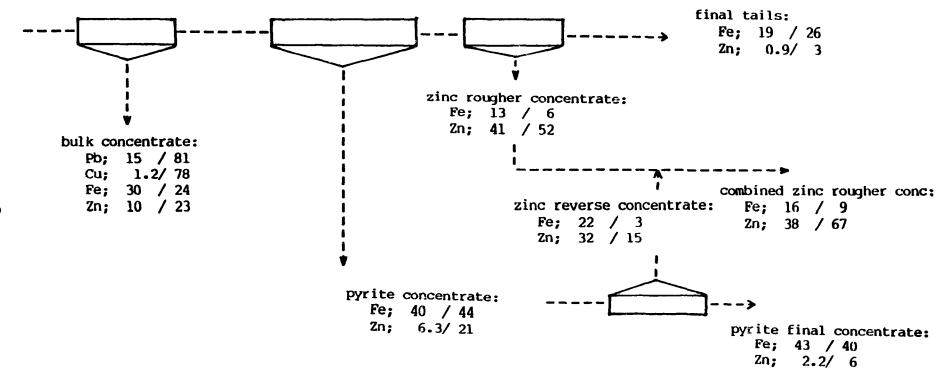


FIGURE 4.7 - Fe & Zn metallurgical balance from test including zinc reverse cleaning.

Labels illustrate: metal; % grade/ % recovery

4.1.8 Preliminary analysis of the chemistry of nitrogen flotation.

Nitrogen is an inert gas and is therefore not likely to play a chemically active role in the promotion of pyrite. To confirm this argon was used in the place of nitrogen. Test 10-501 represents one result from this work. It shows that argon also promoted the flotation of pyrite, 60% of the iron being recovered using 20 g/tonne of xanthate, at a grade of 38% iron.

Some zinc was again floated off the pyrite concentrate, producing a 33% zinc concentrate at 26% zinc recovery. The results are therefore similar to those achieved at McGill University using nitrogen, indicating that pyrite flotation in argon follows the same mechanism as in nitrogen. This confirms that nitrogen plays no direct role in the promotion of pyrite flotation.

The promotion of pyrite flotation is unlikely to be potential related. Pulp potential should fall with the use of nitrogen, which should depress pyrite, according to current electrochemical knowledge of flotation. Figures 4.8 and 4.9 show how pulp potential varies with iron recovery during flotation tests using air and nitrogen during the pyrite flotation stage. Nitrogen slowly drops the pulp potential while at the same time promoting pyrite flotation. The rate at which the pulp potential drops is slow, dropping less than 100mV throughout the 25 minutes of nitrogen conditioning and flotation. Its effect is small in comparison to the effect of copper sulphate or variations in the pH.

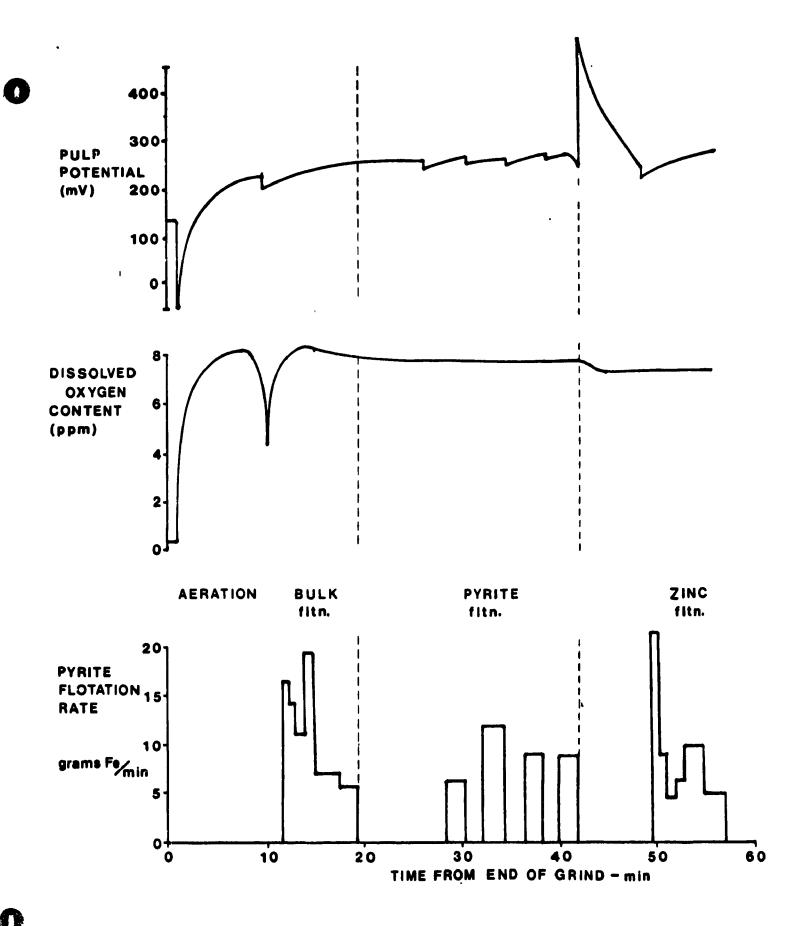


FIGURE 4.8: variation of pulp potential, dissolved oxygen content and pyrite flotability through test including bulk, pyrite and zinc flotation in air.

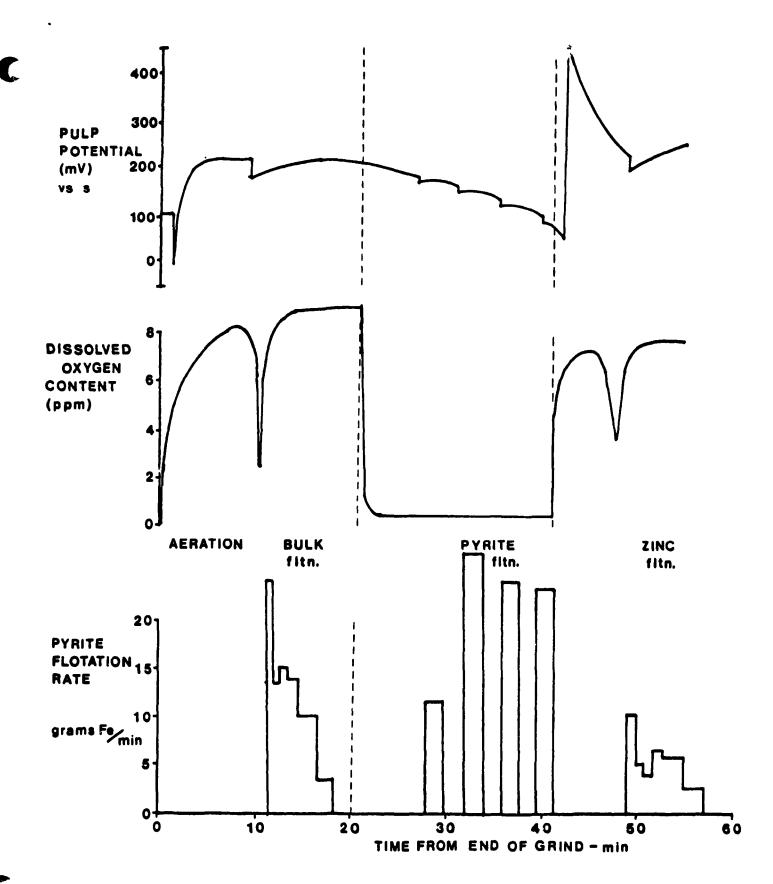


FIGURE 4.9: as figure 4.8 except pyrite flotation is in nitrogen.

4.2 PLOTATION TESTWORK ON MATTABL MINES ORE

Unlike at Brunswick Mining, zinc flotation does not pose any serious problems at Mattabi. However, copper, lead bulk flotation and subsequent separation are difficult. This testwork describes the effect of collectorless flotation on the ore, and assesses the effect of varying the collector dosage rate. Some work examining the effects of sulphite ions and nitrogen on pyrite flotation is included.

Results show that:

- * collectorless flotation recovers more than 80% of the chalcopyrite and 40% of the galena, while rejecting 90% of the pyrite and sphalerite.
- * Sulphite ions aid pyrite rejection from a bulk concentrate floated with collector.
- * Mitrogen promotes pyrite flotation although its effect is limited by the presence of sulphite ions.

Tests performed recently in the laboratory (56) have confirmed some of the observations made above. Some results from these tests, performed on ore from Mattabi's Lyon Lake deposit are included in this report.

4.2.1 Development of standard conditions.

Consistent selective flotation of a copper, lead concentrate proved difficult and the combined copper/lead grade in the concentrate (see Table 4.18) never exceeded 14%. This is lower than the 25% normally achieved on the plant. The lower copper, lead bulk concentrate grades were due to flotation of excess pyrite.

TABLE 4.18: Comparison of the grade/recovery characteristics of copper, lead, zinc and iron from the flotation of the bulk copper, lead concentrate during Test 11-103 and at Mattabi mines.

	Copper/lead Lab.standard		bulk rougher plant data	
	grade/	recovery	grade/	recovery
	(%)	(%)	(%)	(%)
copper	7.80/	70.4	13.00/	75.0
lead	6.79/	55.8	12.00/	75.0
zinc	11.63/	7.2	10.00/	3.4
iron	26.38/	5.4	15.55/	1.8

Zinc flotation results from the laboratory tests were more consistent with plant performance (see Table 4.19). They also show that sphalerite can be floated quite effectively from the pyrite.

TABLE 4.19: Comparison of metal grades and recoveries, zinc flotation in test 11-103 and on the plant at Mattabi Mines.

	-	zinc rougher	r concentrat	te
	test 11-103		plant data	
	grade/	recovery	grade/	recovery
	(%)	(%)	(%)	(%)
copper	0.55/	17.0	0.40/	14.5
lead	1.10/	30.9	0.40/	15.8
zinc	42.43/	90.4	43.00/	92.7
iron	14.98/	10.6	15.70/	11.3

The standard tests reflect the metallurgical problems experienced processing Mattabi ore on production scale. Bulk copper, lead flotation generally yielded poor combined copper, lead grades.

Zinc flotation however produced good grades and recoveries.

The testwork, therefore, concentrated on the problems associated with copper and lead flotation, while also assessing the ability of nitrogen to promote pyrite flotation.

4.2.2 The effect of collector on the characteristics of copper, lead bulk flotation of Mattabi ore.

In test 11-301 a collectorless concentrate was floated after grinding. High recovery of chalcopyrite was achieved (85%) while only 45% of the galena was floated as illustrated by Figure 4.10. Little flotation of sphalerite and pyrite was observed.

Following-up these results, M. Leroux at McGill University (56) has recently tested collectorless flotation on ore from the Lyon Lake deposit, now being mined by Mattabi Mines Ltd. Once again collectorless flotation recovers more than 80% of the copper

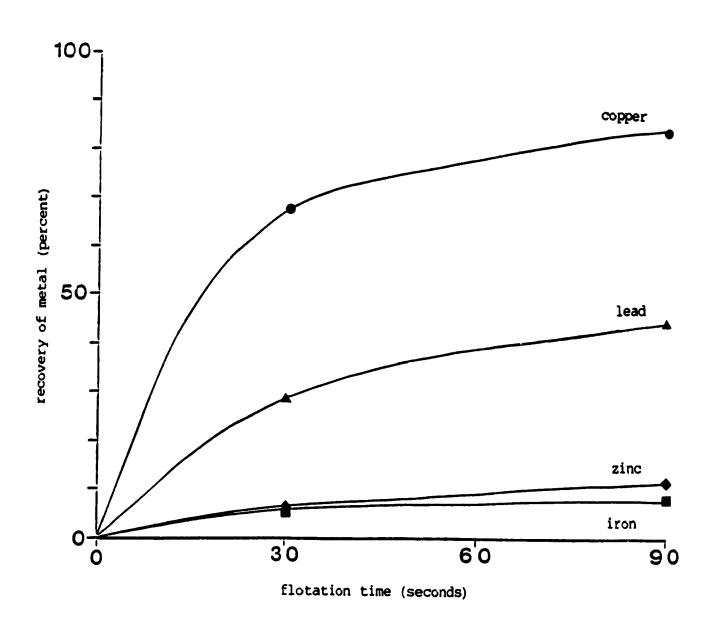


FIGURE 4.10 - Collectorless bulk flotation. Recoveries of Cu, Pb, Zn and Fe against flotation time.

minerals, while recovering 60% of the galena. Pyrite and sphalerite did not respond to collectorless flotation, and the resulting copper, lead combined grade was superior to that using collector. The respective copper rougher grade-recovery relationships for flotation with and without collector are illustrated by Figure 4.11.

The same series of tests revealed that collectorless flotation using nitrogen also recovered almost 90% of the copper minerals. The use of nitrogen appeared to reduce the flotation recovery of galena (see Figure 4.12).

At 5 g/tonne xanthate addition, the recovery of galena was raised to 76%, that of copper, iron and zinc sulphides remaining virtually unchanged (Figure 4.13). At 40 g/tonne xanthate collection the recoveries of iron and zinc sulphides begin to rise (Figure 4.14).

4.2.3 Nitrogen flotation of Mattabi ore

In Test 11-204, 20 minutes nitrogen conditioning preceded bulk flotation in nitrogen. Zinc sulphate was added to the mill at pH 9.5 as the pyrite/sphalerite depressant. A bulk concentrate containing 44% iron at 84% Fe recovery was floated. The bulk flotation iron grade/recovery relationship is illustrated by Figure 4.15.

Under identical conditions, pyrite flotation in air (Test 11-203) is much less effective. Table 4.20 compares the two results:

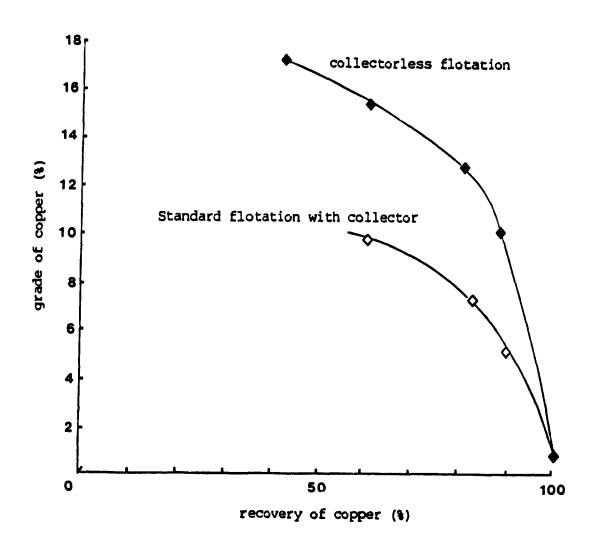


FIGURE 4.11 - Comparison of copper rougher grades and recoveries achieved through collectorless and standard flotation tests at McGill University. Lyon Lake ore.

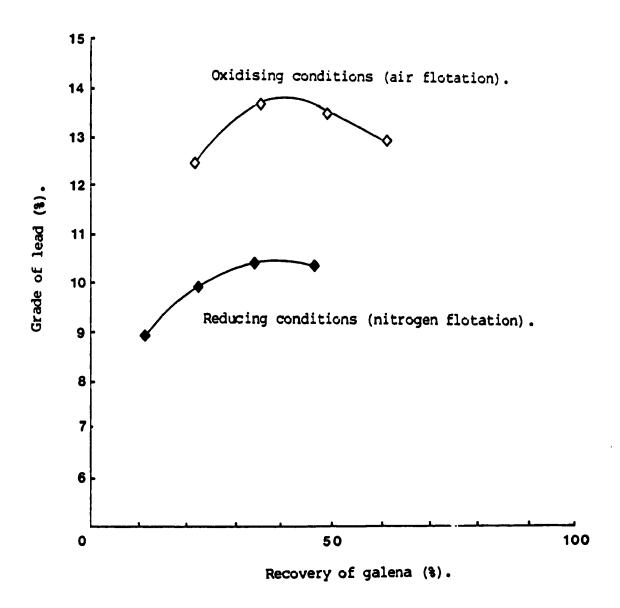


FIGURE 4.12 - Comparison of the effect of oxidising and reducing conditions on the collectorless flotation of galena from Lyon Lake ore.

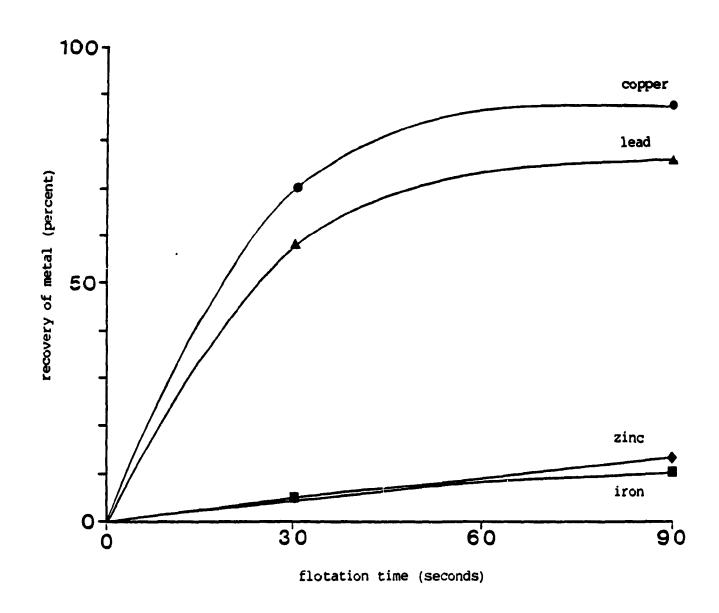


FIGURE 4.13 - Bulk flotation using 5 g/t collector (combination cyanamid 325/241). Recoveries of Cu, Pb, Zn, Fe against flotation time.

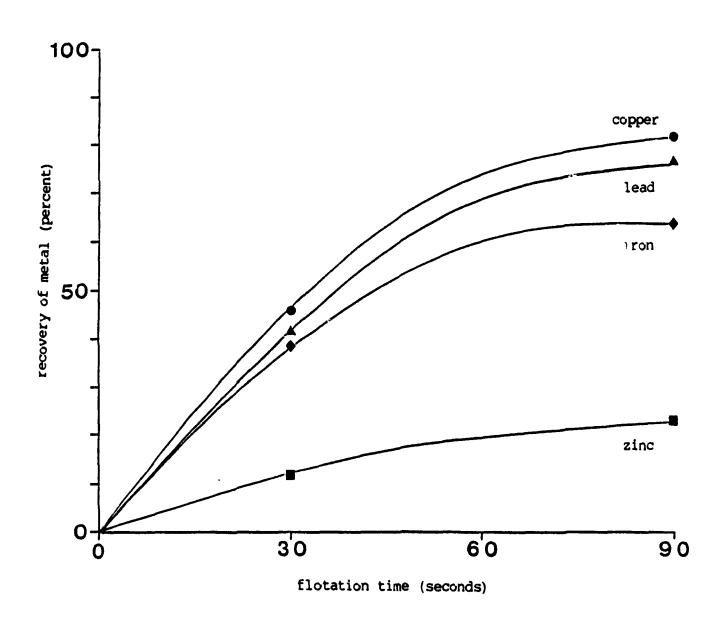


FIGURE 414 - Bulk flotation using 40 g/t collector (combination cyanamid 325/241). Recoveries of Cu, Pb, Zn, Fe against flotation time.

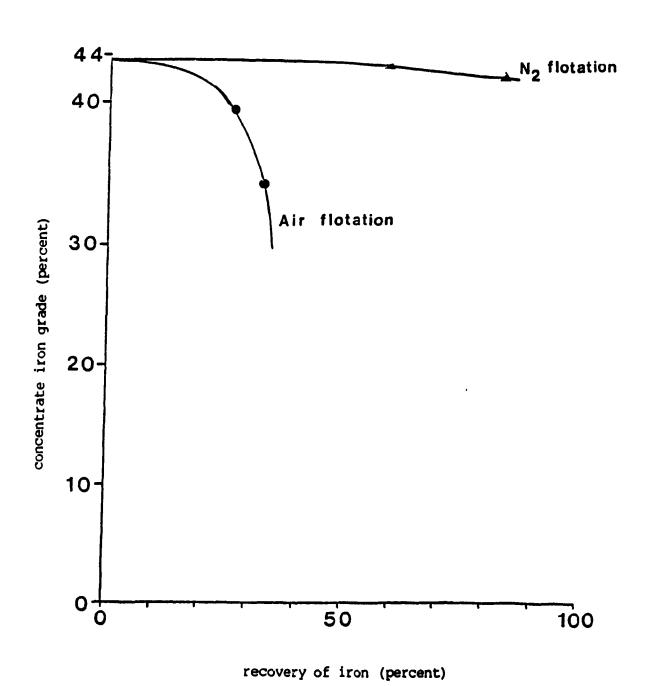


FIGURE 4.15 - Effect of nitrogen on the flotation of pyrite. Grade/recovery relationship of pyrite in Fe/Cu/Pb bulk flotation.

TABLE 4.20: Comparison of results from standard copper, lead flotation using nitrogen and air as the gas phase in flotation.

	Copper/lead using nitrogen		bulk rougher using air	
	grade/	recovery	grade/	recovery
	(%)	(%)	(%)	(%)
copper	0.63/	52.9	2.29/	84.0
lead	0.75/	61.2	1.84/	62.8
zinc	2.15/	11.9	4.77/	11.9
iron	44.24/	83.8	40.22/	33.7

Copper recovery to the bulk concentrate was reduced during nitrogen flotation as illustrated in Table 4.20. Lead and zinc recoveries were unaffected and pyrite recovery more than doubled. This result confirms the observations made during Brunswick Mining testwork that nitrogen can promote the flotation of pyrite.

The use of zinc sulphate as a zinc depressant, maintained as standard for tests at McGill University, resulted in very poor copper, lead grades in the bulk concentrate. Combined copper, lead grades were 4-6% despite recoveries of 55-85% as a result of relatively strong flotation of pyrite. A typical example of this is illustrated by Table 4.20.

Sodium sulphite was much more effective at reducing pyrite flotation and led to copper, lead combined grades closer to (but still lower than) those obtained on the plant as illustrated by Test 11-103 (Table 4.18). Combined copper/lead grades rose to 10-14% at similar (55-85%) recoveries. Figures 4.16 and 4.17 illustrate more clearly the effect of sodium sulphite on pyrite depression. In both cases excess xanthate was added to

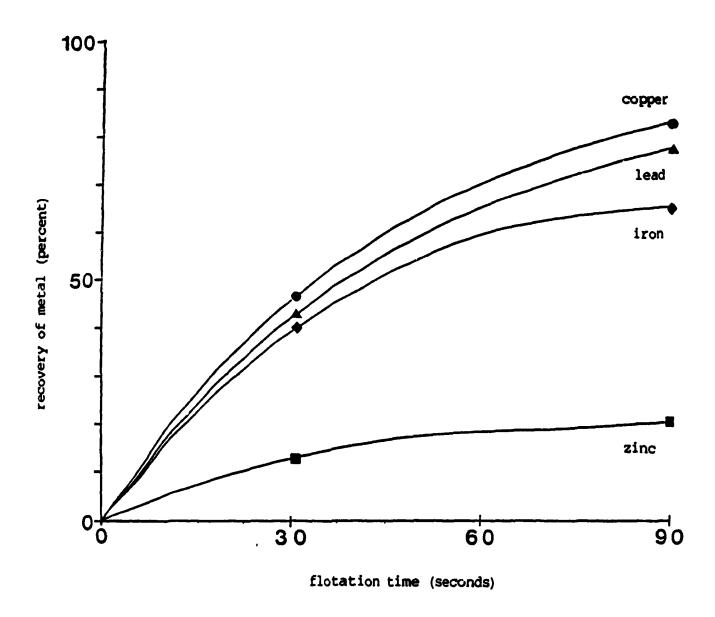


FIGURE 4.16 - Air flotation of Mattabi ore using 1000 g/t sodium sulphite in grind (bulk flotation only). 1000 g/t sodium sulphite is equivalent to 500 g/t SO2.

% Cu, Pb, Zn, Fe recoveries vs flotation time (sec) using excess collector (40 g/t)

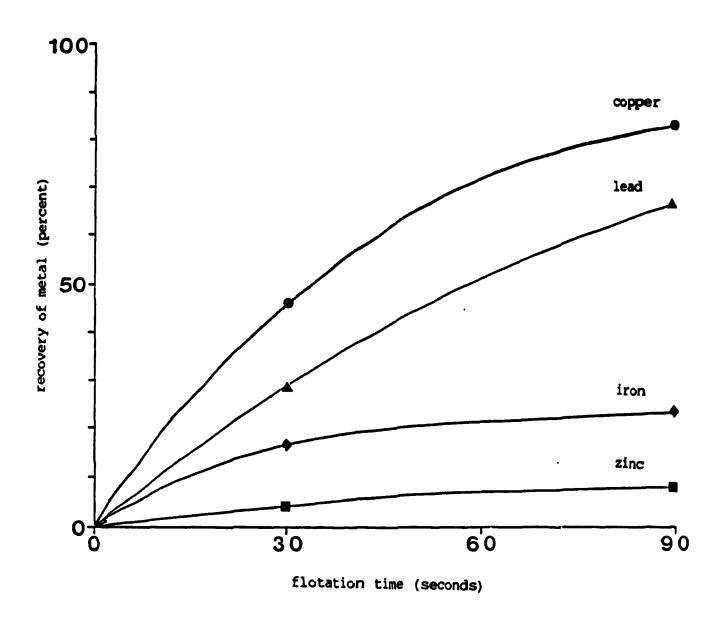


FIGURE 4.17 - Air flotation of Mattabi ore using 3000 g/t sodium sulphite in grind (bulk flotation only). 3000 g/t sodium sulphite is equal to 1500 g/t SO2.

* Cu, Pb, Zn, Fe recoveries vs flotation time (sec) using excess collector (40 g/t) deliberately float the pyrite and this led to a larger differential between recoveries. It is clear that sodium sulphite can strongly, and selectively depress pyrite.

Tests using sulphite ions in air flotation of Lyon Lake ore also showed that pyrite could be selectively depressed, this time at lover dosage rates.

The effect of sodium sulphite on nitrogen flotation of pyrite is complex. With sodium sulphite added to the mill, if bulk flotation is conducted in nitrogen, the recovery of pyrite is relatively low. However, if flotation is performed in air after nitrogen conditioning then significantly more pyrite is recovered. Table 4.21 summarises these results:

TABLE 4.21: Effect of nitrogen conditioning and flotation on the metal grade/recovery relationships of bulk copper, lead, iron flotation - after 1000 g/tonne sodium sulphite addition to the mill.

condition	ing		
gas flotation	nitrogen	nitrogen	air
gas	nitrogen	air	air
	grade/recovery	grade/recovery	grade/recovery
	(%)	(%) (%)	(%) (%)
copper	2.23/ 53.8	1.85/ 77.9	5.26/ 88.1
lead	2.23/ 58.5	1.55/ 60.1	4.80/ 76.0
zinc	6.05/ 9.3	3.89/ 11.1	11.60/ 13.4
iron	37.99/ 19.4	41.06/ 39.6	28.77/ 10.9

Maintaining a nitrogenated environment throughout seems, in this case, to depress both pyrite and chalcopyrite flotation. This is contrary to earlier results using zinc sulphate. However it does follow results from similar tests on Brunswick Mining ore, where nitrogen flotation without any air contact was found to be

erratic. It therefore seems that under certain conditions nitrogen fails to promote pyrite flotation. The effect of sulphite addition and nitrogen on pyrite flotation is illustrated by Figure 4.18. This plots lines of equal pyrite recovery at varying levels of nitrogen conditioning and sulphite addition and shows that nitrogen is only effective when the sulphite addition rate is low.

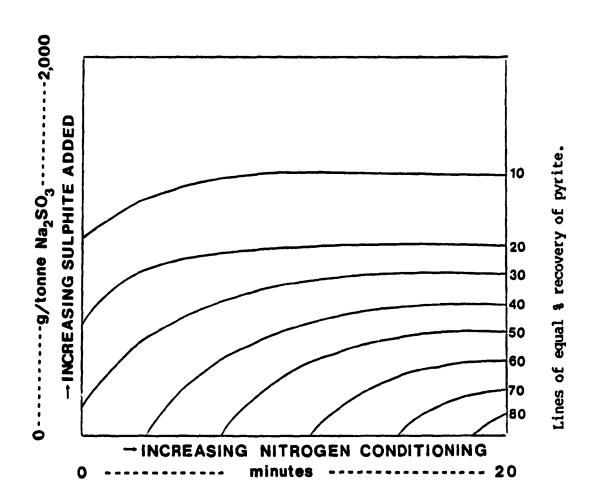


FIGURE 4.18 - Effects of nitrogen conditioning time and sulphite addition on the floatability of pyrite from Mattabi ore.

4.3 TESTWORK ON KIDD CREEK ORE

"D" division at Kidd Creek Mines treats a particularly difficult ore. Both copper and lead concentrates are low grade and pyrite reverse flotation is included in the zinc circuit to remove a highly floatable pyrite constituent.

With pyrite flotation already incorporated into the circuit, the potential use of nitrogen to promote pyrite flotation is of particular interest. Impure nitrogen is available locally so the effect of impure nitrogen was tested on the Kidd Creek ore.

All tests include a standard bulk copper, lead flotation stage followed by flotation of pyrite under different conditions, and finally the flotation of sphalerite in air. As with the Brunswick Mining tests, the presence of the bulk flotation stage prior to pyrite flotation helped to check the reproducibility of conditions such as ore, water and reagent quality, grind and flotation cell performance.

The test results included in this section show that gases with up to 5% oxygen still promote pyrite flotation.

4.3.1. Check on the reproducibility of conditions and comparison with plant results.

Figure 4.19 illustrates the copper and lead grade/recovery bands, during bulk flotation in the four tests reported. They show that the results from the four tests agree reasonably well, implying that flotation conditions remained the same throughout the testwork.

A significant loss of sphalerite to the bulk flotation concentrate was observed. The recovery of sphalerite to the bulk concentrate varied from 18% to 36% leading to zinc grades of 13.5% to 15% in the bulk concentrate. Pyrite flotation to the bulk concentrate was consistently 11-12% except for test 12-101 when it reached 19%. The iron grade was 24-25%.

In each case, the zinc flotation feed varied in composition so it is impossible to directly compare zinc flotation performance. In general, recoveries are low due to the mis-direction of zinc to bulk and pyrite flotation concentrates.

4.3.2. Effect of nitrogen content on the selective flotation of a pyrite concentrate.

Table 4.22 illustrates the effect of varying the nitrogen, or oxygen content of the flotation gas on pyrite recovery and the dissolved oxygen content in the pulp liquor.

FIGURE 4.19 - Grade/recovery relationships for copper and lead flotation to the bulk concentrate. Graphs show copper floated before lead. (Kidd Creek Ore)

TABLE 4.22: Testwork on Kidd Creek Mines, "D" division ore, showing effect of nitrogen content on pyrite flotation and dissolved oxygen content in the pulp.

test number	nitrogen content in gas (%)	pyrite recovery (%)	dissolved oxygen in pulp (ppm)
12-101	80	7.50	7.8
12-104	90	6.80	5.7
12-103	95	22.14	2.5
12-102	100	35.97	<0.5

The use of gases with 5% oxygen or less promotes the flotation of pyrite. This further confirms the observations made from testwork on the two other ores. Table 5 also shows the effect of varying the oxygen input on the dissolved oxygen content of the pulp. In each test, the dissolved oxygen level adjusted to reach an equilibrium dependent on the oxygen content of the gas bubbled into the cell, which is further illustrated in Figure 4.20.

Table 4.23 shows the effect of varying the nitrogen content on the grade and recovery of sphalerite to the pyrite concentrate. It shows that the zinc recovery to the pyrite concentrate is largely unaffected.

TABLE 4.23: The effect of nitrogen content in the flotation gas, on the grade and recovery of sphalerite to the pyrite concentrate.

test number	nitrogen content in gas (%)	sphalerite grade (%)	sphalerite recovery (%)
12-101	80	12.53	10.38
12-104	90	16.96	14.01
12-103	95	10.70	21.52
12-102	100	4.73	15.24

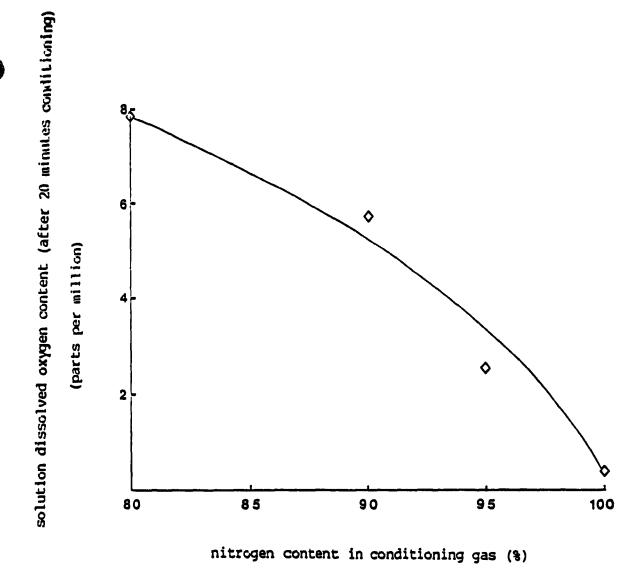


FIGURE 4.20 - Effect of nitrogen enrichment of the conditioning or flotation gas on the dissolved oxygen content of the pulp.

The degree of scatter in zinc recoveries to the bulk and pyrite concentrates makes it difficult to compare zinc flotation grade/recovery relationships based on plant feed. As in previous cases, zinc selectivity is best shown by comparing grades achieved with recoveries based on zinc flotation feed. Figure 4.21 plots the zinc concentrate grades achieved as a function of recovery based on minc flotation feed. Two tests are highlighted, incorporating prior pyrite removal ... rough flotation and one incorporating prior pyrite removal through air flotation. It shows that removal of pyrite through the use of nitrogen results in an increase in zinc flotation selectivity. The same also applies to impure nitrogen with up to 5% oxygen which is probably as effective as pure nitrogen at floating sufficient pyrite to improve zinc flotation selectivity. The use of 95% nitrogen to float pyrite before zinc flotation, in Test 12-103, led to a zinc concentrate grade of 47% at 83% recovery (based on zinc flotation feed).

4.3.3: The effect of nitrogen conditioning on the dissolved oxygen content of the pulp.

Figure 4.22 shows that nitrogen rapidly purges dissolved oxygen from the pulp. The dissolved oxygen content is reduced to 0.5 ppm in less than one minute.

Not all the oxygen is removed by the nitrogen. Figure 4.22 compares the rate at which dissolved oxygen is purged from (1) water and (2) a pulp containing Kidd Creek ore. Oxygen is removed faster from the pulp than from the water. Conversely, when the

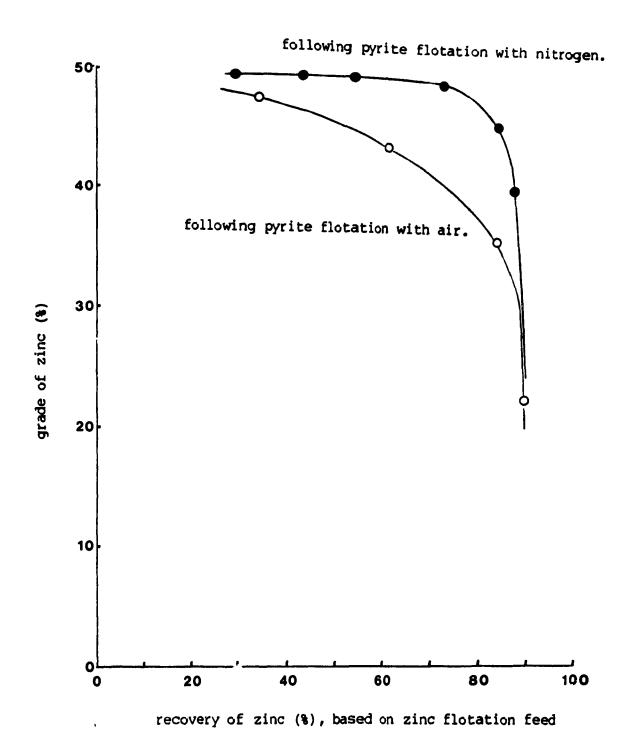


FIGURE 4.21 - Effect of prior removal of pyrite using nitrogen and air, on the selectivity of zinc rougher flotation of Kidd Creek ore. Graphs show the concentrate grades achieved at respective recoveries based on zinc flotation feed.

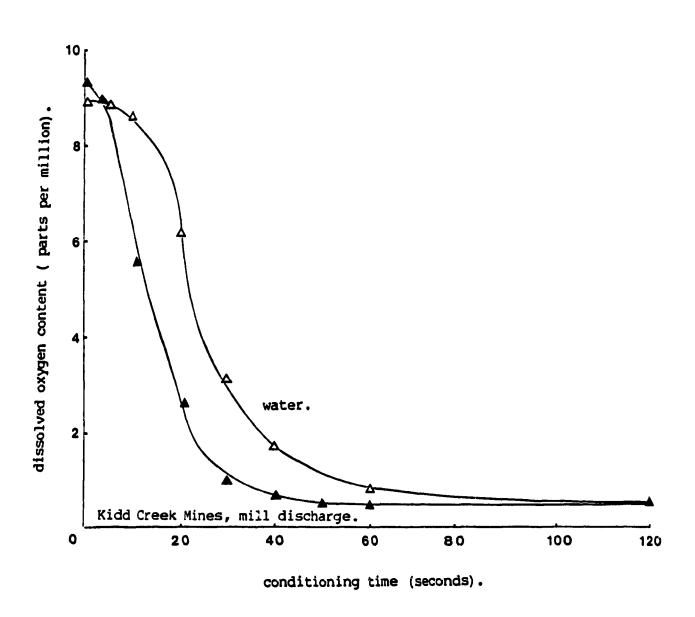


FIGURE 4.22 - Effect of nitrogen conditioning on dissolved oxygen content of water and pulp containing Kidd Creek Mines ore (laboratory mill discharge).

oxygen is re-dissolved through aeration, the dissolved oxygen level rises faster in the water than in the pulp (Figure 4.23).

Both figures suggest that pyrite is constantly drawing oxygen out of solution, accelerating the drop in oxygen content from the pulp during nitrogen conditioning, and retarding the dissolution of oxygen into the pulp water during aeration. Calculations outlined by Appendix 5 suggest that the oxygen demand of the freshly ground ore is roughly 15 ppm dissolved oxygen/minute. This is consistent with other work on similar ores (94).

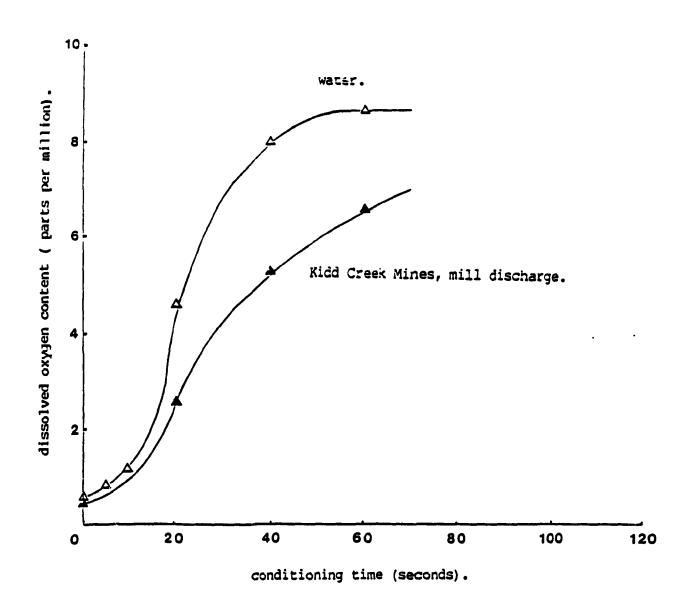


FIGURE 4.23 - Effect of aeration on dissolved oxygen content of water and pulp containing Kidd Creek Mines ore (lacoratory mill discharge).

5.DISCUSSION

5.1 COLLECTORLESS FLOTATION

The results described on three Canadian shield ores of apparently similar mineralogy show a wide variation in collectorless flotation response:

- * Copper minerals from the Lyon Lake deposit (probably largely chalcopyrite) are highly responsive to collectorless flotation. Both copper concentrate grades and recoveries are superior to those when collector is used. Good recoveries and grades were found under both oxidising and reducing conditions.
- * Chalcopyrite from the Mattabi deposit also floated without collector. The recovery was over 80%.
- * Little collectorless flotation of chalcopyrite from the Brunswick Mining ore was observed, although sufficient aeration did improve copper recovery from 15% to 40%. This suggests that chalcopyrite may respond to collectorless flotation under sufficiently oxidising conditions.
- * Other minerals generally showed poor response to collectorless flotation. Galena can float without collector, but its flotation is slow particularly under reducing conditions (reflecting perhaps that the surface oxidation necessary only occurs slowly.).

This work confirms Finkelstein's observations (22) that the collectorless floatabilities of mineral sulphides vary from ore to ore. Several mechanisms have been proposed to explain collectorless floatation (25,36-7,55,58-9,99,102,107-8). These were described in Section 2.1.1 and are summarised by Figure 5.1.

There are enough mechanisms available to explain the collectorless flotation of minerals, consequently it was considered that a better object of this work would be to propose an application for collectorless flotation rather than to explain it.

Section 2.2 describes the main metallurgical problem facing metallurgists at Mattabi Mines namely copper/lead separation. In all tests so far performed on Mattabi and Lyon Lake ores, strong collectorless flotation of chalcopyrite has been accompanied by slover but significant flotation of galena. A process using collectorless flotation is proposed by Figure 5.2.

While collectorless flotation of chalcopyrite may follow several mechanisms (55,99,107-8), that of galena is likely to be the result of formation of elemental sulphur or some other oxidation product on the mineral surface.(99,102) Figure 4.12 compares the effect of floating galena without collector in reducing conditions (using nitrogen) and oxidising conditions (using air). It confirms that oxidising conditions aid the flotation of galena and so supports the assumption that it must be oxidised to float.

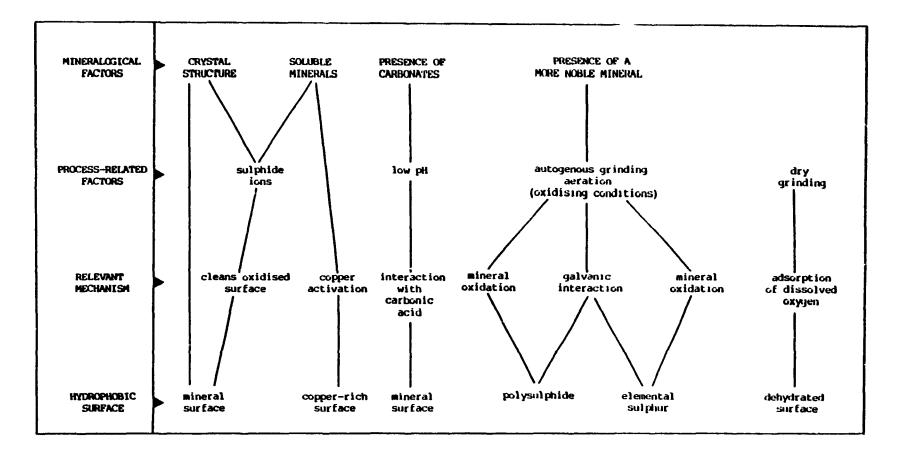


FIGURE 5.1 - Summary of mechanisms behind collectorless flotation as described in Section 2.1.

FIGURE 5.2:Proposed process leading to the production of selective copper and lead concentrates including use of collectorless flotation.

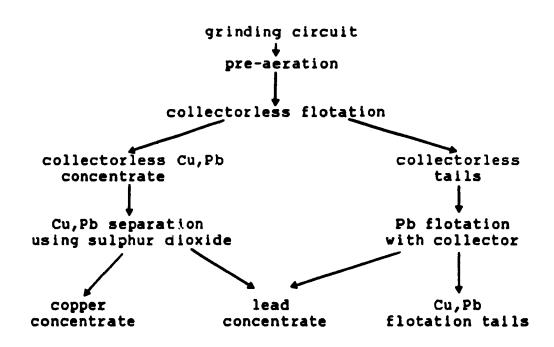


Table 5.1 further shows that oxygen is important, by relating the degree of oxidation of the galena to (1), its collectorless floatability and (2), its susceptibility to sulphite depression:

J

TABLE 5.1: Effect of degree of oxidation on collectorless flotation and response to sulphite depression.

	degree	of	oxidation	
	high		low	
collectorless floatability	good		poor	
response to sulphite depression	good		poor	

Thus collectorless flotation is a means by which chalcopyrite and oxidised galena can be separated from clean galena. Table 5.1 also describes that the galena most difficult to depress by

sulphite depression is clean galena (91) a fact supported by several aspects of the metallurgy at Mattabi Mines.

- * The collectorless flotation of galena is slow, suggesting that galena in the Mattabi ores oxidises only slowly. So much of the galena in the copper/lead separation stage is clean, explaining the poor separability of copper and lead using sulphur dioxide.
- * Oxidation of galena would be accelerated by raising the pulp temperature which may help to explain the positive effect of steam.
- * Copper/lead separation suffers if the pulp is left to age.
 Oxygen demand from the pyrite will tend to make conditions
 more reducing as the pulp ages. This may clean the galena
 of its oxidation products and reduce copper/lead
 separability. This was observed during on-site column
 flotation studies in 1986 (21).

Galena floating with the chalcopyrite in the collectorless concentrate is oxidised and will be effectively depressed using sulphur dioxide. The clean galena left unfloated during collectorless flotation will float well using starvation quantities of collector. Aided by the extra aeration provided by the collectorless float, pyrite will remain depressed with the sphalerite and a high-grade galena concentrate should result, which can be combined with the copper/lead separation tails.

The viability of the process hinges on the consistency of the collectorless flotation of chalcopyrite. It is hoped that regular collectorless flotation tests will be performed on cyclone overflow by personnel at Mattabi Mines to check on the reliability of the mechanism.

As galena flotation to the collectorless concentrate is apparently dependent on oxidising conditions, addition of a reducing agent (such as sodium sulphide) may improve copper collectorless flotation selectivity. Extensive testwork by Yoon and co-workers has shown that sodium sulphide can also promote the collectorless flotation of chalcopyrite (58-9,107-8,99).

5.2 Pyrite flotation using nitrogen

5.2.1 Theoretical aspects.

The initial object of the testwork using nitrogen was to assess whether the use of nitrogen would prevent flotation of pyrite by reducing the pulp potential and the dissolved oxygen content thereby blocking the pyrite-collector interaction.

Comparison of Figures 4.8 (for pyrite flotation in air) and 4.9 (for pyrite flotation in nitrogen) in Section 4.1.8 shows that:

- a) Nitrogen does slowly reduce the pulp potential
- b) Nitrogen very quickly reduces the dissolved oxygen content, but
- c) Nitrogen strongly promotes the flotation of pyrite.

When considering only the collector-mineral interaction,

observations (a) and (b) seem to contradict (c). According to the electrochemical theory of mineral-collector interaction, more reducing conditions should tend to send equilibria 5b and hence send 5a to the left (24,60-1) This infers that the mineral-collector reaction would be retarded.

$$MS + X - = MS(X) + e -(5a)$$

$$H_2^0 + 1/2.0_2 + 2.e- = 2.(OH-).......(5b)$$

Without analysing the mineral surface it is not possible to confirm this.

So another mechanism is altering the hydrophobicity of the mineral surface. With only a fraction of the surface covered by a collector, the hydrophobicity of a mineral surface is likely to be a balance between the amount of collector adsorbed and the hydrophobicity of the mineral surface without adsorbed collector. This balance is particularly applicable to pyrite. Gaudin (28) suggested that a pyrite surface is constantly changing, through formation of hydrophilic oxides of sulphur and possibly iron. Fuerstenau attributed the depression of pyrite with permanganate to the adsorption of hydroxyl ions onto the pyrite surface through reaction with ferric ions in the mineral. (23). Hoyack and Raghavan(43) proposed a similar mechanism for the depression of pyrite using sulphite ions.

The possible role of galvanic interactions altering the mineral surface has been proposed by a number of investigators (40,44,49,79). Galvanic interactions have been used to explain

effects of grinding media (1,14,44,51,79,97) on flotation performance. Similar galvanic effects are likely to occur between pyrite, the most noble mineral, and other base metal sulphides (49,60,75-9). A model of such a mechanism is illustrated by Figure 2.3. Pyrite, being more noble, will tend to draw electrons from sphalerite. This will oxidise sphalerite and promote formation of iron hydroxides on the pyrite surface through reduction of dissolved oxygen. Majima (60) notes that such a galvanic interaction has been found to increase the rate of oxidation of a sulphide mineral through association with pyrite by eight to twenty times. This leads to the formation of hydrophobic, elemental sulphur on the mineral surface.

Such oxidation will aid sphalerite flotation and hydrophilic iron hydroxides on the pyrite surface will depress pyrite. So the effect would be to improve selective flotation of sphalerite from pyrite.

By stopping the supply of oxygen, nitrogen flotation blocks the mechanism. Iron hydroxides will no longer form on the pyrite surface. The increased hydrophobicity of the pyrite will then allow flotation of pyrite with only minimal dosages of collector.

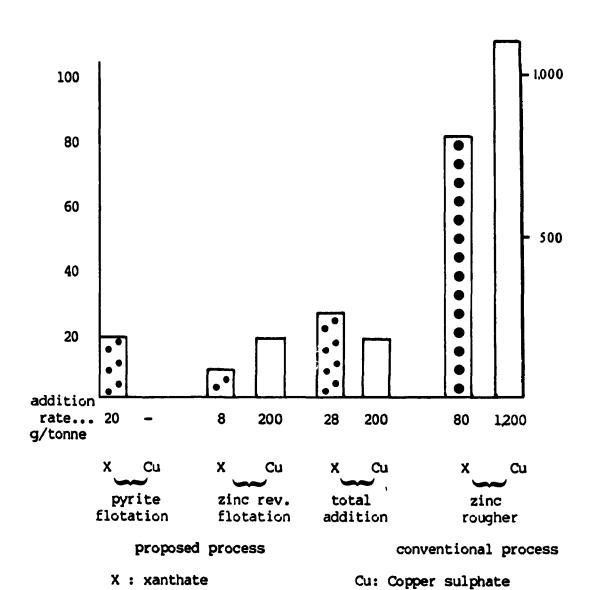


FIGURE 5.3 - Exposure of pyrite to collector and copper sulphate activator, in the conventional and proposed processes.

5.2.2 Operational Aspects.

The results section showed that a relatively high grade pyrite concentrate could be floated from the bulk flotation tails. It also showed that entrained zinc could be selectively recovered from the pyrite concentrate by reverse flotation with collector following copper activation, in air and at pH 10-11 (although other pH levels were not tested). An overall metallurgical balance for the proposed process is illustrated by Figure 4.7.

It shows that the proposed process produced a zinc rougher concentrate grade 50% higher than that produced by the conventional route, at a similar recovery. While this is unlikely to improve the overall grade/recovery relationship of a circuit it may make the cleaner circuit simpler and easier to operate.

Inspection of Figure 5.3 reveals that the proposed process largely avoids the contact of pyrite with copper sulphate. While the potential for copper sulphate activation of pyrite is not well recognised in North America, Yamomoto (106) notes this use and copper sulphate is used to activate pyrite in flotation circuits in South Africa (67). Pyrite would also be less exposed to collector. This would suggest several advantages:

- * Easier overall pyrite rejection (indicated by the improved zinc rougher grades).
- * Less reliance on pH for pyrite rejection (with the resultant saving in lime costs).

* Reduced consumption of copper sulphate and xanthate by eliminating their wasteful adsorption onto the pyrite surface.

The nature of the zinc reverse flotation concentrate suggests that it may isolate locked particles. This would make it a particularly good stream to regrind.

The main problem associated with the use of the proposed flowsheet is the supply of nitrogen. Based on laboratory tests, 400 tonnes of fresh nitrogen gas will be needed per day for a plant the size of Brunswick Mining (treating 10,250 tonnes per day). For both economic and environmental reasons, some form of recirculation system will be needed (i.e. closed cells). In this case air could be used as the carrier gas, oxygen being quickly consumed by the pyrite. Some nitrogen supply may be necessary to ensure a positive pressure in the flotation cell, stopping air from leaking into the system, but this is unlikely to be more than 20 tonnes per day. Conditioning could be performed in a closed tank. Conditioning gas may not be necessary. If it is then the gas can be recirculated in an enclosed system. The supply of extra nitrogen could be manipulated automatically by dissolved oxygen levels at the end of pyrite flotation. Tests so far suggest that the dissolved oxygen level should be below 2 ppm.

Another aspect of the effect of nitrogen, or oxygen deficient gases on the flotation of pyrite concerns the use of flotation columns. Calculations outlined in Appendix 5 suggest that the

amount of air injected into a column is not enough to satisfy the demand for oxygen from a pyritic ore. Therefore, as air rises through the cell it becomes deficient in oxygen and will tend to increase the floatability of pyrite. So, while this needs to be checked it suggests that the geometry of the cell could affect the chemical environment of the pulp, the flotation chemistry and the grade/recovery relationship of a flotation process. This may limit the use of columns in processing streams which have a high oxygen demand.

5.2.3 Economics of the process (Balance based on Brunswick Mining)

As mentioned in the previous section, use of fresh nitrogen cannot be economically justified. The capital cost of the plant would be in the region of \$8 million and the running costs \$16,000 per day.

So the balance developed in this section is based on 20 tonnes of nitrogen a day, enclosing the flotation cells and circulating the gas. At Brunswick Mining the cells (Outokumpu) are virtually enclosed already and it is assumed the gas recirculation system can be home-built at negligible cost.

The cost of a nitrogen plant to produce 20 tonnes of nitrogen a day is approximately \$750,000. Its running costs are \$800 per day. Based on straight-line depreciation and not including tax considerations, the capital cost over a 15 year period is equivalent to 1.3 cents per tonne of ore. Running costs are equivalent to 7.6 cents per tonne of ore. The total cost of the

nitrogen is therefore 8.9 cents per tonne.

Figure 5.4 compares approximate reagent costs in a zinc circuit with the cost of nitrogen. It shows that the cost of 20 tonnes/day nitrogen can be justified by a 12% saving in lime or 19% saving in copper sulphate consumption. The cost of nitrogen is unlikely to be justified by savings in collector (requiring a 57% saving).

Operating costs (power, maintenance etc.) of the zinc flotation circuit are normally about 60 cents per tonne of ore (2,42). The use of nitrogen can then be justified by a 15% reduction in overall zinc flotation capacity.

At this stage, it is not possible to predict that nitrogen flotation is economical. The cost comparisons listed above do, however, show that the process is relatively cheap in comparison to other flotation costs. This suggests that any improvement in zinc/pyrite separation efficiency which results in the use of lower pH and less reagents will probably justify the extra costs of the process. The process could be particularly attractive to more isolated operations where transportation of lime and copper sulphate becomes very costly (88).

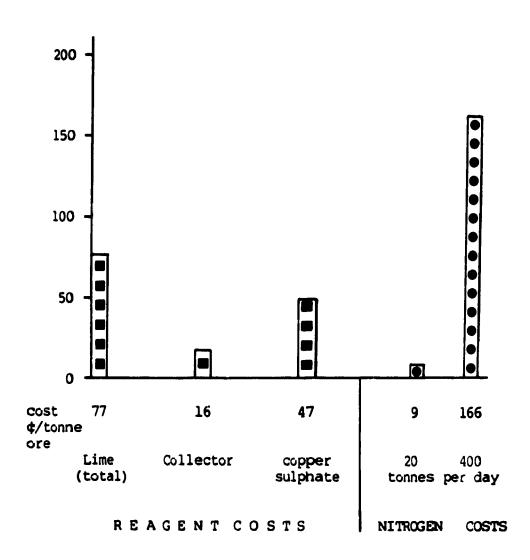


FIGURE 5.4 - Comparison of approximate plant reagent costs with expected costs of using nitrogen (based on 400 and 20 tonnes per day).

6. CONCLUSIONS

6.1 COLLECTORLESS FLOTATION

- The collectorless floatability of chalcopyrite varied from ore to ore, though all ores tested showed some response to collectorless floatation. The collectorless recoveries of chalcopyrite in ores from Mattabi are similar to those using collector.
- Some collectorless flotation of galena was observed from ores from Mattabi Mines, particularly under oxidising conditions.
- * A process is proposed whereby chalcopyrite and oxidised galena are floated without collector. These are subsequently separated by sulphur dioxide depression of galena. The remaining untarnished galena is floated with collector.
- * Pyrite and sphalerite showed no response to collectorless flotation in all ores tested.

6.2 NITROGEN PLOTATION.

- * The use of nitrogen strongly promotes the flotation of pyrite. Pyrite flotation is selective from zinc-bearing minerals but not from copper- and lead-bearing minerals.
- * Nitrogen conditioning and flotation reduces the pulp potential. The level of dissolved oxygen in solution drops to below 0.5 ppm. within 1 minute, through the use of nitrogen.
- * Pyrite flotation is probably linked to the removal of dissolved oxygen. This blocks a galvanic interaction between pyrite and sphalerite, which otherwise serves to depress the pyrite through OH- formation.
- * Nitrogen flotation can be used to float pyrite after bulk copper, lead flotation and before zinc flotation. This improves the rougher concentrate grade achieved by subsequent zinc flotation.
- * Aerating the pyrite concentrate pulp depresses the pyrite Copper activation and reverse zinc flotation with collector in air selectively recovers zinc minerals from the pyrite concentrate.
- * The proposed process incorporating nitrogen flotation and reverse flotation of zinc-bearing minerals reduces the exposure of pyrite to collectors and copper sulphate. This may explain the improved zinc rougher grades achieved. The

process may also isolate locked zinc-pyrite particles in the reverse concentrate, in which case regrinding could liberate extra zinc minerals.

* The process is probably viable economically at 20 tonnes nitrogen per day if savings in lime of 17% are realised.

7. RECOMMENDED FURTHER WORK

7.1 COLLECTORLESS FLOTATION

7.1.1 To assess the potential of the proposed process:

- Regular on-site collectorless flotation tests should be performed to check the reliability of collectorless flotation.
- Copper, lead separation of the collectorless flotation concentrate should be investigated. Galena should be depressed using sulphur dioxide.

7.1.2 To improve on the process:

- As collectorless flotation of gaiena seems to be based on surface oxidation, collectorless flotation with a reducing agent (e.g. sodium sulphite) may result in a selective copper concentrate.
- Alternatively, flotation recovery of galena could be maximised. While galena floats slowly without collector, it may be selectively recovered from pyrite and sphalerite if allowed to float to the limit of recovery.

7.2 NITROGEN PLOTATION

7.2.1 To increase understanding of nitrogen flotation.

- Nitrogen testwork should be performed on Mattagami ore, which contains mainly pyrrhotite. Pyrrhotite is less noble

than pyrite but has a higher oxygen demand.

- A flotation column should be used to check in whether the mis-direction of zinc is the result of entrainment.

7.2.2 To assess commercial potential of nitrogen flotation.

- One of the potential rewards of removing pyrite before zinc flotation is a saving in lime costs. To check this, flotation of a zinc concentrate after pyrite flotation under less alkaline conditions should be tested.
- Continuous tests should be performed to optimise pyrite flotation.
- The process is only likely to be viable if oxygen deficient gas is produced by enclosing the cells and recirculating the gas. The production of sufficient 'fresh' nitrogen is likely to be too expensive. Tests should be run using an enclosed cell to establish whether pyrite can be promoted in this way.
- Mineralogical analysis of the reverse zinc concentrate should be undertaken to identify the concentration of locked particles in the sample.
- Nitrogen should be tested on the pyrite reverse flotation stage which is currently in some zinc circuits.(e.g Kidd Creek Hines).
- Most ways of producing nitrogen also produce oxygen. As a

logical extension of nitrogen flotation to float pyrite and aeration to depress pyrite, oxygen-enriched aeration could improve pyrite rejection in the copper, lead bulk float. The use of oxygen enriched gases during pre-aeration and bulk flotation should be tested.

7.2.3 Other related testwork.

- Tests should be run which check the dissolved oxygen content and perhaps pulp potential of the pulp at different levels in a flotation column, treating an ore with a high oxygen demand.

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9: APPENDIX 1

GRINDING TEST DATA

Brunswick Mining	vet	sieve ana	l ys is				
grind time	per	cent pass	ing				
(minutes)	7	4 µm	37µm				
20	8	3	44				
30	9	5	67				
40	10	0	92				
Mattabi Mines	vet	sieve ana	lysis				
grind time	per	cent pass	ing				
(minutes)	300µm	150µm	74µm				
. 30	100	100	95				
20	100	99	77				
10	98	86	48				
Kidd Creek Mines	vet	sieve ana	lysis				
grind time	per	cent pass	ing				
(minutes)	7 4 µ	m	45µm				
20	73		20				
30	76		64				
40	79		71				

10. APPENDIX 2 - BRUNSWICK MINING ORE TESTWORK

COLLECTORLESS FLOTATION

- 10-101 Collectorless flotation in air, followed by bulk copper, lead flotation and zinc flotation.
- 10-102 Collectorless flotation in nitrogen, followed by bulk copper, lead flotation and zinc flotation in nitrogen.
- 10-103 Collectorless flotation then zinc flotation in nitrogen.
- 10-104 Collectorless flotation then zinc flotation in air.

NITROGEN FLOTATION

- 10-201 Bulk copper, lead flotation in air, pyrite flotation in air and nitrogen, and zinc flotation in air.
- 10-202 Bulk flotation in air, pyrite flotation in air and zinc flotation in air.
- 10-203 Bulk copper, lead flotation in air, pyrite flotation in air then nitrogen and zinc flotation in air.
- 10-204 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.
- 10-205 Bulk copper, lead, pyrite flotation in nitrogen with stage addition of collector, then zinc flotation in nitrogen.
- 10-206 Bulk copper, lead, pyrite flotation in nitrogen, single addition of collector, zinc flotation in nitrogen.

- 10-207 Copper activation then zinc flotation in nitrogen only.
- 10-208 Bulk copper, lead, pyrite flotation in nitrogen, zinc flotation in nitrogen.
- 10-209 Nitrogen conditioning, then copper, lead, pyrite flotation in air and zinc flotation in air.
- 10-210 Nitrogen conditioning, then copper, lead, pyrite flotation in air, folowed by zinc flotation in air.
- 10-211 Bulk copper, lead, pyrite flotation in nitrogen then zinc flotation in air.
- 10-212 Bulk copper, lead flotation in air, then pyrite flotation in nitrogen with zinc reverse cleaning, finally zinc flotation in air.
- 10-213 Bulk copper, lead flotation in air, then pyrite flotation in nitrogen with zinc reverse flotation, finally zinc flotation of pyrite tails in air. (repeat of test 10-212)
- 10-214 Bulk copper, lead flotation in air, pyrite flotation in nitrogen, zinc flotation in air.
- 10-215 Bulk copper, lead flotation in air, pyrite flotation and zinc flotation in air.

STANDARD PLOTATION

10-301 Standard copper, lead flotation and zinc flotation in air.

PLOTATION TESTS ON-SITE AT BRUNSWICK MINING.

- 10-401 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation.
- 10-402 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.
- 10-403 Bulk copper, lead flotation in air, pyrite flotation in nitrogen with zinc reverse cleaning and zinc flotation of pyrite tails in air.
- 10-404 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air. (McGill Std procedure).

ARGON FLOTATION

10-501 Bulk copper, lead flotation in air, pyrite flotation in argon and zinc flotation in air.

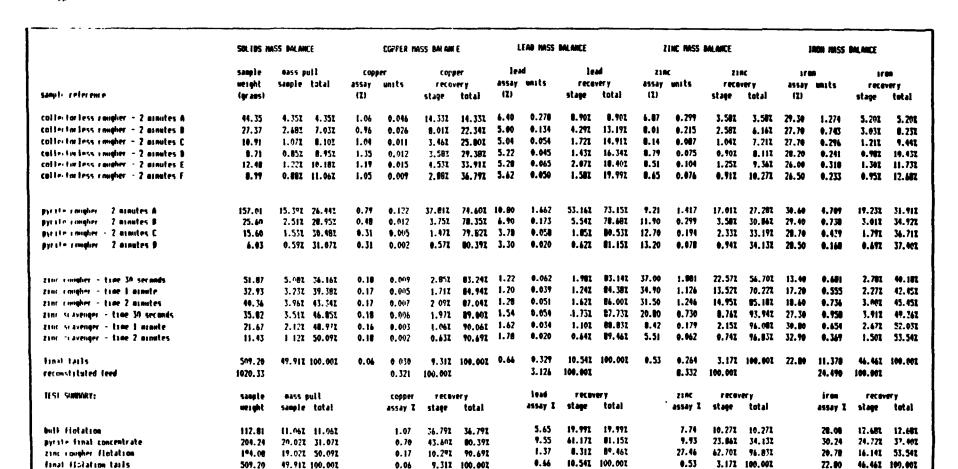
	SOI 195 N	ASS DALANC	E		COPPER IM	SS BALANC	ŧ	1	LEAD MASS	BALANCE		1	INC MASS I	MLANCE		1	non mass 1	M. MICE	
	sample	0455 PK	li	Copi	er	(099	er	le.	ıđ	lea	nd .	zia	c	210	c	170		ire	-
	nesilyt	sample	total	assay	units	recov	ery	assay	wits	reco	rer y	8558Y	wits	recov	er v	assay	mits	/ ecov	
sample reference	-			(2)		stage	total	(2)		stage	total	(2)		stage	total	(2)		stage	total
collecter less raughe 2 ainutes A	40.45	4.412	4 412	0.30	9.017	4.991	4.992	5.20	0.233	7.092	7.092	6.0 1	0.354	4.102	4.102	27.70	1.223	4.762	4. 76
collectories cougher - 2 acoutes 8	17.20	1.072	6.30I	0.38	0.007	2.13Z	7.122	4.89	0.071	2.75 t	7.04Z	7.07	0.171	1.992	4.091	24.70	0.503	2.041	7.00
collect riless cougher - 2 acoutes C	12 99	1.421	7.721	♦.36	0.005	1.521	8.642	4.42	0.045	1.992	11.032	7.24	0.131	1.52%	7.412	25.00	9.366	1.491	8.40
celler liv less rougher - 2 minutes D	7.67	1.06Z	8.77%	0.36	9.004	1.132	9.772	4.36	9.044	1.40Z	13.232	7.01	0.075	1.111	0.72%	24.60	0,260	1.051	7.54
collector less raugher - ? asnutes E	6.83	0 752	9.521	0.37	0.003	0.02I	10.591	4.42	0.033	1.00I	14.232	7.32	0.047	9.01Z	9.52%	24.80	0.195	0.751	10.29
collectoriess rougher - 2 minutes f	6.47	9.732	19.75%	C.37	9.003	0.801	11.391	4.30	●.03l	0.951	15.101	7.12	0.064	0.772	10.291	24.30	0.177	0.771	11.00
collectorless rougher - 7 aroutes 6	4.05	0 442	10.672	0.37	0.00?	0.491	11.001	4.40	0.019	0.571	15.771	7.52	0.012	0.472	10.702	24.90	0.119	0.45I	11.45
pyrate emiglier - 2 assutes A	132.07	14 412	25.092	9.47	0.068	20. 14I	32.021	6.44	0.928	20.211	43. 98 2	2.11	0.304	3.532	14.312	39.40	5.532	22.432	33.00
pyrate cougher - 2 acoutes 0	84.52	7.41L	34.542	0.45	0.042	12.642	44.652	4.30	9.406	12. 34Z	54.322	2.60	0.245	2. 0 5%	17.162	37.40	3.729	15. 00 7	48.17
pyrate complex Zazautes C	54.98	5 992	49.571	9.51	0.031	7.00Z	53.74%	3.84	0.730	4. 99Z	63, 312	3.02	O. 181	2. 10E	17.261	37.40	2.371	7.672	50.59
pyrate rangher - 2 ainutes D	55.00	4.011	46.532	0.50	0.030	0.742	42.48I	3.20	0.197	5.992	40.312	3.15	0.187	2.201	21.462	37.00	2.392	•.791	40.20
zene raugher - Lene 30 serands	84.70	9.761	55. 8 92	0.49	0.044	13.232	75.90Z	4.72	0.437	13.291	e2.402	19.30	1.695	17.68Z	41.142	20.10	2.603	10.561	78.94
eine reugher time 30 seconds	46 43	5 072	60.86I	♦.39	9.079	5.001	81.7 8 1	2.50	0.127	3.052	86 45Z	25.40	1.207	14.742	56.072	21.40	1.084	4.401	03.23
zine empler time I minute	20.59	2.25%	43.112	0.37	0.008	2.472	84.75%	2.04	0.046	1.412	87.841	30.40	0.483	7.932	44.021	17.30	0.474	1.761	84.77
zam siavenner - time 30 seronds	49.08	5. 161	60.471	0.34	0 017	5.731	89.991	3.08	0.165	5.017	92 872	33.00	1.767	20.52%	84.542	10.40	0.785	4.001	80.71
zinc scarenger - time i minute	31.32	3 472	71.002	0.34	0.012	3.462	93.44Z	2.04	0.070	2.121	94.971	24.90	0.051	7,002	94.421	10.50	0.632	7.54I	71.55
zime seavenger - time 7 mimites	0.65	0.742	72.831	0.32	0.003	0.90I	94.34Z	1.90	0.018	0.55z	95.54 %	16.40	0.155	1. 9 07	94.212	17.20	0.181	0.732	92.29
final tails	249 03	27.171	100.00I	0.07	0.019		100.001	0.54	0.147		100.001	1.20	0.326		100.007	7.00	1.902		100.00
recoustilled feed	916.46				0.334	100.001			3.209	100.001			8.412	100.001			24.660	100.001	
TEST SIMMARY:	sample	0255 P			copper	recov			lead	recev	- •		2180	LECOM			1700	recev	
	weight	saople	tetal		assay I	stage	total		assay 1	stage	total		assay l	st age	totał		assay I	stage	tetai
Duth flutation	93.41	10 672			0.37	11-802			4.85	15.771	15.77%		0.47	10.702	10.701		26.42	11.451	
pyrite final commentrate	328.50		46.532		0.48	50.801	62.691		4.91	53.532	69.317		2.57	10.60I	21.462		37.10	56.832	60.29
zinc rimpher flotation	244.47	26.271			0.40	31.472			3.21	26. 231	95.541		23.90	74.75%	76.212		21.03	24.012	
final flotation tails	249.03	27.171	100.00Z		0.07	5.661	100.001		0.54	4.462	100.001		1.20	3.742	100.00I		7.00	7.71%	100.00

10-101 Collectorless flotation in air, followed by bulk copper, lead flotation and zinc flotation.

FLOTATION CONDITIONS

	pH	activator	Pre-condi	tianing	coll	ector	flotation
		added			type	add1t1on	gas
		g/t	gas	time		g/t	•
collectorless rougher - 2 minutes A	8.5	-	nıtrogen	25 s ın	-	•	nitracen
collectorless rougher - 2 minutes B	8.5	-	•	-	-	_	nitrogen nitrogen
collectorless rougher - 2 minutes C	8.5	-	-	_	_	_	_
collectoriess rougher - 2 minutes D	8.5	-	•	-	-		nitrogen
callectoriess raugher - 2 minutes E	8.5	-	-	_	_	-	nitrogen
collectoriess rougher - 2 minutes F	8.5	-	•	_	•	_	nitrogen
collectorless rougher - 2 minutes 6	8.5		-	-	_	_	nitragen
•						-	nitragen
pyrite rougher - 2 minutes A	8.5	-	-	-	SIPY	50	nitragen
pyrite rougher - 2 minutes B	8.5	-	-	-	-	_	n:trogen
pyrite rougher - 2 minutes C	8.5	-	-	-	-	-	nitrogen
pyrite rougher - 2 minutes D	8.5	-	-	•	-	•	nitragen
zinc rougher - 30 seconds	10.0	1200	none	5 a in	SIPX	50	nitrogen
zinc rougher - 1 minute	10.0		•	-	-	-	nitrogen
zinc rougher - 2 minutes	10.0	-	-	-	•	-	nitrogen
zanc scav enger - 30 seco nds	10.0	_				- 4	- •
zinc scavenger - I minute	10.0			-	SIPX	30	nitrogen
zinc scavenger - 2 minutes	10.0		-	-	-	-	nitragen
The Scavenger - 2 Bindles	10.0	-	-	-	-	-	nitrogen
			GRINDING	CONDITIO	NS:		
			30 minut	es. mild	steel c	harge	
			1200 a/t	: 21NC SU	lohate		

10-101 Collectorless flotation in air, followed by bulk copper, lead flotation and zinc flotation.



10-102 Collectorless flotation in nitrogen, followed by bulk copper, lead flotation and zinc flotation in nitrogen

FLOTATION CONDITIONS

Sample reference	pН	activator added	Pre-con	ditioning		ector	flotaties
John College		g/t	gas	time	type	addition g/t	ģas
collectorless rougher - 2 minutes A	8.5	•	air	25 mir	-	•	alr
collectorless rougher - 2 minutes 8	8.5	-	_	•	-	-	317
collectorless rougher - 2 minutes C	8.5	-	-	-	-	-	alr
collectorless rougher - 2 minutes D	8.5	-	-	-	-	•	215
collectorless rougher - 2 minutes E	8.5	•	•	-	-	-	air
collectorless rougher - 2 minutes F	8.5		-	-	-	-	alr
pyrite rougher - 2 minutes A	8.5	•	-	•	SIPI	50	alr
pyrite rougher - 2 minutes B	8.5	-	-	-	-	-	217
pyrite rougher - 2 minutes C	8.5	-	_	-	_	-	alr
pyrite rougher - 2 minutes D	8.5	-	•	-	-	-	air
zinc rougher - 30 seconds	10.0	1200	none	5 min	SIPI	50	alf
zinc rougher - 1 minute	10.0		-	-	-		air
zinc rougher - 2 minutes	10.0	-	-	-	-	•	air
zinc scavenger - 30 seconds	10.0	. -	-	-	SIPI	30	air
zinc scavenger - 1 minute	10.0	-	-	-	-	-	215
zinc scavenger II	10.0	•	-	-	-	•	air

GRINDING CONDITIONS:

30 minutes. mild steel charge

1200 g/t zinc sulphate

10-102 Collectorless flotation in nitrogen, followed by bulk copper, lead flotation and zinc flotation in nitrogen

	SOLIDS IN	ASS BALAM	CE		COPPER N	ISS PALAM	E	LEAD MASS DALMICE				1	INC MASS I	BAL ANCE		IRON MASS BALANCE				
	sample	8855 p		cap	107	C 000	per	ìe	M .	1e	46	210	•	210	AC	ire		ire		
Sample reference	we i ght	Sample.	total	4224 A	units	reco stage	v er y total	85589	units	rece	very total	455 8 y	units	rece stage	very tetal	8558Y	units	recen	very total	
						31000	(0(8)			2.092				2444	(8/41			ar after	(0.0	
collectoriess rougher - 2 minutes	54.3*	5.782	5.781	0.54	170.0	7. 301	9, 302	4.34	0.344	11.742	11.741	7.00	0.407	5.472	5.492	34, 20	1.977	7.051	7.05	
collectivitess scavenger ~ 2 minutes	25.78	2.642	8.422	0.71	0.01*	5.592	14.90Z	5.04	0.133	4.261	16.001	0.19	0.216	3.012	0.492	32.70	0.064	3.001	10.13	
the employ - time 30 seconds	35 15	3.601	12.021	1.20	790.0	12.091	27.782	7.52	0.271	8.481	24.681	37.99	1.345	18.971	27.472	i1.40	0.411	1.41	11.4	
rinc rougher - time 1 minute	38.99	3.901	15.932	1.12	0.044	13.032	40 872	7.20	0.359	11.501	76 1 9 2	30.90	1.206	16.762	44.43	15.30	0.577	2.132	13.7	
zanc rimigher - tame 2 maintes A	46.70	4.172	20.10X	0.90	0.038	11 192	52.012	0.06	0. 70	11.047	48.071	74.50	1.105	15.342	39.79%	10.70	9.780	2.701	14.5	
inc rimpher - time 2 minutes 0	13.86	1.422	21.52%	9.70	0.013	3.011	55.022	6.06	9.084	2.761	50.771	22.10	0.314	4.361	44.152	21.20	0.30i	1.07%	17.3	
tinc sravenger - time Th seconds	99. 74	10. 182	31.702	● 50	0.051	15.101	71.00I	6.04	0.415	19.701	70.471	17.50	1.702	24.76%	90 712	29.99	2.943	10.501	28.6	
inc scavenger - time 1 minute	54 57		37.301	0 41	0.023	6.84Z	77 812	J	0.172	5.521	75.901	5.40	0.302	4.201	73.112	37.00	7.101	7.701	33.0	
anc reavenger - time 2 minutes A	66.84		44.15%	0.25	9.017	5.112	87.942	7.00	0.137	4.391	80.371	2.40	9. 164	2.291	93.392	42.00	2.877	10.261	44.1	
rinc scavenger - time 2 minutes B	7.04	0.932	45.072	0.25	0.092	9.472	03 A32	2.12	0.020	0.432	81.00I	7.61	0.024	0.341	95.731	41.50	9.305	1.371	47.4	
inal tails	525.90	54.931	100.001	0.10	0.055	16.371	100,001	1.00	0.593	17.00Z	100.001	0.54	0.308	4.27%	100.001	26.90	14.720	52.511	100.0	
econstituted feed	975.66				0.335	100.001			3.125	100.001			7.197	100.001			20.036	100.002		
TEST SIMMARY:	sample	0355 (wil		c opper	recev	ery		lead	recov	er y		ZIRC	FPCOV	er y		17 0 0	recev	e ry	
	wei ght	samte	totai		assay l	stage	totai		assay X	st age	total		assay I	stage	total		assay l	stage	tota	
collectorless flotation	82.17		0.42Z		0.57	14.902	14,901		5.93	16.001	16.001		7.43	8.472	8.402		33.73	10.132	10.1	
conc rougher flotation	127.80		21.521		1.05	40.772	55.022		0.29	34.771	50.771		30.47	55. 462	64. 15 1		15.95	7.452	17.3	
tinc scavenger flotation time 0-1.5 min	153.91	15.772	37.302		0.47	22.011	77.831		4.77	25.211	75.991		13.71	20.762	73.112		32.40	19.791	35.8	
zinc scavenger flotation time 1.Smin+	75.88	7.792	45.071		0.25	5. BOZ	93.432		2.01	5.021	B1.00%		2.43	7.621	95.732		41.94	11.432	47.4	
final flotucion taits	535.90	54.932	100.002		0.10	16.372	100,001		1.00	17.00Z	100.00I		0.54	4.271	106.001		26.00	52.511	100.0	

10-103 Collectorless flotation then zinc flotation in nitrogen.

FLOTATION CONDITIONS

Sample reference	pH		Pre-condi	tioning	coll	ector	flotation
semple reverence		added g/t	gas	tine	type	addition g/t	gas
collectorless rougher - 2 minutes collectorless scavenger - 2 minutes	8.0 8.0		air	10 min	-	-	air
-	5.0			_	•	-	air
zinc rougher - 30 second	10.0	1200	nitrogen	20 min	SIPX	50	nitrogen
zinc rougher - 1 minute	10.0	-	-	-	-	-	nitrogen
zinc rougher - 2 minutes A	10.0	, -	-	-	-	-	nitrogen
zinc rougher - 2 minutes B	10.0	-	-	-	-	•	nitrogen
zinc scavenger - 30 seconds	10.0	-	-	-	SIPX	30	nitrogen
zinc scavenger - 1 minute	10.C	-	-	-	-	-	nitrogen
zinc scavenger - 2 minutes A	10.0	-	-	-	-	-	nitragen
zinc scavenger - 2 minutes B	10.0	-	-	-	-	-	nıtrogen

GRINDING CONDITIONS:

30 minutes, stainless steel charge

1200 g/t zinc sulphate

	SGLIDS MASS BALANCE				COPPER MASS BALANCE				LEAD MASS BALANCE					BALANCE		IRON MASS DALANCE				
est number 224 sample		2255 a	eass pull		per	COGI	ner	le	ıd	lead		ZINC		ZINC		100	_	ires		
	meraht	Sample		8558Y	units	reco		45587	units	reco			units	reco			units			
sample reference		200611		4224,	4.11.		total	•		stage	total	,		_	total	e236y	m1(2	stage	very total	
collectorless reweher - 2 ainstes	69.26	6.781	6.782	€.61	0.041	12.472	12.472	4.14	0.418	13.112	13.112	7.36	0.477	7.00Z	7.007	34.00	2.306	8.142	8,14	
collectorless scavenger - 2 minutes	24.48	2.431		0.81	0.020	5.94Z	19.417		0.125	3.921	17.031	7.07	0.221	3.072	10.071	32.40	0.700	2.781	10.73	
zinc raugher - time 30 seconds	74 89	7.44%	14.662	0.92	9.068	20.641	39.051	7.14	0.680	21.342	30.371	35.40	2.449	37.142	47.23Z	13.00	0.947	3,421	14.34	
zanc congher - tage I azaute	67.36	4.497	23.35%	0.86	0.058	17.35%	56.40X	8.76	0.584	18.402	56.771	27.90	1.867	26.181	73.402	10.00	1.205	4. 251	18.57	
zinc rayalor - time 2 minutes A	50.33		29 142	0.74	0.043	12.931	49.331		0.386	12.112	68.88%	17.40	1.020	14.30X	87.762	24.80	1.553	5. 481	24.00	
zinc rougher - time 2 minutes B	21.18		31.252	0.57	0.012	3.74%	73. 00 %	5.60	0.105	3. 30Z	72.101	10.20	0.215	3.01Z	90.712	31.00	₹.652	2.301	26.30	
zinc scavenger - time 30 seconds	43.53	4.332	35.572	0.54	0.023	7.042	8 6.121	5.20	0.228	7.171	79.35I	7.15	0.309	4.34Z	95. 0 51	35.30	1.527	5,392	31.77	
zinc scavenger - time 1 minute	24.83	2.672	38.24Z	0.39	0.010	3.132	93.25%	3.50	0.095	2.991	82.342	3.14	0.084	1.17%	94.221	38.00	1.013	3.502	35. 35	
zanc scavenger - time 2 minutes A	17.14		90.14Z	0.30	8.004	1.72%	24.971	2.78	0.053	1.462	84.001	2.24	0.043	0.40Z	96.822	30.30	0.720	2.572	37.92	
zinc scavenger - time 2 minutes B	11.64	1.162	41.302	0.25	0.003	0.872	8 5. 842	2.46	0.028	0. 0 92	8ª. 70%	1.64	0.022	0.30I	97.121	38.00	0.439	1.551	39.47	
final tails	590.00	58.70 z	100.001	9.00	0.047	14.162	100.001	v. 82	9.401	15. 10Z	100.00I	0.35	0. 205	2.002	100.00I	29.20	17.141	60.532	100.00	
reconstituted feed	1006.44				0.332	100.001			3.107	100.00Z			7.133	100.001			20.320	100.001		
TEST SUMMARY:	sample weight	aass (copper assay I	recov stage	ery total		lead assay I	recov stage	ery total		ZIRC assay I	recev			irm	recev		
	merdur) make 1.5.	(W/a)		essey .	arada	totai			•			assay 4	stage	total		assay 1	stage	total	
collectorless flotation	92.74	9.211	7.211		9.66	10.417	18.41Z		5.87	17.032	17.032		7.81	10.072	10.091		33.50	10.932	10.73	
zinc rougher flotation	221.76	22.93Z	31.25%		♥.#2	54.662	73.09Z		7.78	55. 151	72.181		26.10	80.421	90.712		17.87	15.442	24.30	
zinc scavenger flotation time 0-1.5 min	70.36	6.99I	30.242		0.48	10.197	83.25I		4.43	10.162	82.34%		5.62	5.512	96.221		34.33	8.772	33. 33	
zinc scavenger flotation time 1.5min+	39.78	3.062	41.302		0.20.	2.572	85. 94 Z		2.66	2.551	84.702		2.10	0.90Z	97.121		30.17	4.121	37.47	
final flotation tails	590.80	50.701	100.00I		0.08	14, 151	100,001		0.02	15. loz	100.00I		0.35	2.891	100.001		29.20	40.532	100.00	

10-104 Collectorless flotation then zinc flotation in air.

Sample reference	Нq	activator	Pre-con	ditioning		ector	flotation
June 1 Ci Ci Cii Ci		added g/t	gas	tine	type	add:tion g/t	g a s
collectorless rougher - 2 minutes	B. 0	-	alr	10 min	-	-	air
collectorless scavenger - 2 minutes	8.0	-	-	-	-	-	alr
zinc rougher - 30 second	10.0	1200	MOAL	Snix	SIPI	50	air
zinc rougher - 1 ainute	10.0	-	-	-	-	-	air
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	a1 <i>r</i>
zinc rougher - 2 minutes B	10.0	-	-	-	-	-	alr
zinc scavenger - 30 seconds	10.0	-	-	-	SIPX	30	air
zinc scavenger - 1 minute	10.0	-	•	-	-	•	air
zinc scavenger - 2 minutes A	10.0	-	-	-	_	-	air
zinc scavenger - 2 minutes B	10.0	-	-	-	-	•	air

GRINDING CONDITIONS:

30 minutes. Stainless steel charge

1200 g/t zinc sulphate

10-104 Collectorless flotation then zinc flotation in air.



	SOLIBS II	MSS BALANCE		COPPER N	IASS DA M	CE 33	t	EAD MASS	MLANCE		ı	INC MASS I	MLMICE		1	RON MASS I	ML MICE	
	sample	eass pull	cop	per	CO	pe r	les	4	lea	ıđ	210	c	218	c	170	•	ire	IA
samle reference	wezght	sample total	455 a y	units	reco stage	total	assay	units	recon	very total	8558 y	enits	recev stage	ery total	85587	wits	recev	ery total
					3/mg/c				areds				20040					
																•		
triff compher time 1.5 proutes	67.06	7.312 7.312	2.56	0.187	54.672	54.432	24.30	1.774	51.281	St. 281	7.11	0.520	5.742	5.741	24.20	1.769	7.002	7.001
bulk stavenger - time 1.5 minutes	44.70	4.05% 14.16%	1.48	0.047	13.592	48.227	7.60	0.457	18. 98Z	70.262	10.80	0.740	8.172	13.922	31.00	2.170	8.772	15.012
pyrite rampher - time 30 seconds	44.52	4.712 18.071	0.25	0.012	3.441	71.661	3.26	0.154	4.442	74.70 z	6.66	0.314	3.472	17.302	38.10	1.833	7.341	23.152
pyrite rougher - time I aimite	29.77	3.152 27.022	0.32	9.019	2.942	74.692	3.85	0.121	3.501	78.201	8.46	0.267	2.751	20.331	34.50	1.150	4.612	27. <i>1</i> 52
pyrite raugher - tipe 2 minutes (A)	46.07	4.762 26.782	0.20	0.014	4.06Z	70, 66Z	3.31	0.164	4.742	82.94Z	10.00	0.534	5.921	24.751	33.70	1.473	-	34.451
pyrite raugher - time 2 minutes (B)	13.33	1.411 28.401	0.23	0.003	• •51	70.617	2.63	0.037	1.07%	84.021	12.20	●.172	1,902	28. 15%	.70.10	9.425	1.70Z	34. 15I
pyrite scavenger - time 1.5 minutes (C)	58 14	4.15% 34.55%	0.08	0.005	1.442	81.042	1.02	9.063	1.012	05.03E	3.66	0.238	2.671	30.782	42.68	2.585	10.351	44.502
pyrite scavenger - time 1.5 minutes (D)	85.69	9 011 43.562	0.08	0.007	2.101	83. ISX	1.11	0.100	2.892	89.712	3.38	0.304	3.341	34.147	42.40	3.837	15.371	41.872
pyrite scavenger - time 1.5 minutes (E)	52.25	5.531 49.091	0.07	0.005	1.45Z	84,401	1.02	0.054	1.631	70.342	4.12	0.220	2.521	34.462	41.50	2.295	7.172	71.66Z
pyrite scarenger - time f.5 minutes (F)	42.24	4.471 53.561	4.08	0.004	1.041	85.64%	1.00	9.048	1.392	71.743	5.54	0.248	2.74Z	39. 40Z	37.40	1.770	7. 0 ~1	70.15%
pyrite scavenger - time 1.5 minutes (G)	24.75	2.621 56.181	0.07	0.007	0.672	86.332	1.13	0.030	0.052	*2.591	8.56	0.224	2.461	41.002	34.90	0.714	3.441	81.922
zinc rougher - time 30 seconds	15 99	1.692 57.072	0.20	0.003	0 992	87, 321	6.75	0.013	0.372	92.961	50.00	0.846	9.35I	51.232	9.47	9.160	9.642	12.41
zinc reigher - time & ainute	18.63	1.97% 59.84%	0.22	0.004	1.271	88.591	0.78	0.015	0.442	73.40I	48.90	0.764	10.662	41.001	9.71	0.171	0.772	83.221
zinc rnigher - time 2 minutes	31.75	3.361 63.202	9.17	0.906	1.862	90.45I	0.75	0.025	●.732	74.132	44.90	1.574	17.421	79.30I	10.26	0.343	1.371	84. 6 01
zinc scavenger - time Eminute	19.67	2.081 65.282	0.24	0.005	1.462	91.91Z	0.95	0.020	0.571	94.701	39.60	9.824	9.111	80.412	13.30	0.277	1.112	
zinc reavenger - time 2 minutes	18.76	2.011 67.2°Z	0.24	0.005	1.412	93.321	1.16	●.♦23	0.672	95.371	31.90	0.440	7.072	95.48 1	14.00	0.321	1.291	84. 99Z
final fails	307.03	32.711 100.001	0.07	0.023	4.461	100.00I	0.47	0.160	4.632	100.001	1.25	0.407	4.521	100.001	7.73	3.246	13.011	100.002
test ford foromstatuted)	944,77			0.343	100.001			3.444	100.001			7.050	100.001			24.167	100.001	
TEST SIMMARY:	Sample	mass pull		copper	reces	ref Y		lead	rece	v e r y		zinc	recev	ery		zinc	FPCO	rer y
	weight	sample total		assay Z	stage	total		assay 1	stage	total		assay l	stage	total		4954y I	stage	tetal
bull flotation	133.76	14.167 14.167		1.65	68 221	68.272		17.19	70.261	70.261		0.87	13.922	13.021		27.00	15.012	
pyrite rougher flotation in air	134.51	14.242 28.402		0.27 -	11.371	79.612		3.35	13.752	84.02%		7.05	14.242	20. 157		35.46	20.351	34.157
pyrite scavesger flotation in mitrogen	262.47	27.79% 56.18%		0.00	6.732	84.331		1.07	0.501	92.592		4.47	13.721	41.001		41.04	45.442	
ZIAC rougher fintation	46.37	7.027 63.702		0.20	4.122	90.45Z		0.76	1.542	74.132		40.21	37.422			7.87	2.70Z	
zinc stavenger flotation	30.63	4.091 47.292		0.24	7.667	73.322		1.05	1.24Z			35.02	14.107			14.63	5.345	
final tails	309.03	32.712 100.002		0.07	6.601	100.00Z		0.47	4.432	100.001		1.25	4.527	100.00I		7.73	13.011	100.00

10-201 Bulk copper, lead flotation in air, pyrite flotation in air and nitrogen, zinc flotation in air.

Sample reference	pH	activator added	Pre-condi	tioning		ector	flotation
		g/t	gas	time	type	addition g/t	gas
bull rougher - 1.5 minute A	8.5	-	air	10 min	SIPX	50	٠
bulk rougher - 1.5 minute B	8.5	-	-	-	SIFX	20 20	air air
pyrite rougher - 30 seconds	9.5	-	•	•		-	air
pyrite raugher - 1 minute	0.5	-	•	-	-	-	315
pyrite rougher - 2 minutes A	8.5	-	-	-	_	-	air
pyrite rougher - 2 minutes B	8.5	-	-	-	-	-	31 <i>f</i>
pyrite scavenger - 1.5 minutes E	8.5	-	nitrogen	20 min	-	_	
pyrite scavenger - 1.5 minutes D	9.5	-	-	-	-	-	nitrogen nitrogen
pyrite scavenger - 1.5 minutes E	8.5	•	•	-	•	-	•
pyrite scavenger - 1.5 minutes F	9.5	•	_	_	_		nitrogen
pyrite scavenger - 1.5 minutes 6	8.5	-	-	-	-	-	nitragen nitragen
zinc rougher - 30 second	10.0	1290	none	5 s in	SIPX	50	31.6
zinc rougher - 1 minute	10.0	-	_	-		JV -	air air
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	alf
ZINE scavenger - 1 minute	10.0	-	-	-	SIPX	30	air
	19.0				-	94	211

GRINDING CONDITIONS:

30 minutes. mild steel charge

1200 g/t zinc sulphate

10-201 Eulk copper, lead flotation in air, pyrite flotation in air and nitrogen, zinc flotation in air.



	SBL 105 H	ASS BALAN	CE		COPPER NA	SS BALANC	E	LEAD MASS	DALANCE		t	INC MASS I	MLANCE		I	RON HASS I	ML MICE	
	sample	#a55 p	ol 1	Copp	er	cabb	er	lead	lea	Id	210	C	216	ĸ	ire		ire	
	werght	Sample	total	2553Y	units	FECEV	ery 455	ay units	rece	er y	assay	mats	79001	er y	85547	units	reco	ver v
sample reference						stage	total		stage	total	•		stage	tetal				total
bull rougher - time 1.5 minutes	91.76	7.591	9.59Z	2.02	0.194	57.531	57.532 20.2	0 1.934	59.401	59.601	8.16	0.792	9.832	8. 83Z	26.50	2.540	10.167	10.107
bulk snavenger - time 1.5 minutes	59.90	6.20Z	15.072	0.58	●.034	10. 0 21	68.361 7.4	7 0.467	14.442	74.04Z	10.60	9.446	7.523	14.352	33.20	2.005	0.362	18.531
pyrite rougher – time 30 seconds	56.17	5.90Z	21.772	0.20	0.012	3.512	71.062 2.5	4 0.150	4.612	78.65Z	5.00	0.347	3.921	20.27%	39.70	2.342	1.302	27.92
pyrite rougher - time 1 minute	26.49	2.782	24.552	0.78	0.008	2.312	74.102 3.4	4 0.0%	2.751	61.40Z	6. 80	0.247	2.791	23.06Z	35.70	0.773	3.901	31.90
pyrate rougher - time 2 mantes (A)	22.70	2.382	26.932	0.29	0.007		74.232 3.3		7.422	84.02Z	11.40	●.272	3.071	26.132	31.90	0.761	3.051	34. 751
pyrite rougher – time 2 minutes (B)	16. 29	1.711	20.642	●.27	9.005	1.377	77.602 3.7		1 177	86.022	12.40	0.212	2.392	28.521	27.50	0.504	2.021	36.971
pyrite scavenger - time 1.5 minutes (E)	3.65	0.302	29.032	0.32	0.061	0. 34I	77.971 2.9		0.351	86.371	11.80	0.045	0.51Z	27.932	26.60	0.102	0.41%	37.37
pyrite scavenger - time 1,5 minutes (B)	0.00	0 002	27.032	0.00	9.000	0.00I			0.00Z	86.372	0.00	9.000	0.00Z	27.031	0.00	0.000		37.37
pyrite scavenger - time 1.5 minutes (E)	0.00	0.001	29.031	0.00	0.000	0.90Z			0.00Z	96.371	9.00	0.000	0. 0 01	27.031	0.00	9.000	0.091	37.37
pyrate scavenger - time 1.5 minutes (F)	9.00	0.001	27.032	4.99	8.009	9.98L			0.001	86.371	0.00	0.000	0.001	27.031	0,00	1.000	0.002	37.37
pyrite scavenger - time 1.5 minutes (6)	0.00	0.001	29.03I	0.00	9.000	0.002	77.971 0.0	0,000	0.001	66.371	1.00	1.100	0.001	27.03%	1.00	0.900	0.001	37.37
zenr enigher - tage 30 seconds	10.87		31.012	0.47	●.●10	7.89I			●.711	87.0EZ		0.721	10.391	39.432	17.10	0.339		30.73
zincemigher - time 1 minute	22.76		33.402	0.20	0.005	1.171			0.762	87.84Z		1.100	12.42%	31. 64 Z	12.70	0.300	1.242	• •
zini emigher - time 2 ainutes	30. 47		37.441	9.25	0.010	3.001			1.371	87.23Z		1.405	17.032	70.071	15.10	1.610	2. 44Z	
zuer emigher - time 2 minutes	23.84	2.501	39.951	●.17	9.004	1.767	96.541 [.]	4 0.029	0.802	70.11Z	34.40	0.712	10.292	01.16Z	10.10	0.453	1.02%	44.23
zanc sravenger - time 1 minute	9.95		40.772	0.18	0.002				0.352	70.461		0.3.	4.181		19.20	0.201		45.03
zim sraven ger - time 2 minutes	15.44	1.621	42.621	0.70	●.●03	0.741	00.971 1.0	6 0.017	♦.53 2	90.992	30.70	0.501	5.442	70.771	21.50	0.349	1.40%	46.43
final tails	546.30	57.301	100.001	0.07	0.040	11.932	100.901 0.5	il 0.273	7.01I	100.001	1.39	0.798	7.012	100.002	23.30	13.371	53.571	100.00
test lend freemstituted)	952.00				0.337	100.001		3.249	100.061			8.857	100.001		35.70	24.960	100.001	
BEST SUMMARY:	sample.	0055	•		copper	19001	- •	lead	recev	ery		zinc	recev	ery		iron	recov	ery
	weight	sample	total		assay I	stage	total	8554y Z	stage	total		assay I	stage	total		assay I	st age	total
bulk flatations	151.06		15.072		1.45	49.361	_	15.16	74.042			7.13	16.351	14.352		27.15	10.532	
pyrite rougher flotation in air	121.65		20.611		0.24	9.251		3. 65	11.767			8.44	12.17%	28.521		36.01	10.43I	
pyrite scavenger flotation in mitrogen	3.45		27.032		0.32	0.361		2.95	0.351			11.00	0.512			26.60	0,412	
zinc rougher flotation	103.96		39.951		0.26		86.541	1.11	3.741			42.28	52.132	01.147		15.47	1.01	44.2
zini scavenger flotation	25.39		1 42.627		0.19	1.521		1.00	0.001	70. 4 72		32.66	7.841			20.64	2.201	-
final tails	544.30	57.30	Z 100.00Z		●.07	11.731	100.00Z	●.51	9.012	100.00Z		1.37	7.012	100.00Z		23.30	53.578	100.0

10-202 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation in air.

The state of the s

FLOTATION CONDITIONS pH activator Pre-conditioning collector flotation sample reference added type addition gas g/t tine g/t bulk rougher - 1.5 ainute A 8.5 311 10 min SIPI 317 bulk rougher - 1.5 sinute B SIPX 315 pyrite rougher - 30 seconds 8.5 **a**11 pyrite rougher - ! sinute 8.5 315 pyrite rougher - 2 minutes A 8.5 217 pyrite rougher - 2 minutes B 8.5 215 pyrite scavenger - 1.5 minutes D **B.5** 415 zinc rougher - 30 second 19.9 1200 none SIPX 415 zinc rougher - I minute 10.0 116 zinc rougher - 2 minutes A 10.0 415 zinc rougher - 2 minutes B 10.0 air zinc scavenger - 1 minute 10.0 SIPY 415 Zinc scavenger - 2 minutes 10.0 317 GRINDING CONDITIONS: 30 minutes, mild steel charge 1200 g/t zinc sulphate

10-202 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation in air.

	50L105 F	MASS PALANCE		C o ffer I	MSS BALM	ICE .		EAO MASS	BALANCE		ı	INC MASS	BALANCE		1	IRON MASS	MLMICE	
	Sample	mass pull	CO	per	cog	per	le	nd .	le	ad	218	C	211	N C	170	36	ire	•
Samule reference	≈ 21 ght	sample total	assay	units		very	assay	units	reco	•	8558 y	units	FECO		45587	mits	reco	-
					stage	totai			stage	total			stage	total			stage	total
bulk rougher—time 1.5 minutes	73.70	7.272 7.272	2 65	0.173	55. 452	55. 452	22.80	1.459	47.641	47.642	6.4*	0.472	5.721	5.271	25.94	1.004	7.401	7.991
bulk scavenger—time 1.5 minutes	*1.19	8.9°Z 16.26Z		0.051	14.742	70.192	8.39	0.754	21.67%	6°.312	7.57	9.840	7.522	14.75%	35.40	3.201	12.701	29. 182
pyrite raugher - time 30 seconds	23 42	7.241 23.501	0.20			14 250												
syrite compher time 1 aimite	49 40	4 84Z 30.35Z		0.014	4.161	74.352	2.78	0.201	5.701	75.092	5.34	6.387	4.202	17.032	44.60	2.095	11.44	31.672
pyrite rougher time 2 minutes (A)	35 75	3.522 33.872	0.20	0.014	3.94Z	78 29%	2.48	0.183	5.272	20.352	0.11	0.418	4.631	23.452	38.70	2.440	10.511	42, 10%
pyrite ringher fine 7 ainutes (B)				0 (10)	2.512	80.822	3.04	●.107	3.081	83.432	8.91	0.314	3.482	27.131	35.30	1.244	4. 94 Z	47.121
blivir combine tac t attidf62 (8)	11.84	1.172 35.04Z	9 25	0.003	0.847	81.667	3.07	0.036	1.032	84.461	11.60	0.138	1.52%	28.451	30.00	0.341	1.432	40.552
pyrafe scavenger - fame 0.5 minutes (C)	40 93	4.942 39 012	n, (;0	0 net	1 642	22 712	1.29	0.052	1.501	85.762	3.69	0.149	1.651	30, 301	42.10	1.699	4.742	55, 301
Pyrite sia eugur - time 1 minute (B)	10 .3°	8.712 47.792	0.09	0.000	7 757	84 *47	1.12	0.078	7.801	86.762	4.17	0.365	4.042	34.342	41.40	3.425	14.393	47.491
pyrite scavenger - time 2 minutes (E)	55.55	5.482 53.262	9.09	0.005	1.421		1.17				-							
pyrite scavenger time 2 minutes (F)	20.32	2.001 55 271	0.10	0.002	9 581			0.064	1.041		6.24	0.342	3.782	36.122	38.50	2.109	0.372	79.051
Pyrite scavenger - time 2 minutes (6)	15.62	1.54% 56.81%	0.10	0.002	0 402	87.44Z	1.37	0.027	0.792	91.391	7.03	0. 181	2.001	40.132	34.30	0.687	2.732	80.78Z
pyrite scavenger—time 2 minutes (N)	19.22	1.8°% 58.70%		0.002	0.551	87.992	1.43	0.022 0.032	0.63Z 0.91Z	*7.021 *2.931	11.10 11.30	0.171 0.214	1. 07 1 2.371	42.021 44.392	31.30 30.50	0.482 0.578	1.911 2.291	82.472 84.772
ZINC rougher time 10 seconds	20.85	2.062 40.762				•• •••												
ZIAC Emiglier tion Laguite	31.67	3 122 AT ERY	0 18	0.004	1.062	89.051	0.54	0.011	0.321	93.252	55.60	1.143	12.45%	57.042	7.03	0.161	0.442	85.432
ZIRC raugher tree 2 grantes	31.07 21.96		0 10	0.006	1.712	90.767	0.43	0.070	0.571	73 812	\$2.10	1.627	10.001	75.047	9.17	0.286	1.147	96.74Z
ZINC rougher time 7 grantes	16.97	2.172 66.052	0.22	0.045	1. 172		0.82	0.018	0.517	94.321	45.00	0.902	10. 72	86.021	11.20	9.242	0.762	
The strate of th	16.77	1.671 67.721	0.24	0.001	1.167	93.292	1.00	4.418	0.52%	94.84%	35.60	9.596	4.592	92.512	14.40	0.241	0.762	10. MT
zinc sca enger - time 2 minutes A	18 55	1.832 49.552	0 27	0.005	1 422	94.712	1.41	0.024	0.74Z	95.58%	10.40	0.355	3.972	94.542	20.30	0.371	1.472	90.162
ZINC Sta Pright time 2 minutes B	17.91	1.77% 71.31%		0.004	1.17%		1.56	0.028	0.791	74.371	7.81	0.138	1.532	78. 04Z	17.60	0.344	1.371	
final tails	290.96	28.671 190.001	0.05	0.014	4.132	100.001	0.44	0.126	3.632	100.001	0.61	0.175	1.941	100.001	7.44	2.134	0.471	100.002
test feed treconstituted)	1014.28			0.34	100.002			3.481	100.002			7.035	100.001			25.175	100.001	
IEST SUMMEY:	Sample	asss sell		Copper	recov			lead	recev			ZINC	recov			1700	recev	
	wezght	sample total		assay 1	stage	total		assay 1	stage	total		assay I	stage	total		assay I	stage	total
bulk finistron	164.97	14 347 44 349			** ***	** **=			40 5									
pyrite rimidier flotation in air		16.262 16.262		1.50	70.192	70.191		14.83	49.312	67. 31Z		8.17	14.752	14.752		31.26	20. 102	20.181
Pyrite cra pager flutation to mitrogen	199.41	10.772 35.042		0.21	11.472	#1 66Z		2.01	15. 152	84.462		6.47	13.912	20.652		38.00	20.371	48.55%
histra are mides attribution to uttrodes	240.03	23.67% 58.70%		0.0	4.332	87. 991		1 75	9 477	až ažš		6.Ct	15 742	44 325		38.79	36.432	
zinc sta enger flotation	74.48	9 072 67,772		0.20	2 291	93.29%		0.44	1.912			51.22	40.222	92.612		9.39	3.472	
final tuis	76.46	3.592 71 312		0.25	2.592			1.40	1.531	76.37%		13.71	5. 45z	78.06Z		17.76	2.052	91.532
Three 1 1115	290.96	20.491 160.001		0.05	4.132	100.00Z		0,44	3.432	100.00Z		0.41	1.742	100.00Z		7.44	0.472	100.00Z

10-203 Bulk copper, lead flotation in air, pyrite flotation in air then nitrogen and zinc flotation in air.

	pН	activator	Pre-cond:	tioning	coll	ector	flotation					
sample reference		added			type	add1t1on	qas					
		g/t	gas	time		g/t	•					
bulk rougher - 1.5 minute A	8.5	-	aır	10 a ın	SIPI	50	air					
bulk rougher - 1.5 minute B	8.5	-	-	-	SIPY	30	air					
pyrite rougher - 30 seconds	8.5	-	-	-	-	-	air					
pyrite rougher - 1 minute	8.5	-	-	-	-	-	air					
pyrite rougher - 2 minutes A	9.5	-	-	-	-	-	alf					
pyrite rougher - 2 einutes B	8.5	-	-	-	-	-	air					
pyrite scavenger - 30 seconds C	8.5	-	nitrogen	20 min	SIPI	10	nitrogen					
pyrite scavenger - 1 minute D	8.5	-	-	-	-	-	nitrogen					
pyrite scavenger - 2 minutes E	8.5	-	-	•	-	•	nitrogen					
pyrite scavenger - 2 minutes F	8.5	-	-	-	-	-	nitrogen					
pyrite scavenger - 2 minutes 5	8.5	-	-	-	-	-	nitrogen					
pyrite scavenger - 2 minutes H	8.5	-	•	-	•	•	nitrogen					
zinc rougher - 30 second	10.0	1200	none	5 min	SIPX	50	air					
zinc rougher - 1 minute	10.0	-	-	-	-	•	air					
zinc rougher - 2 minutes A	10.0	•	•	-	-	-	arr					
zinc rougher - 2-minutes B	10.0	•	-	-	-	•	air					
zinc scavenger - 1 minute	10.0	-	-	-	SIPI	30	aır					
zinc scavenger - 2 minutes	10.0	-	-	-	-	-	air					
	GRINDING CONDITIONS:											
	30 minutes. mild steel charge											
	1200 g/t zinc sulphate											

10-203 Bulk copper, lead flotation in air, pyrite flotation in air then nitrogen and zinc flotation in air.



	SOLIDS N	ASS BALANCE		COPPER M	ASS BALAM	Œ	(LEAD MASS	MLANCE		1	INC MASS	DALANCE		ı	ROM MASS (MLMICE	
	520017	eass pull	cop	per	copy	per	lea	ıd	lea	ıd	210	C	216	ĸ	170		100	
	wezght	sample total	a 55 a y	unsts	Fec b	4 27 y	9229À	wats	F 2C 01	very	assay	units	F 2C 0	rer y	45549	mats,	1900	very
sample reference					stage	tetal			stage	total			stage	total		,	stage	total
bull remaker - time 2 assetes	56.21	5.531 5.5	I 2.88	9.159	47.282	47.20Z	5.72	0.289	8.372	8.372	4.13	0.339	4.102	4.102	32.99	1.820	4.442	4, 441
bulk scavenger - time 2 ainutes	132.35	13.032 18.5		0.087	26.992	76.272		1.772	51.351	59.731	7.05	0.718	11.312	15.472	31.00	4.037	14.29%	20.731
pyrite rangher - time 1 minute (A)	32.30	3.10% 21.7		0.006	1.072	78.142		0.245	7.101	66. 82 1	7.20	0.293	3.402	17.072	31.90	1.014	3.571	24.32
pyrite raugher - time 1 minute (B)	29.74	2.732 24.6		0.004	1.271	79.412		0.120	3. 48 Z	70.30I	7.25	0.212	2.612	21.712	22.90	1.051	3.7 71	28.04
pyrite rougher - time 1 minute (C)	23.56	2.321 26.9		0.003	0.791	80.201		0.054	1.417	71.917	4.44	0.154	1.90Z	23.40Z	34.30	0.842	2.962	31.01
pyrite enigher - time & minute (8)	21.05	2.071 29.0	12 0.09	9.002	0.581	80.77 1	1.78	0.037	1.071	72.981	4.55	0.134	1.672	25.271	22.00	0.725	2.571	33.50
pyrite scavenger - time 1 minute (A)	231.50	22.002 51.0	5Z 0.06	0.014	4.231	85.001	1.54	0.351	10.18Z	83 16Z	2.80	0.638	7.842	33.132	42.00	9.756	34.521	40. 10
pyrite scavenger - time minute (B)	83.93	8.172 60.0		0.006	1.771	86.77%		0.141	4.072	87.231	4.82	0.374	4.852	37.991	40.30	3.335	11.001	77.90
pyrite scavenger - time 1 minute (C)	38.06	3.751 63.7	71 0.19	●.004	1.167	87.93L	1.90	0.071	2.041	8 9.301	8.54	●.321	3.952	41.742	35.70	1.345	4.762	84.63
zine e nigher - time 30 seconds	62.42	4.19Z 69.9	Z 0.18	0.011	3.472	91.352	1.12	0.067	1.992	91.202	46.10	2.832	34.001	76.821	11.70	0.731	2.591	87.2
zinc edigher - time I minute	33.64	3.312 73.2	32 0.23	0.008	2. 36I	93.712	1.50	0.452	1.522	92.811	41. j0	1.361	14.742	93.5 8 Z	17.60	0.583	2.062	
zinc rimpher - time 2 minutes	26.41	2.62% 75.8		_	1.862			0.050	1.467	94.26Z	13.20	0.346	4.762	97.84Z	23.10	0.605	7.14 1	
zinc rmigher - time 2 minutes (B)	11,20	1.101 74.9	51 0.22	0.002	0.751	96.321	2.14	0.024	0.68 2	94.951	4.42	0.049	0.601	98.442	23.70	0.261	0.972	92.37
zinc scavenger - time 30 seconds	12.48	1.237 70.1	BZ 0.16	0.002	0.411	94.932	1.62	0.020	0.58I	95.53I	1.75	0.024	0.302	70.742	27.00	0.332	1.173	73.5
zinc =ravenger = time minute	9.39	0.921 79.1		0.002	0.491	97.412	1.78	0.016	0.4BZ	76.00%	1.16	0.011	0.131	98.871	23.40	0.216	0.772	
final tails	212.30	29.901 100.0	01 0. 04	9.006	2.591	100.002	0.55	0.138	4.002	100.002	0.44	0.092	1.131	100.002	7.70	1.407	5.692	100.00
test feed (reconstituted)	1015.92			0.323	100.001			3.450	100.007			8.117	100.001			29.264	100.001	
IESI SIMMARY:	sample	mass pull		Copper	reco	er y		lead	FECOV	er y		zinc	recov	ery		1100	recev	ery
	weight	sample tota	1	assay I	stage	total		assay I	stage	total		assay I	stage	total		assay I	stage	total
bulk flotation	188.56	10.567 18.5	49	1.33	74 974	76 271		11.10	59.731	59.732		6.78	15 407	15.492		31.57	20.731	20.73
pyrite rougher flotation in air	106.65	10.562 18.3 10.502 29.0		0.14	4.501			4.36	13.262	72.98%		7.57	7.782	25. 271		34.40	12.05%	
syrite scaveneer flotation in mitrogen	352.47	39.712 43.7		0.07	7.162			1.62	16.312	89.301		3.70	16.662	41.742		41.50	51.071	
zinc rimeher flotation	76.06	9.462 73.7		0.20		93.712		1.28	3.512	92.812		44.35	51.452			13.90	4.452	
zinc scavenger flotation	57.40	5.871 79.1		0.20	3.712			1.00	3. 201	76.00I		10.96	5. 291					
final tails	212.30	20.901 100.0		0.29 0.04	2.592	7/. 412	•	9.44	3. 201 4. 0 01			10.96 0.44	J. 274	78.0/4		24.0 0 7.7 0	3.001 5.471	94.3 100.0

10-204 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.

	рH	activator	Pre-cond:	tioning	coll	ector	flotation
sample reference		added			type	addition	gas
		g/t	gas	time		g/t	•
bull rougher - time 2 minutes	8.5	•	air	10 min	SIPX	50	air
bulk scavenger - time 2 minutes	8.5	-	-	-	SIPX	30	air
pyrite rougher - time 1 minute (A)	8.5	-	nitrogen	4 a ın	-	-	nıtrogen
pyrite rougher - time 1 minute (B)	8.5	-	nitrogen	4 min	-	•	nitrogen
pyrite rougher - time 1 minute (C)	8.5	-	nitrogen	4 min	-	-	nitragen
pyrite rougher - time 1 minute (D)	8.5	-	nıtragen	4 410	-	-	nitrogen
pyrite scavenger - time 1 minute (A)	8.5	-	-	-	SIPX	10	nitrogen
pyrite scavenger - time 1 minute (B)	8.5	-	-	-	-	••	nitragen
pyrite scavenger - time ! minute (C)	8.5	-	-	-	-	-	nıtrogen
zinc rougher - time 30 seconds	10.0	1200	none	5 a ın	SIPX	50	air
zinc rougher - time 1 minute	10.0	-	-	-	-	-	air
zinc rougher - time 2 minutes	10.0	-	-	-	-	-	alf
zinc rougher:- time 2 minutes (B)	10.0	₹.	-		-	-	air
zinc scavenger - time 30 seconds	10.0	-	-	-	SIPX	30	air
zinc scavenger - time 1 minute	10.0	•	-	-	-	-	air

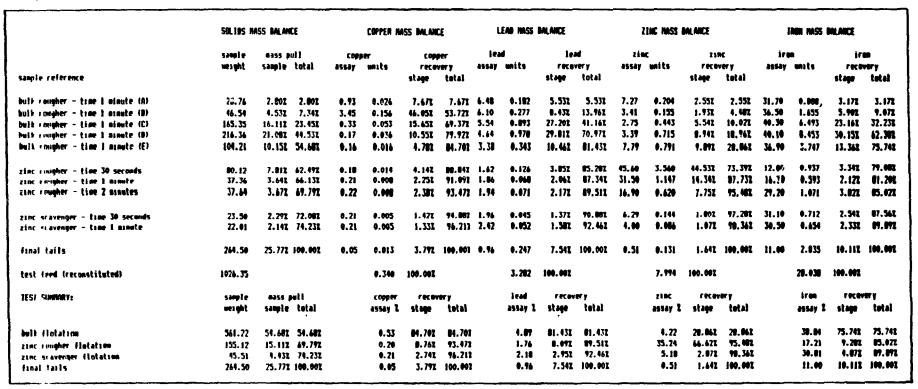
GRINDING CONDITIONS:

50 minutes, stainless steel charge.

Zinc Sulphate, 1200 g/t

10-204 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.

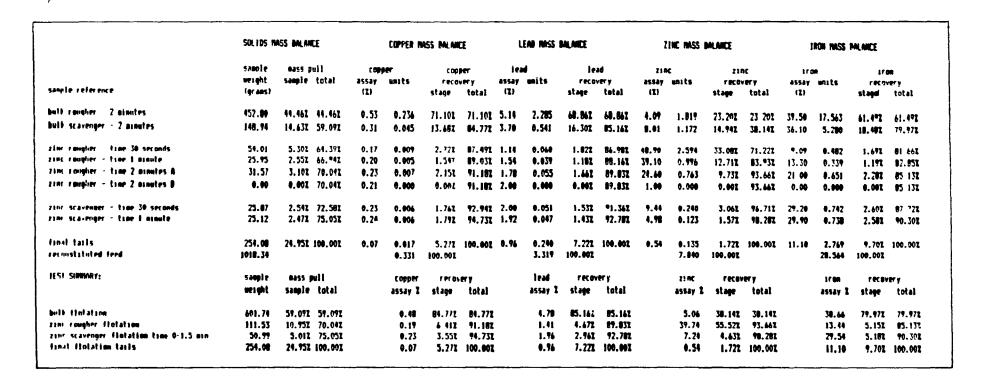




10-205 Bulk copper, lead, pyrite flotation in nitrogen with stage addition of collector, then zinc flotation in nitrogen.

	FL	r T A	TION	C	0 N D	1 7 1	0 N S				
Sample reference	pH	activator add ed	Pre-condi	tioning	call type	ector addition	flotation gas				
Jampie i erei enec		g/t	gas	tiae	-,,	g/t	,				
bulk rougher - 1 minute A	8.5	-	nıtragen	3 a in	SIPI	20	nitrogen				
bulk rougher - 1 sinute B	9.5	-	nitrogen	3 m in	SIPX	20	nitrogen				
bulk rougher - 1 minute C	8.5	-	nitragen	3 min	SIPI	20	nitrogen				
bulk rougher - 1 sinute D	8.5	-	nitrogen	3 min	SIPY	20	nıtrogen				
bulk rougher - 1 minute E	8.5	-	nitrogen	3 810	SIPX	20	nitrogen				
zinc rougher - 30 seconds	10.0	1200	none	5 min	SIPX	50	air				
zinc rougher - I minute	10.0	-	-	-	-	-	31 0				
zinc rougher - 2 minutes	10.0	•	-	-	-	-	a 17				
zinc scavenger - 30 seconds	10.0	-	-	-	SIPX	30	air				
zinc scavenger - 1 minute	10.0	•	-	•	-	-	317				
	GRINDING CONDITIONS:										
			30 s inu	tes. mil	d steel	charge					
			1200 g/	t zinc s	ulphate						

10-205 Bulk copper,lead,pyrite flotation in nitrogen with stage addition of collector, then zinc flotation in nitrogen.



10-206 Bulk Cu,Pb,pyrite flotation in nitrogen, single addition of collector, zinc flotation in nitrogen.

samnle reference	pH i	activator	Pre-con	ditioning	coll	ector	flotation					
sample reference		added g/t	gas	time	type	addition g/t	gas					
bulk rougher - 2 minutes	8.5	-	-	-	SIPX	100	nitrogen					
bulk scavenger - 2 minutes	8.5	-	-	-	-	-	nitrogen					
zinc rougher - 30 second	10.0	1200	none	5 m in	SIPX	50	air					
zinc rougher - 1 minute	10.0	-	•	-	-	-	31 <i>r</i>					
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	alr					
zinc rougher - 2 exnutes B	10.0	-	-	-	-	-	air					
zinc scavenger - 30 seconds	10.0		-	-	SIPX	36	alr					
zinc scavenger - 1 minute	10.0	-	-	-	-	-	alr					
			6RIND1	N 6 CONDITIO	NS:							
	30 minutes. mild steel charge 1200 g/t zinc sulphate											

10-206 Bulk Cu,Pb,pyrite flotation in nitrogen, single addition of collector, zinc flotation in nitrogen.

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	SOLIDS M	ASS BALAN	CE		COFFER I	IASS INLAM	£	ι	EAD MASS	BALANCE		1	INC MASS	BALANCE		1	ROW MASS	MALMICE	(
	sample	8455 P	ol l	cop	er	£ 0p;	er	lea	4	lea	i	210)C	218	c	110	M	iro	98
	⊯esgfit	sample	total	assay	units	reco	ery	assay	units	recov	er y	assay	units.	recov	er y	2553y	units	recov	very
Sample reference						stage	total			stage	total			stage	total			stage	total
zinc rougher - time 30 seronds	24 64	2.501	2.50%	0 -3	0.008	2.642	2.641	7.5C	0.198	6.342	6.34Z	4.84	0.172	2.392	2.392	31.50	0.789	2.781	2.797
zinc rougher - time I minute	77.16	2.752	4.762	0.35	0.909	2.571	5.15%	5.60	0 126	4. 761	10.60%	7.60	0.171	2.381	4.762	32.00	● 721	2.541	5.321
zinc scavenger – tiee 1 minute ([)	27.72	2.021	7.572	0.37	0.010	3.332	8.48Z	4,24	0.117	4.032	14.432	9.27	6.761	3.632	8.3°Z	29.10	0 820	2.891	9, 213
zinc scalenger - time Lainute (II)	18.39	1.072	4.447	0 40	0.007	2, 392	10.862	3,94	0.074	2,492	17.112	10.90	0.202	2 802	11.172	27.40	0.517	1.012	10.02
ZINC SCAVENGER - time minute ([[])	18.68	1.902	11.342	0.41	0.008	2.482	13 352		0.072	2,422	19.542	13.90	0.262	3.642	14.632	26.10	0.4%	1.75%	11.77
zinc scavenger - time Eminite (1V)	24 31	2.472	13.017	9.41	0 010	3.232	16.582	3.40	0.066	Z. 90Z	22.44Z	11.90	0.294	4 091	18.912	25.00	9.419	2.182	13 95
zinc scavenger - time E minute (V)	14.81	1.511	15.321	0.45	0.007	2.161	18.742	3.82	0.058	1.742	24.381	17.00	0.256	3.552	22.47%	23.30	0.351	1.241	15.18
zinc scavenger - time i minute (VI)	22.43	2,28%	17.69Z	0.45	0.010	3.271	22.012	3.52	9,080	2.712	27.091	15.30	0.340	4.847	27.312	23 99	0.545	1.921	17.10
zinc scavenger - time 1 minute (VIII)	31 67	3.721	20.822	0.46	0.015	1.721	26.741	4.14	0.133	4.502	31.57%	17.50	0.563	7.82%	35.132	26.80	0.863	3.041	20.15
•	779.00	79.182	100.00Z	0.29	0.230	73.262			2.027	68. 41Z	100.00I	5.70	4.6/2	64.872	100.001	28.60	22.696	79.05%	100.00
recnistated feed	783.81				0.313	100,002			2.763	100.001			7.207	100.00Z			28.360	100.001	

10-207 Copper activation and zinc flotation in nitrogen only.

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	pH a	activator	Pre-condi	tioning	colle	ector	flotation			
sample reference		added			type	addition	gas			
		g/t	gas	time		g/t				
zinc rough er - 30 seconds	10.0	1200	nitrogen	5 min	SIPX	50	nitrogen			
zinc rougher - 1 minute	10.0	•	-	-	•	-	nitrogen			
zinc scavenger - I minute I	10.0	-	-	-	SIFX	20	nitrogen			
zinc scavenger - 1 minute II	10.0	•	-	-	SIPI	20	nitrogen			
zinc scavenger - 1 minute III	10.0	-	-	-	SIPX	20	nıtrogen			
zinc scavenger - 1 minute IV	10.0	-	-	-	SIPY	29	nitrogen			
zinc scavenger - 1 minute V	10.0	-	-	-	SIFY	20	nitrogen			
zinc scavenger - 1 minute VI	10.0	-	•	-	SIPX	20	nitrogen			
zinc scavenger - 1 minute VII	10.0	-	-	-	SIPX	50	nitrogen			
			6RINDIN6	CONDITI	ons:					
	30 minutes, stainless steel charge									
				t zinc s						

10-207 - Copper activation and zinc flotation in nitrogen only.

	SOLIDS N	ASS BALAME	Œ		COPPER M	55 BALANC	ī	i	EAD MASS I	MLANCE		2	INC MASS I	ALANCE		10	ROW MASS 2	MLMICE	
	saeple weight	mass po sample		COPP assay		recov		lea	d units]ea		2101 25524		Z L P	-	iro assay		176	
sample reference	- tynt	30mp15	(014)	43347	- 1.1(3	stage				stage	total	,				43347	mus (2	recor stage	,
bulk rougher - 2 minutes	53.54	5.221	5.221	2.66	0.139	43.352	43.351	5.06	0.264	7.391	7.302	4.73	0.351	5.101	5.102	33.22	1.733	6.12%	6.122
bulk scavenger - 2 ainutes	130.70	12.742	17.96I	0.53	0.068	21.092	64.44Z	2.98	0.380	10.632	16.022	3.93	0.501	7.271	12.371	39.10	4.981		23.701
zinc rougher - time 30 seconds	33 22		21 191	0.37	0.012	3 74 2	68.181	7.24	0.234	4.567	24.58I	30.60	0.991	14.302	26.751	16.10	0.521	1.842	25 531
zinc rougher - time 1 minute	36.52		24. 75Z	0 34	0 012	3.781	71.962	8.98	0.320	0.951	33 531	28.40	1.011	14.691	41.432	17.70	0.437	2.251	27.781
zanc rougher - tame 2 manutes A	20 24	7.751	27 501	0.2 9	0.008	7.492	74,451	8.48	0.733	4.532	40.061	25.19	9.671	10.03I	51.462	17.90	0.548	1.731	29.721
zinc rougher - time 2 minutes B	17.98	1.751	29.261	●.27	0.005	1.481	75.931	6.78	0.117	3.331	43.392	23.30	0.408	5.931	57.392	21.10	0.370	1.301	31.022
zinc scavenger - time 30 seconds	220.41	77.761		0.14	0.031	7.732	85.671		0.815	22.017		8 11	2.001	29.061	86.45Z	36.00	8.014	28.281	50.301
zinc scavenger - time 1 minute	117.55	11 652	63.172	0.15	0.C17		91.13%	2.70	0.315	0.011	75.00%	5.09	0.593	0.617	95.061	39.40	4.591	14.202	75.511
zinc scavenger - time 2 minutes A	48.42	4 721	67.89Z	9.15	0.007	2.211	93.342		0.107	3.067	78.07%	3.20	0.151	7.197	97.26%	37.90	1.883	4.452	02 151
zinc scavenger - time 2 minutes B	21.19	2.071	69. 95 2	0.16	0.003	1.032	94.371	2.14	9.044	1.242	79.312	2.46	0.051	0.742	97.991	30.30	0,791	2.791	84.94
final tails	300.30	30.051	100.00Z	0.06	●.018		100.001	2.46	0,739		100 002	0.46	0.138		100.002	14.20	4.267	15. C&Z	100.001
reconstituted feed	1026.07				0.320	100.00I			3.572	100.001			6.007	100.00Z			29.335	100.001	
TEST SHIMARY:	sample	#355 f			copper	recov	•		lead	recov			2180	recov	•		1ren	recov	
	weight	saaple	total		assay l	stage	total		assay l	st age	total		assay l	stage	total		assay 1	stage	total
bull flotation	194.24		17.96Z		1.15	64.442			3.58	18.072			4.74		12.371		37.39		23.701
zinc rougher flotation	115.96		29. 262		0.33	11.497	75. 93%		8.02	25.371			27.44	45.021	57.391		10.37	7.332	
zinc scavenger flotation time 0 1.5 min	347.94		63.172		0.14	15.177	71.132		3.33	31.427			7.65	37.672			37.17	44.48Z	
ginc scazenger flotation time 1.5min*	67.61		69.95%		0.15	3.242	94.372		2.27	4.302			2.97	2.931			39.41	9.44I	
final flotation tails	308.30	30.052	100.004		9.06	5.631	100.00I		2.44	20.692	100.901		0.46	2.017	100.00Z		14.20	15.041	100.00

10-208 Bulk Cu,Pb, pyrite flotation in nitrogen, zinc flotation

	FL	0 T A	T I O N	C C	0 N D	I T I	O N S
	pH		Pre-cond:	tioning		ectar	flotation
sample reference		added g/t	gas	time	type	addition g/t	gas
bulk rougher - 2 minutes A	8.5	-	nitrogen	20 min	SIPX	50	ni trogen
bulk rougher - 2 minutes B	8.5	-	-	-	SIPX	20	nitrogen
zinc rougher - 30 second	10.0	1200	none	5 010	SIPX	50	nitrogen
zinc rougher - 1 minute	10.0	-	•	-	-	-	nitrogen
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	nitrogen
zinc rougher - 2 minutes B	10.0	-	-	-	-	-	nitrogen
zinc scavenger - 30 seconds	10.0	-	-	-	SIPX	30	nitrogen
zinc scavenger - 1 minute	10.0	-	-	-	-	-	nitrogen
zinc scavenger - 2 minutes A	10.0	-	-	-	-	-	nitrogen
zinc scavenger - 2 minutes B	10.0	-	-	•	-	-	nitrogen
		•	6RINDIN6	CONDITIO	ONS:		
			30 minu	tes, sta:	inless st	teel charge	
			1200 g/	t zinc si	ulphate		

	50L105 M	ASS BALAN	CE .		COPPER IN	165 BALAM	Œ	L	EAD MASS	BAL MICE		1	INC MASS E	AL ANCE		i	PC4 MASS 8	MANCE	
	5amo] e	Bass p		Coc	ser	COD	ger .	lea	-	iea	-	218		218	E	170		111	3 0
sample reference	es i âp s	sa op le	total		units	reco			wasts	rece	•	8558Y	mm1 t s	recen		assay	mits	recev	
sechit i asti fură				(2)		stage	total	(1)		stage	tetal	(3)		st age	total	(2)		st age	total
bull rougher - 2 groutes	386 38	39.722	39 731	0 60	0.238	71.132	71 132	5.24	2.082	47 8°Z	67.891	4.52	1.7%	24.382	24.381	40.10	15 933	54.072	54 PFI
bulk scavenger - 2 minutes	41.90	4.312	44.04Z	9 47	9.030	8.872	8 0.001	5 82	0.251	9.182	76.071	5.01	9.345	4.681	29 G&I	35.30	1.521	5.242	60.132
zinc rougher - time 30 seconds	62 50	6 442	50 481	0.23	0.015	4 472	84.421	2.06	0.133	4.321	90.391	45.20	2.90*	39.491	68.561	12 70	0.017	2 821	62 042
zinc rougher - time 1 minute	33.45	3.442	57. 972	0 25	9.007	2.572	86 -91	2.44	9.004	2.741	93.13%	35.20	1 211	14.442	84.991	16.90	0.550	1.901	44 847
zinc rougher - time 2 minutes A	24.92	2 561	54.481	0 24	9.006	1.842	88.821	2 40	0.062	2.011	85 13E	21.10	P.541	7.341	92.331	25.60	0.454	2.261	67 102
zinc rougher - time 2 minutes 8	14.75	1.521	58. 00 t	0.22	0.003	1.001	89.81%	1.76	0.027	0.872	86.00I	12.50	0.190	2.571	94.912	30.40	0.464	1.401	48.70%
zimc scavenger - time 30 seconds	10.80	7 051	60 041	0.17	6 003	1.042	90.851	1.74	0.040	1.291	07.30Z	4.60	0 094	1 281	%. 19I	37 20	0.761	2.522	71 221
zinc scavenger - time I minute	Z 4. 14	3. 90 %	63.94Z	9.16	0.005	1.432	72.201	1.90	0.05 7	1.841	8 7. 151	2.97	0.087	1.212	•7.392	37.30	1.110	3.051	75.178
final tasis	359.43	34.961	100.001	0.07	0 026	7.721	100.001	0.70	♦.333	10 851	100.00I	●.52	0.102	2.612	100.00I	19.50	7.207	24.032	100 001
rermstituted feed	•72.45				•. 335	100.001			3.667	100.001			7.367	100.00I			29.028	100.00I	
TEST SUPPARY:	sample	0159 #	wi i		COPPET	recev	ery		lead	recov	ery		2100	FPCOV	ery		Lr on	FECOV	er y
	ne i âșt	sample	total		assay I	stage	total		assay l	stage	total		assay I	stage	total		assay I	stage	total
bulk fletation	470.20	44.041	44.04Z		9.41	80.00I	80.001		5.30	76 CTL			4.86	29.061	29.041		39.63	60.132	871 00
zinc raugher flotation	135.71	13.76X	50.001		0.24	9.812	89.811		2.18	7.742	66.00%		34.76	65. 84Z	94.912		17.83	8.571	101 63
zinc scavenger flotation time 0-1.5 min	49.03	5.041	63.042		0.16	7.471	72.281		1.92	3.151	87.151		3.63	2.492			37.26	6.47%	
final flotation tails	359.43	34.761	190.00I		9.97	7.72%	100.001		0.70	10.851	100.00Z		0.52	2.617	100.001		19.50	24.831	100.001

10-209 Nitrogen conditioning, then Cu,Pb,pyrite flotation in air and zinc flotation in air.

	FL	0 T A	T 1 0	N C	0 N C	1 7 1	0 N S				
sample reference	pH	activato	r Pre-cana	ditioning	coll type	ector addition	flotation gas				
		g/t	gas	ti ne		g/t	·				
bulk rougher - 2 minutes	8.5	-	nitroge	15 sec	SIPX	50	air				
bulk scavenger - 2 minutes	8.5	-	•	-	SIFY	30	air				
zinc rougher - 30 second	10.0	1200	none	5 min	SIPX	50	alf				
zinc rougher - ! minute	10.0	٠-	-	-	-	-	a 17				
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	a 17				
zinc rougher - 2 minutes B	10.0	•	-	-	-	-	a 16				
zinc scavenger - 30 seconds	10.0	-	-	-	SIPX	30	¥17				
zinc scavenger - 1 minute	10.0	•	-	-	-	-	31 7				
			6RINDIN	6 CONDITIO	NS:						
	30 minutes. mild steel charge										
			1200 n	/t zinc si	olohat#						

10-209 Nitrogen conditioning, then Cu,Pb,pyrite flotation in air and zinc flotation in air.

	SOL 185 M	ASS PALANCE		COFFER !	MSS BALAN	Œ	1	LEAD MASS	BALANCE		1	INC MASS	BALANCE			IPON MASS	BAL NUCE	
	500019	eass pull	COS	per	coe	90 7	le	aď	le	24	•••							
	weight	sample total	4554v	units	rece			units			218		211	-	15	Des .	17	De
Sample reference	(grans)		(2)		st	,	(\$)	611173	reco stage	total	4554v (Z)	MULES	reco- stage	vers total	4558y (1)	enits	reco stage	•
bulk rougher - 2 minutes	417.70	41 022 45 021	0 57	9.245	75 762	75 761	5.24	2.306	(8 (30				•				rege	(0.41
hull-scavenger - 2 minutes	138 70	14.292 57.312	0.21	0.030	7.271			0.554	68.672 16.472	60.621 85.111	3.73 9.47	1. 675 1.353	71 492 18.112	21 491 39 601	30 90 34.50	16 73 <u>6</u> 4,929	59 061 17 391	
zinc routher - time 30 seconds	72 68	7 4°1 64 801	0 20	0 015	4.632	B*.661		0 0%	2.051	87.961	44 60	3 33•	44 701	84.301	17.70	0.913		79 67
zinc rougher - time 1 minute zinc rougher - time 2 minutes A	27.48	2 871 67 631	9.27	0.008	7 362			7.061	1.027	89.782	27.00	0.764	10.231	74.542	21 90	0.410	7 172	
	22.42	2 311 69.941	0.26	9.00	1.051	93.871		0.061	1.602	91 58I	6 77	0.203	2 712	97.251	31.60	0.730	2.581	
zinr rougher - time 2 minutes B	12.94	1.332 71.272	0.21	0.903	0.841	94.74%	2.40	0.032	9.952	72 531	3.21	0.043	0.57%		31.40	0.425		B5 *4
zinc scavenser - time 30 seconds	14 11	1 451 72 721	0.15	0.002	0 672	95.412	1.82	9 026	0.791	93 321	2.03	0.030	A 407	9 221	75 14			
zinr scavenger - time 1 minute	10.75	1 111 73.832	0.14	9 002	0.551	75. %1	1.84	●.020		93.932	1.16	0.013	0.171		35 10 32. 60	0.510 0.361		87 74 87.01
final fails	254.00	26.171 100.001	0.05	0 013	4.642	100.001	0.70	0.204	6.072	100.002	0.46	0.120	1 417	100,001	4. 55	•		
reconstituted feed	970.86			0.324	100.00I			3.361	100.001		••••	7.469	100.001	100.301	11.70	3.114 20.330	10.992 100.002	100.00
TEST SUMMARY:	sample	eass pull		copper	recov	er y		lead	recav	êr v		2100	recov			iron		
	wer ght	sample total		assay I	stage	total		assay I		total		assay 1	Stage	total		assay 1	recov Stage	
bulk flotation	556.40	57.311 57.311		0 48	8 5 031	0 5.031		4,77	#5.117	85.111		5.16	39,402	70				
zenr rougher flotation	135.52	13 % 71 271		0 23	•.713			1.79	7.423	92.531		31.15	50.22%			37.80		7£ 45
zami scavenger flotation time 0 1.5 min	24.B6	7.561 73.812		0.15	1 22%	95 961		1.83		93.912		1.65	0.571	97.02% 90 39%		19.26	• 492	
final flotation tails	254.06	26 17 1 100 001		0.05	4.042	100.001		0.78		100.001		0.46		190.00Z		34.02 11.90	3.071 10.991	87.0

10-210 Nitrogen conditioning, then Cu,Pb,pyrite flotation in air, followed by nitrogen conditioning and zinc flotation in air.

\$ 3

FLOTATION CONDITIONS

	pH		Pre-condi	tioning	colle		flotation
sample reference		added g/t	gas	time	type	addition g/t	gas
bulk rougher - 2 minutes	8. 5	-	nitrogen	3 a ln	SIPX	50	alr
bulk scavenger - 2 einutes	8.5	-	nitrogen	3 e in	SIPX	30	alf
zinc rougher - 30 second	10.0	1200	none	5 min	SIPX	50	air
zinc rougher - 1 minute	10.0	-	-	-	-	-	air
zinc rougher - 2 minutes A	10.0	-	•	-	-	-	air
zinc rougher - 2 minutes B	10.0	-	-	-	-	-	air
zinc scavenger - 30 seconds	10.0	, -	-	-	SIPX	30	air
zinc scavenger - i minute	10.0	-	-	-	-	-	3 1r

GRINDING CONDITIONS:

30 minutes, mild steel charge

1200 g/t zinc sulphate

NOTE:

Nitrogen added during xanthate collection period

in zinc flotation stages. (see #)

10-210 Nitrogen conditioning, then Cu,Pb,pyrite flotation in air, followed by nitrogen conditioning and zinc flotation in air.

	SOLIDS W	ASS BALAN	CE		COPPER M	ASS DALAM	Œ	l	eap mass (MLANCE		I	INC MASS B	ial ance		1	ROM MASS 8	AL MICE	
	samie	0445 3	u! l	Cobt	ier	COP	er	lea	d	lea	đ	218	t	216	ιξ	110		ire	,
	weight	sample	total	2552y	enits	reco	ery		units	recov		assay	ensts	18001	r e r y		wnits	16000	er v
sample reference	(graes)			(2)		stage	total	(2)		st age	total	(1)		stage	total	(2)		stage	total
oile rougher - 2 minutes (A)	160 55	16 212	16 711	1 46	0.237	72 261	72 261	13 50	2.188	67.112	67 112	9.05	1 467	19 242	19 242	30 10	4 879	17 221	17 22
bulk rougher - 2 ainutes (8)	49 07	6 9'1	23.181	0 29	0 020	6 172	78.432	4 14	0 287	8.851	75.971	10.10	0.704	7 242	28.481	33.70	2.350	8 301	25 52
on1+ scaveng∞c - 2 minutes (A)	B1 21	8 201	31 301	0 13	0 011	3.251	81 671	2 06	0.169	5 182	81.151	7.19	0 5%	7. '32	36.21%	37 10	3.042	10 741	36 25
bull scavenger - 2 minutes (D)	15.05	1 521	32.901	0.13	0 002	0.601	0 2.291	2.34	0.636	1.091	82 241	11.50	0.175	2 201	38.501	29.30	0 445	1 571	37 63
zini rougher - time 38 seconds	50.75	5 122	18 0"1	0 17	6 004	2.661	84 951	0 99	0.046	1 412	83.65%	46.30	2.373	31.117	49 421	15 30	0 784	2 771	40 5
zanr rougher - time 1 minute	42 01	4 24%	42 271	0 16	0 007	2.071	87 92%	1 20	0 054	1 667	85 31Z	31.20	1.323	17 362	86 972	20.00	0.848	2 992	43 5
zinc rougher - time 2 minutes A	3 <u>0</u> 72	3 912	46.18Z	0.15	0.006	1.791	88 BII	1 42	0 063	1 942	87 261	17 40	0 680	8 921	95 892	24.20	1 142	4 031	47 63
zinr rougher - time 2 minutes B	22.10	2 231	48.412	0.15	0.003	1.022	89.841	1.74	0.039	1.191	88 451	4.67	0.103	1.352	97.251	34.30	0.810	7 851	56 4
zanc scavenger - time 30 seconds	40.67	4.112	52.521	0 00	0.004	1.131	9 0 961	0.90	0.037	1.131	89.581	1.08	0.044	0 581	97 832	43 10	1.770	6 252	56.7
zinc scavenger - time 1 minute	36.56	3.691	56.21%	0.07	0.003	1.011	91.092	6.90	●.033	1.021	90 40I	0.81	●.030	0.392	98.221	42.70	1.576	5 561	62.29
final taris	433 70	43.791	100 00I	0.06	9.024	8.021	100 001	0.70	0.307	7.401	100.001	0.31	0.136	1.781	100,001	24.40	10 485	37.711	100.0
reconstituted feed	990 39				0.328	100.001			3.241	100.001			7.425	100.002			20.332	100.001	
TEST SIMMARY:	sample	B445 p	w11		copper	f e CDV	ery		lead	recov	er y		210C	recov	er y		1100	recov	iery
	weight	sample	total		assay 1	stage	total		assay I	stage	total		assay l	stage	total		assay I	st age	total
helf flotation	229 62	23.181	23.101		1.12	78.431	78 432		10 48	75.971	75.977		7.37	28 482	28 481		31 10	25.521	25.5
pyrite flutation	96.76	9 721	32.901		0.13	3.862	82.297		2.10	6.272	82.24Z		7.86	10.021	3801		35 98	17.312	37.8
zinc rougher flotation	153 58	15.511	48.41X		0 16	7.541	89 841		1.31	6.717			28.89		97.252		23.11	12 65%	
zinc scavenger flotation time 0 1.5 min	77.23	7.89%	56.217		0.09	2.14%	91.087		0.70	2.151			0.95	0.971	98.221		42.71	11 811	62.2
fanat fintation tarks	433.70	43.792	100.002		0.04	8.021	100 00I		0.70	7,402	100.001		0.31	1.701	100.002		24.40	37.711	100.0

10-211 Bulk copper,lead,pyrite flotation in nitrogen then zinc flotation in air

\$ 1

	FL	OTA	T 1 0	N C	O N D	ITI	O N S
, ,	pH a		Pre-con	ditioning		ector	
sample reference		added g/t	gas	time	type	addition g/t	gas
bulk rougher - 2 minutes A	8.5	_		10 📭	SIPX	50	
bulk rougher - 2 minutes B	8.5	+	-	-	SIPX	30	
oulk scavenger - 2 minutes A	8.5	-	-	-	SIPX	50	
bulk scavenger - 2 minutes B	8.5	-	-	-	-	-	
zinc rougher - 30 second	10.0	1200	none	5 min	SIPX	50	aır
rinc rougher - 1 minute	10.0	-	-	•	-	-	alr
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	air
zinc rougher - 2 minutes B	10.0	-	-	-	-	-	air
zinc scavenger - 30 seconds	10.0	-	_	-	SIPX#	30	alf
zinc scavenger - 1 minute	10.0	-	-	•	-	-	air
			5R I ND I	NG CONDITIO	INS:		
			30 mi	nutes. 👊	steel o	charge	
			1200	g t zinc si	ulphate		
	h	OTE:					
	N	litrasen .	added dur	ing kantha	te condi	tioning per	10d (see \$
	,	ir c. odeu	augeu dur	rog zentna	ce condi	croured be.	100 ,256 •

10-211 Bulk copper, lead, pyrite flotation in nitrogen then zinc flotation in air

	50L105 H	MSS BALAN	CE		COPPER N	ASS MALAN	Ci		LEAD MASS	BALANCE		1	LINE MASS	BALANCE		1	ROM MASS	BALANCE	
	sample	8855 9		€09)	per	cop	per	le	ad	le	ad	216	10	21	nr	170	_		
	wesaht	sample.	total	a553y	unsts	reco	very	8558y	wits	reco	very	45547	units	790			units		Cu
Saaple reference	(graes)			(1)		stage	total	(1)		stage	totai	(2)		stage	tetal	(2)	UMIES	stane	total
hulf rougher - 2 arnutes	195 00	18.662	18 447	1 12	0 209	65.721	45.721	10.40	1.740	50.5.1	£4 £10								
hulb cravenger - 2 minutes	115.28	11.037		0.32	9.035	11.102	76.02Z		1.103	20.731	50.531 79.262	7.21 0.52	1.345 0.940	15.042 10.512	15.042 25.542	30.10 32.50	5.616 3.584	23.60Z 15.06Z	23 601 38.661
Pyril+ ronc - In scavenger 1	10.53	1.912	30.692	0.36	0 004	1.142	77.9£I	3 43	0.035	0. 90Z	PO. 162	37.76	4 700						
Pyrile conc - In scavenger 2	6 58	0 63Z	31.321	0 29	0 002	0.571	78.54Z	3.02	0.017	0.501	89 651		0.350	4.251	29.791	16 90	0.159	0.71%	39.37
Dyrite final conc (in scav tails)	328.59	31.441		0.11	0.035	10.951	B9. 421	1.67	0.525	13.672	94,332	38.00	0.227	2.53%	32.322	17.50	0.110	9.46I	39.21
yralm right concentrate	345.70	33.072	••••	0.12	•.•3	12.591	07.424	1.75	4.323	15.072	74.332	4.90 0.39	2.169	24 252 31.032	54.581	35.54	11.475	49.212 49.392	86. 65
rinc rougher - tion 30 seconds	45 49	4.351	67.112	0.17	0.007	2.331	91.741	0.75	0.033	0.851	95.18I	50.40	3 107	44	-				
rinc rmigher tive formite	24.02	2. 301	69.412	0.19	0 004	1.372	93.121	0.94	0.022	9.562	95.74X	40.30	2.193	24.521	01.10Z	9 05	0.750	1.472	80 52
ranc rougher - trae 2 minutes	15.73	1.501	70.912	0.23	0.003	1.091	94.201		0.017	0.50I	96.24 <u>Z</u>	29.30	0 ¶26 0.426	10 352 4.761	91.452 96.212	10.40 13.70	0.239 0.206	1.002	91.39
rinc scavenger - time 30 seconds	13.11	1.25%	72.172	0.22	0.003	0.872	95.07Z	1.47	0.018	0.48Z	96.722	12.30	A						
zinc scavenger - time ! minute	13.06	1.251	73.421	0.17	0.002	0.751	95.021	1.36	0.017	0.441		5.40	0 154 0.070	1.722 0.782	97.947 98 .721	16 BO 15 90	0.211 0.199	0.89Z 0.83Z	93.11
final tails	277.84	26.5 8 Z	100.002	0.05	0 013	4.182	100.00Z	0.41	0.107	2. 847	100 00Z	0.43	0 114	4 300					
reconstituted feed	1045.23				0.318	100.00Z	•		3.839	100.001	100 402	4.13	8.944	100.001	100.v01	6.17	1.640 23.7 99	6.871 100.001	100.00
TEST SIMMARY:	saepie	0055 p	od 1		copper	recov	iery		lead	recov	er y		ZINC	recov	Bru				
	weight	sample	totai		assay I	stage	total		assay I	stage	total		assay 1		total		iron assav I	recov stage	
bulk flotation	310.20	29.69%	29.692		0.82	76.022	76.071		10.25	79.261	70.262		7.70	25,542					
Pyrite flotation-rougher concentrate	345.70	33.072	62.751		0.12	12.591			1.75	15.07%	94.331		0.39	23.34% 31.03%	25.541		20.94	30.642	
zanc rougher flotation	85.24		70.91%		0.19		94. 20%		0.90	1.917	96.241		43.48		56.581		35.54	49.392	
zinc scavenger flotation time 0-1.5 min	26.17	2.50I	73.421		0.21	1.611			1.42	0.921	97.162			37.64Z	96.212		۰, 75	3.341	41 10
final flotation tails	277.84		100.002		0.05		100.002		0.41		100.001		9.96 0.43	2.511	90.721 100.001		16.35	1.72%	93.11

10-212 Bulk copper, lead flotation in air, then pyrite flotation in nitrogen with zinc reverse cleaning, finally zinc flotation in air.

	pH	activator	Pre-cond:	tioning	coll	ector	flotation
sample reference		added g/t	gas	time	type	addition g/t	gas
bulk rougher - time 2 minutes	8.5	_	air	10 min	SIPI	50	air
bulk scavenger - time 2 minutes	8.5	- ,	-	-	SIPX	30	air
pyrite rougher - time 2 minutes A	8.5	-	nitragen	2 -10	SIFX	5	nstrogen
pyrite rougher - time 2 minutes B	8.5	-	nitrogen	2 min	SIPX	5	nitroger
pyrite rougher - time 2 minutes C	8.5	-	nitrogen	2 018	SIPX	5	nitrogen
pyrite rougher - time 2 minutes D	8.5	-	nitrogen	2 min	SIPX	5	nitroger
zinc rougher - time 30 seconds	10.0	1200	-	5 ain	SIPY	50	air
zinc rougher - time I minute	10.0	-	-	•	-	-	alf
zinc rougher - time 2 minutes	10.0	-	-	-	-	-	air
zinc scavenger - time 0.5 minutes	10.0	-	-	-	SIPX	30	air
zinc scavenger - time 1 minute	10.0	-	-	•	-	-	911
zinc reverse rougher - 1 minute	10.0	200	none	5 min	SIPX	50	air
zinc reverse scavenger - 1 minute	10.0	•	•	•	-	-	air

SFINDING CONDITIONS:

30 minutes, mild steel charge

Zinc Sulphate, 1200 g/t

10-212 Bulk copper, lead flotation in air, then pyrite flotation in nitrogen with zinc reverse cleaning, finally zinc flotation in air.

	SOLICE IN	FSS PALAM	:[CUPPER MA	SS MLANC	E	L	EAD MASS I	ALANCE		11	INC MASS B	AL AMEE		Į	RON MASS B	LAICE	
	sample	#355 PI	el l	0000	er	COPP	er	lea	đ	lea	đ	21 N	:	218	t	110	•	110	
	wer ght	sample	total	92294	umits	recov	er y	assay	units	recov	er y	8558Y	units	recov	ery	assay	units	recov	2 7 y
sample reference	(graes)			(Z)		stage	total	(2)		stage	totai	(2)		stage	total	(2)		stage	total
bull emigher 2 arnutes	130 07	17 082	12.08I	1.70	0 705	64.961	-	16.89	2.030	62.202	62.201	8.24	0.996	11 042	11.042	25.40	3.070	12.54%	12.541
bulk scazenger - Z minutés	72.59	6 742	10.832	0.39	0.026	₹.321	73.2 0 T	5.33	0.359	11.012	73.212	11.40	0.769	8.532	17.572	29.60	1.996	6. ISI	20.69
pyrate conc - In scavenger 1	15.52	1.442	20.271	0.21	● 003	0.961	74.242	1.41	0.023	♦.712	73.922	44.20	0.637	7.972	26.641	14.50	0.209	0.852	21.54
pyrite conc - In scavenger ?	13.59	1.267	21.532	0.22	9.003	0. 88 Z	75.122	1.96	0.025	0.761	74.682	33.50	0.423	4.602	31.331	20.50	0.259	1.061	22.60
myrite (ina) conc (2m scav tails)	408.10	37.921	57.451	0 11	0.042	13 192	80.311	1.48	0.561	17.192	91.872	4.86	1.143	20 44Z	51.762	40 50	15.356	62.72%	85.32
pyrile ighr concentrate	437.21	40.421		9.12		15 021		1.50		10.661		7.15		32.192		38.96		64.632	
zine enligher - ti on 3 0 seconds	30.32	3 651	63. 10Z	0.16	0.005	1.051	90.151	0.83	0.030	0.932	92.802	59.10	1.630	20 302	72 062	7.12	9. 773	1.362	86.68
zanc erngher - time 1 minute	42.47	3.952	67.95%	9.19	0.007	2.371	92.527	1.02	0.040	1.231	94.04Z	41.70	1.645	18.251	90.311	12.40	0 489	2.002	88.66
zanc rougher - time 2 minutes	22.56	2.101	69. 152	0.19	0.004	1.261	73.791	1.44	0.030	0.921	94.96I	26.50	0.55 5	4.167	76.472	14.40	0.348	1.421	90. 10
gant sis enger - time 39 seconds	17.67	1.647	10.792	0.22	0.004	1.142	94.932		0.024	0.731		8.06	0.132		97.431		0. 202	1.192	91.29
zinc sravenger - time 1 minute	12.92	1.20%	71. 99 %	9.17	♥. 002	0.65%	95.571		0.017	0.532	96.227	4.56	0.055	0.617	98.54Z	16.60	0.199	0.817	92.1
zanc scavenger - time 2 minutes	0.00	0.091	71.992	0.18	0.000	0.00I	95.571	0.00	9.000	0. 6 02	°6.221	9.00	0.000	0.90I	98.541	0.00	0.000	0.00Z	91.2
final tails	301 50	28,012	100.002	0.05	0.014		100.001	0.44	0.123		100,001	0.47	0.132		100.002	6.90	1.933	7.892	77.1
reconstituted feed	1076.31				0.316	190.00%			3.264	100.00Z			7.018	100.00Z			24.485	100.002	
TEST SUMMORY:	sample	#255	pul l		copper	recov	ery		lead	recov	ery		ZIRC	recov	er y		1100	f e C 0 v	er y
	weight	sample	total		assay Z	stage	total		assay 1	st age	total		assay l	stage	total		assay Z	stage	tota
bult flotation	202.66		10.831		1.23	73.201			12.69	73.212			9.37	10.572			26. •0	20.672	
pyrite flotation	437.21		59. 452		0.12	15.072			1.50	18.662			7.15	32.19%			38.76	64.631	
zinc +nugher flotation	104 35	9.7AT	67.152		0.10	5.4BT			1.04	3.092			41.58	44.70%			12.07	4.782	70.1
zinc scalenger flotation time	30.59	2.842	71.99%		0.20	1.792	95 572	!	1.45	1.262			6.58	2.071	98.54 ₹		17.20	7.011	•7.1
final flotation tails	301.50	28, 612	100.002		0.05	4.431	100.001	l l	0.44	3.701	190.002		0.47	1.462	100.007		6.90	7.092	100.0

10-213 Bulk copper, lead flotation in air, then pyrite flotation in nitrogen with zinc reverse cleaning, finally zinc flotation in air. (repeat of test 10-212)

sample reference	pH	activator added	Pre-cond:	tioning		ector	flotation
		g/t	gas	time	type	addition g/t	gas
oulk rougher - time 2 minutes	8.5	-	air	10 ain	SIPX	50	air
oulk scavenger - time 2 minutes	8.5	•	-	•	SIPX	30	air
ovrite rougher - time 2 minutes A	8.5	-	nitrogen	2 min	SIPI	5	nitragen
pyrite rougher - time 2 minutes B	8.5	•	nitrogen	2 -10	SIPX	5	nitragen
pyrite rougher - time 2 minutes C	8.5	-	nitrogen		SIPI	5	nitrogen
pyrite rougher - time 2 minutes D	8.5	•	nıtrogen		SIPX	5	nitrogen
ovrite rougher - time 2 minutes E	8.5	-	natrogen		SIFX	5	nitragen
pyrite rougher - time 2 minutes F	8.5	-	nıtrogen		SIPX	5	nıtragen
pyrite rougher - time 2 minutes 6	8.5	-	nitrogen		SIPI	5	nitrogen
pyrite rougher - time 2 minutes H	8.5	-	nitrogen	2 a ın	SIPX	5	nıtrogen
zinc rougher - time 30 seconds	10.0	1200	_	5 min	SIPI	50	air
rinc rougher - time 1 minute	10.0	-	-	-	- -	-	air
zinc rougher - time 2 minutes	10.0	-	-	-	-	-	217
zinc scavenger - time 0.5 minutes	10.0	•	-	-	SIPX	30	air
zinc scavenger - time 1 minute	10.0	•	-	-	-	-	air
zinc reverse rougher - 1 minute	10.0		none	5 a in	SIPX	B	air
zinc reverse scavenger - 1 minute	10.0	•	•	-	-	-	air
			6RINDINS	CONDITIO	NS:		
			30 minut	es, eild	steei o	targe	
			11nc 51	phate. 1	700 /+		

10-213 Bulk copper, lead flotation in air, then pyrite flotation in nitrogen with zinc reverse cleaning, finally zinc flotation

יות ביין אר אין ארוניים מים ביי

	SOLIDS W	ASS BALAK	CE		COFFER M	ASS BALAN	EE		LEAR MASS	MLANCE		;	ZINC MASS	BALANCE		1	PON MASS	PALANCE	
	saecie weight	Mass p Sample		1907		Cop rece		le	ad units	-	ed .	211		21	ınc	iro	n	17.	ne .
Sample reference	(grams)	2400.1		42741	4 11713	stage	•	B3367	autr2	reco stage	total	3 55 8 7	unsts	reco Stage	total	45544	units	reco	1 ery
bulk rounter time 1.5 minutes	20 30	2 502	2 50Z	1 43	0.036	9.672	9.672	12.50	0.313	4.621	4.622			_				,.	
bulk scavenger - time 1.5 minutes	68.59	8.447	19.942	1.93	0.163	44.062	53.731	43.40	3.463	54.99%		7.13 7.8 9	0.178 0.835	1.512 8.932	10.842		0 688 1 823	2.831 7.471	17 22
Byrite rousher - time 2 minutes A	58.60	7.222	10.161	0 53	0.038	10.35%	64.082	11.60	0.852	12.501	71.291	11.30	4 4						
Byrste congher - fime Z minutes B	48.00	5.911	24 087	9.30	6.018	4.801	48.861	6.11	0.361	5.342	76 637	•. 44	0.816	8. 731			2.224	•. 147	19.46
Pyrate compler - tree 2 pinutes C	116 70	14.382	38 462	0.16	0.023	6 221	75.10X	2.88	0.414	6.172	82.74I	7.05	0.558	5.471	25.542		7.047	8.412	27 81
Byrste rougher - time 2 minutes B	82 00	10.231	48 692	0.14	0.014	3.872	78.972	2.72	0.227	3.352	85.107	7.05 5.80	1.014	10.84I		32 70	4 702	17.372	47.20
									*****	J. JJ.	03.174	3.89	0.503	4.351	42. 332	39.00	3. *68	14.31%	63.51
zinc rougher — fine 38 seronds	29 60	3.651	57.3"	0.17	0.006	1.681	90.652	2.02	0.074	1.002	e*.101	45.00	1.674						
zinc rougher - time 1 ainste	32 10	4.032	56 '62	0.21	0 008	בַּכַרַ בַ	87.942	2.36	0.005	1.402	98.592	44, 29		17.912	• • • • • •	4.60	0.761	1.497	65 0
zinc romher - time 2 minutes	26 30	3.242	50.601	0.20	8.009	2 541	85.48I	3.37	0.109	1.612		31.40	1.781	19.052		11.50	0.467	1.973	65.7
				_							3.204	31.40	1.018	In Bal	90.572	15.60	0.504	2.061	£9.90
zinc scazenger - fine 0.5 minutes	•.70	1.701	Pu Bus	0 36	0.904	1.157	86.642	4.49	9.054	0.792	00.002	16.10	9.193	2.061	83 435	** **			
zinc scavender – time 1 minute	€. 10	1.121	61.072	0.33	0.004	1.007	87.64I	4.14	0.046	0.491	°1.697	22.70	0.255	2.771			0 Zo2	1 201	
												*****	V. 233	2.721	€ 5.361	21.70	0.243	1.002	71.20
final tails	3119.03	38.061	100.002	0 12	0.046	12.362	100.002	1.48	0.564	8.32X	190.001	1.14	0.434	4.44Z	100.002	18.40	7.097	29.801	190.50
test food (reconstituted)	011.53				0.370	100.00I			6 772	100.001			7.348	100.002			24,320	100.001	
TEST SHMMARY:	saeple	0355	ol l		Copper	FECOV	er v		lead	FECOV	er,		****						
	weight	sample	total		assay 1	stage	total		assa) I	stage	total		ZIAC assay 1	7007			17.08	FECO	
	•				•	•			-, -				43367	st age	total		assay I	st age	total
hulb flotsbyon	98.89	19.942	10.047		1.02	52 772	53.732		36.34	58.712	58.71%		•. 26	10.847	10.847		22.05		
pyrite rougher flotation in nitrogen	306.30	37.74%	48.691		0.25	25.741	78.972		4.01	27.391	86.10Z		7.90	31.001			22.05	19.322	10 3
RENC COMPINE FEOTISTION	88.60	10.922	59.501		0.22	4.511	85.48Z		2.55	4.102	99.292		40.97	47.852	90.572		34.20	53.102	67.51
zinc sravenori fintation	18.89	2. 122	61.972		0.75	7.162	87.64Z		4, 22	J. 487	°1.687		17.20	4.782	95.36I		12.22	5.491	70'W
final tells	309.03	38.081	100.001		0.12	12.361	100.002		1.48		100.001		1.14		73.361 100.002		23.14 18.40	7. 202 20. 801	71.70

10-214 Bulk copper, lead flotation in air, pyrite flotation in nitrogen, zinc flotation in air

FLOTATION CONDITIONS		L	0	T	A	T	I		D	×		C	0	N	D	I	Ţ	I	0	N	S	
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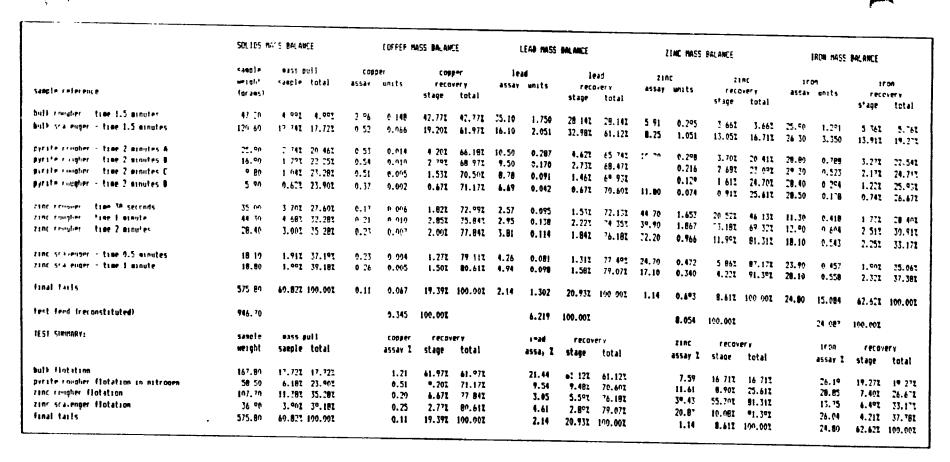
	рH		Pre-cond:	tioning		ector	flotation
sample reference		added g./t	gas	time	type	addition g/t	gas
bulk rougher - time 1.5 minutes	8.5	-	aır	10 min	SIPX	50	alr
bulk scavenger - time 1.5 minutes	8.5	-	-	-	SIPX	30	air
pyrite rougher - time 2 minutes A	8.5	-	nitrogen	4 min	SIPY	6	nıtrogen
pyrite rougher - time 2 minutes B	8.5	-	nitrogen	4 min	SIPY	6	nitrogen
pyrite rougher - time 2 minutes C	8.5	-	nitrogen	4 min	SIPX	6	nitrogen
pyrite rougher - time 2 minutes 9	8.5	-	nitrogen	4 a in	SIFX	6	nitrogen
zinc rougher - time 30 seconds	10.0	1200	-	5 min	SIPI	50	air
zinc rougher - time ! minute	10.0	-	-	•	-	-	alr
zinc rougher - time 2 minutes	10.0	-	-	-	-	-	alf
zinc scavenger - time 0.5 minutes	10.0	-	-	-	SIPY	30	alr
zinc scavenger - time 1 minute	10.0	-	-	-	-	-	air

GRINDING CONDITIONS:

30 minutes, mild steel charge

Zinc Sulphate, 1200 g/t

10-214 Bulk copper, lead flotation in air, pyrite flotation in nitrogen, zinc flotation in air



10-215 Bulk copper, lead flotation in air, pyrite flotation & zinc flotation in air.

F	L	0	Ţ	A	Ţ	I	0	N	C	0	N	Đ	I	Ŧ	I	0	N	S	
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sample reference	pH	activator added	Pre-coi	nditioning	coll type	ector addition	flotation
Sample (e.e.e.e.e.		g/t	gas	time	type	g/t	gas
bulk rougher - time 1.5 minutes	8.5	-	air	10 min	SIPX	50	air
bulk scavenger - time 1.5 minutes	9.5	-	-	-	SIPY	30	a 15
pyrite rougher - time 2 minutes A	8.5	•	air	4 min	SIPX	6	air
pyrite rougher - time 2 minutes B	9.5	-	air	4 @10	SIPX	Ł	315
pyrite rougher - time 2 minutes C	8.5	-	air	4 min	SIPX	6	air
pyrite rougher - time 2 minutes D	8.5	-	alf	4 min	SIPX	6	alf
zinc rougher - time 30 seconds	10.0	1200	-	5 e in	SIPX	50	alr
zinc rougher - time i minute	10.0	-	-	•	-	-	air
zinc rougher - time 2 minutes	10.0	-	•	-	•	-	air
zinc scavenger - time 0.5 minutes	10.9	-	-	-	SIPY	30	air
zinc scavenger - time 1 minute	10.0	-	-	-	•	-	air

GRINDING CONDITIONS:

30 minutes, mild steel charge

Zinc Sulphate, 1200 g/t

10-215 Bulk copper, lead flotation in air, pyrite flotation & zinc flotation in air.

	SOLIDS A	ASS BALAN	CE		COFFEP N	ASS PALAM	C.E.		LEAD MASS	BALANCE		1	INC MASS	BAL AMEE		1	IPON MASS	BAL ANCE	
	sample	0255 p		cop		COD	•	le		le	ad	218	ıc	210	B.C	174			
	weight	sampie	tota!		units	reco			units	reco	VEF V	3553y	units	reco			units	17	-
sample reference	(graes)			(2)		stage	total	(1)		stage	totai	(1)		stage	total	(2)	umit's		verv total
ultrougher Toundes	178 55	14 191	14 392	1.48	0 213	67 232	67 232	13 60	1 957	65 532	e5.531	6. 70	0.964	17 485		••		•	
wik scavenger - 2 minutes	65.77	6.032	21 22%	9.41	0.029	8 842	76 07E	4.20	0.292	9 791	75.321	9.30	0.464	13.002 0.622	13.082 21.691	35.16 31 20	5.052 2.132	1º 5ex 0.25x	
rinc religher. Time 30 Seconds	71 17	7 797	28 67I	0.15	0.011	3 501	79 572	1.34	0.099	3 322	79.64Z	35.69	2.632	75	F3				
rane emiglier - take flaanste	52 89	5. 492	34.112	0 18	0.019	3 122	87 602	1.60	● 068	2.942	81 58I	20.60	1.502	35.402	57.307	14.70	1.007	4.211	
zanc ronigher - 1100 Z name - A	50 21	5.271	In sil	0 18	0 009	2.062	85 66Z	1.70	● 089	2.971	84.552	15.60	0 814	21.462	78 832	19 70	1.082	4.192	35.2
zanc imigher - fame 2 manufes B	20.52	2.962	42 70"	0 18	0.005	1 681	87.34%	1.64	0.040	1 632	86.162	9.90	0.264	11.03Z 3.58Z	89.872 93.442	25.70 27.00	1.340 0.800	5.1°2 3.102	91.46 94.56
ting scarenges - time IO seconds	20 95	2.181	44,472	0 17	0.004	1 173	80.51%	1.50	0.033	1.092	87.271	5, 20	0.113	1.572	84 801	•• ••		_	
rinc reavenger - time I minute	Z9 65	3.121	47.592	0 16	0.005	1 :02	*0 98Z	1.41	0.044	1.472	68.742	3.49	0.113		94. 982	34.40	0.749	2.901	_
zinc scaveliger - time 2 minutes A	37.48	3.892	51.497	0.14	0.005	1 '21	91.80%	1.35	0.053	1.762	90.501	2 41	0.094	1.492	76.452	35.10	1.695	4. 242	51.6
zinr stavenger - time 2 minutes D	27.49	2.852	54.372	0.11	0.003	9.997	92.79%	1.14	0.032	1.091	91.591	1.40	0.040	1.272 0.542	97.732 98.271	33.40 33.50	1.3(0 0.753	5.062 3.692	56 76 60.3
final tasts	419 76	45.682	100.002	0.45	0.073	7.712	100.002	♦.55	0 251	8.412	100.001	0.20	0.128	1 779	100.00Z	33 44			
reconstituted fees	962.67				0.317	100 001			Z.987	100.001		****	7.375	100.001	107.014	21.40	10.231 25.029	3°.612 100.002	109.0
TEST SYMMEY	saepte	9255 B			copper	FPCOV	er y		lead	FECOV	ery		2100	Fecov	et v		11 00	recov	•••
	nesäyf	sample	totai		assay I	stage	total		assay l	stage	total		assay Z	stage	total		assay I		total
bulk flotation	204 32	€.21	21 221		1.14	76 072	76.071		10.60	75. 32%	75.323		7.54	21.492	21.692		33.84	49 4	
zanc rimigher flotation	202.70	0.21	42.29%		0.17	11.772	87.347		1.54	10.862	85.182		25.12	71.751	93.442		_	27.811	
zsuc slavenger flotation time 0-1.5 min	50.98	0.05	47.5%		0.16	2.741	90.081		1.45	2.572	88.74Z		4.19	3.011	70.452		20.46	16.692	44,5
zinc stavenger flotation time 1.5mint	41.89	0.07	54. 323		0.13	2 712	97 192		1.26	2.851	91.591		1.98	1.012	78.772		34.01	7 142	
final fightion tasks	437.70	0.46	100.00I		0.05	7.212	100.00Z		0.55		100.001		0.28		100.007		33.56 22.40	9.762 39.612	

10-301 Standard bulk copper, lead flotation and zinc flotation.

	pH	activator	Pre-cor	nditioning	coll	ector	flotatio
sample reference	·	added g/t	gas	tine	type	addition g/t	gas
bulk rougher - 2 minutes	8.5	•	air	10 a in	SIPX	50	alr
bulk scavenger - 2 minutes	8.5	•	-	•	SIPY	30	air
zinc rougher - 30 second	10.0	1200	none	5 eir	SIPX	50	air
zinc rougher - 1 minute	10.0	-	-	-	-	•	a 1f
zinc rougher - 2 minutes A	10.0	-	-	-	-	-	air
zinc rougher - 2 minutes B	19.9	•	-	-	-	•	317
zinc scavenger - 30 seconds	10.0	-	-	-	SIPX	30	air
zinc scavenger - ! minute	10.0	-	-	-	-	-	31L
zinc scavenger - 2 minutes A	10.0	-	-	•	-	-	a 17
zinc scavenger - 2 minutes B	10.9	•	-	•	-	•	315
			6P.INDI	NE CONDITIO	INS:		
			30 01	nutes. mild	steel	tharge	

10-301 Standard bulk copper, lead flotation and zinc flotation.

sample reference built recipier - 30 seconds & 70.10 built recipier - 30 seconds & 70.10 built recipier - 1 minute & 54 60 built recipier - 1 minutes & 76.70 built recipier - 2 minutes & 31.00 brite recipier - 2 minutes & 31.00 frite recipier - 2 minutes & 31.00	Sample Sample													5	PUR PASS BREAK	ž	
· -	A SEC.	34	, 1	Laddoo	ا يا		,	lead	_	3u t 2		::		5		5	_
-	3,092	(2)	53 tun	recovery stage total	total	A (₹)		recovery stage to	total	4554y an olds (2)		recovery stage tol	fry total	(2)	anits.	recovery stage te	tetal
_	* * * * * * * * * * * * * * * * * * * *	2. (iB	.86 0	26 10g	26.102 3	3,98	1.357	32, 972	17 978		701						
_		82/	9 949	15.47					42. D						. E	200 £	3 042
_	-	6	0 077	11.791						20				_	6.715	1:42	5.711
		44.0	9100	10.67			2				P :				1.142	261 +	100.
	3.871	0	610 0	5 842							. J.				.974	7.23	17.131
		9.38	8 00 6	7.912		3.42	6	2.861	E 632	2 8	6. 127 6. 127		20 602 23.961	32 ES	\$ II	5 262	22. 191 28. 261
		ů.	9 063	0. 991			210 0	1.022	87. 64X	4	£			•		ļ	
	2 65	9.22	9u0 u	2.041	7.6 972		0.067		P4.261		240				287	7.151	<u>.</u>
	2, 101 30 072	0 18	too c	1.301			9 042		95.291		2				١٠:	ř.	707
		9.16	0 003	760 1	17.22	1 62	9.035	1.651	Bé. 131		0.177	1001	101.11	P 2	7 R C	3 217	13. 92
pyrite rahr concentrate 177,80	27.6 8	0.19		5.411		7.07									B	2.152	39.981
tine eviglier - time 29 seconds 152,20	7.687 39 912	0.10	806 v	1 ti			0.071	1.721	-	5	2/3					,	
23116 eruglier time 36 serends 56.10	2.831 42.741	9 13	6,003	1/0 1	82.70I		9.028							_ `	80.	216.5	. 30 I
tine emigher tine I ninite 43 lo	7 197 44 927	<u>-</u>	0 003	0 657	B3.462		0.031						700 00	_		-	1.517
21 ur rangter - frae prante 79.00	168.64 172 1	6 0	to (equ	Ξ	84.77	1.82	0.034	0.872	191.06	2.3					0.	ž.	16.052
truc emigher - time 2 aimiter 90.40	1.562	9 16	00 0	162 :		 \$	0.067								0.20.0	1,7.7	6.3
ring ingler - fine 2 minites 45,70	2 311 53.761	0 18	0 00	1.301	80.372		0.034			2.58	0.060	0.631	78.822	42.16	6 972	3.562	28. 58 28. 58
tinal taits 916 2n	46.24% 100.00%	90.0	9.037	11.632	100.00I	33.0	6.305	7.417	100.001	6 .74	11.0	100					
reconstituted feed 1981.30			0.318	100.001		•	=				=			#.# #	27.279	- 361 - 361 - 361 - 361	3. 3.
1851 CHMMAPY: sample	ling see		Copper	F PC OVPF Y	2	_	lead	FECOVETY		-	70.7						
#EI OPF	sample total		2552y Z	st age	tote1	₹	~	stage	total	ř	**	stage to	total		17 On 4554y 7	stage to	total
Beilf flatation 469.80	7. 267		161	73.012	73.817	-		81.671	81.631		02	נ ניים זר	27. 04.9	•			
pyrite final concentrate 177.80	M.972 32.24		9.19	2.412	17.27				66,137					•			19.07
zinc ringher fintation 389.80	12 222 51.457		0.13	7.852	87.07X				91 782				701				39. vBZ
first flutation tails findl. last In conc.) 961.90	48.551 100.001		9 0.0	12.932	100.001		6. 70	1.222	100.001		. e		100 001	~ 7	2 ; 2 ;		55.701

10-401 Bulk copper, lead flotation in air, pyrite flotation in air

and zinc flotation.

Sample reference	pH a	ictivator added	Pre-cor	nditioning			flotation
sample (E)E) Ence		g 't	gas	time	type .	addition g/t	ĝ϶ċ
bulk rougher - 10 seconds A	°.1	_	317	20 min	SIPY/SAX	4()	air
oulf rougher - 30 seconds B	9.1	_	-	-	-	-	31r
bulk rougher - 1 minute A	0.1	-	-	_	SIPY SAX	5	air
bulk rougher - 1 minute B	9.1	-	-	-	•	-	air
bulk rougher - 2 minutes A	۰.1	-	-	-	SIPY/SAY	5	air
bulk rougher - 2 minutes B	9.1	-	-	-	SIPX/SAY	5	air
pyrite rougher - 2 minutes A	٠.1	-	-	4 #19	SIF 1'SAX	6	air
pyrite rougher - I minutes B	9.1	-	-	4 mis	SIFX/SAX	Ł	alf
pyrite rougher - I minutes C	9.1	-	-	4 #10	SIF t 'SAX	6	air
ovrite rougher - I minutes D	9.1	-	-	4 #10	SIPI /SAX	6	air
zinc rougher - Id seconds A	10.2	600	none	jo sin	SIPY'SAX	25	air
zinc rougher - Iv seconds B	10.2	-	-	-	-	-	air
zinc rougher - 1 minute A	10.2	-	-	-	-	-	31"
zinc rougher - 1 minute B	10.2	-	-	-	SIFI 'SAX	5	alr
zinc rougher - 2 minutes A	19.2	-	-	-	-	-	a1f
zinc rougher - 2 minutes B	10.7	-	-	-	SIPX/SAX	5	air
NGTE:			6RINDII	NG CONDITIO	NS:		
Soda asr esed for pH adjustment dur	ına bulk fi	letation	25 •11	nutes. s :ld	steel char	roe. BMS	a:11

10-401 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation.

	50L105 %	ASS BALANCE		COFFEE NA	SS PALANC	£	Li	EAD MASS B	ALANCE		71	INC MASS B	AL AN(E		Įį	ROM MASS 9	# WACE	
	9,996	eace soll	(001	er	copp	27	lea	1	lea	ć	2180	•	210	:	110		11.00	a
	⊯e ±ght	sample total	Secti	units	recov	ery	4554v	enits	recov	er y	9229A	units	FEEDV	er v	25541	units	re c.	e ry
Sample reference			121		stage	*otal	(Z)		stage	total	(1)		stage	total	(1)		stage	totai
Bull-rougher 30 seconds A	op no	5 281 5 281	1.64	() (14F.)	24 712	24 711	2º 31	1 549	39.911	39 017	8 30	0.479	4 801	4 902	23 44	1 778	4.923	4 871
bull inuglier *A seconds B	45 70	2 467 7 75%	1.76	0.04"	12 172	37.082	19 58	0 482	12.431	52 34%	0.52	0.210	2 302	102	27 88	0 687	7.681	7 501
bull conglier - 1 sinute A	95 70	5 171 12.891	1 04	0.054	15 571	52.591	10 02	0 514	13 252	65 60I	9 86	9 506	5 542	12 642	39 44	1.562	P Jas	13.5=
bulk rougher - I grante B	71 19	4 102 17 972	9 84	0 014	9 56Z	62 151	7.10	e 301	7.752	77 752	10.16	0.4.6	4 561	101	23, 55,	1 -35	5 471	14.01
bull-rowher 2 ainstes A	ive au	5 871 22 941	0 46	זרט ס	7.701	69.851	4 80	0 287	7 262	80 612	19 77	0.690	6 571	77 871	33.72	i cau	7 717	76 17
bulk emplee - 2 armutes #	51 110	2 757 25 692	0 44	9 012	3.451	73.301	4 34	0 119	3.081	81 981	10 50	0 384	3 162	27 021	31 80	0.674	3 412	30 1.
certife counties 2 minutes A	70 6"	3 811 20 491	0.20	(<u>ค</u> .ค	2 171	75 471	2.22	0 .84	2.191	95 861	8 34	0.317	7 497	10 512	31 97	1.215	4, "32	34.5
pritergagger 2 signites 8	175 70	7 37% % B1%	0.16	110 6	3 76%	79.272	1 14	780 0	2.15%	B8 011	5 40	0.395	4 771	34 841	38.19	: 'a'	10 Pal	45 1
pyrite reigher - 2 minutes C	228.80	12 772 49 151	¢ 16	959 0	178 2	84.86Z	1 00	0.123	3.1 0 2	of leg	5 30	0.654	161	4; QQI	38 98	4 509	18 732	64.4
syrite rougher - ? sinutes B	241.50	13 022 62 172	0 15	0 021	5 942	90.80Z	0 00	0.117	3.021	94.211	10 34	1.346	14 742	55 742	35.86	1.660	18 1°Z	82.6
pyrite rahr concentrate	674.60	% 48I	0 17		17.50%		1.17		10.531		7 44		Id JUI		34.95		52.511	
zinc cougher - time 30 seconds	9 2 40	4 98" 67.15%	0 14	0 007	1 991	92.792		0.036	0.921	95.142	46 32	2.307	25 271	82 002	11.78	0 5P7	2 291	64 0
tion anugher - time 30 seronds	39 4 0	2 172 67 272	0 16	0 003	0 971	93.761	0.94	0 018	0.462	05 571	45 38	0.964	10 151	90 561	11.68	0 749	0 974	85.6
zinc cougher - tise minute	21.69	1 167 70 447	9.12	9 991	0.402	74.167	1.16	0.014	0.35I	5 942	34.12	0.397	4 352	95 911	16.78	0 105	1 762	86 5
zanr annaher - tame manute	23 60	1 272 71 712	0.14	0.003	0 511	94.661	1.10	0.015	0.39%	ob 332	7.22	0.092	1 917	97 912	29.00	0 -60	1.442	88 4
zinc emigher - time 2 minutes	44 40	? 307 74 102	9 12	0 003	0 827	95.48Z	1.05	0.025	0 65%	96 78 1	2 B0	0.067	9.732	98 65%	28 50	0 683	2.661	e6 ;
zinc rougher - time 2 minutes	7.20	0 501 74.601	0 12	0 001	0.172	95.651	1.66	0.005	0.14Z	97.121	2.30	0.012	0 132	98.787	28.16	0 140	0.542	*1 7
final table	471. 20	25.402 100.002	0.06	0 015	4 352	190.001	0.44	0.112	2.801	100.00Z	0.44	0.112	1.221	109.00 I	8.80	2.235	8 711	100.0
reconstituted feed	1854.90			0.351	100.001			3.880	100.00I			7.132	100.00Z			25.669	100.00Z	
TEST SIMMARY:	sample	mass pull		copper	recov	,		lead	recov			ZINC	recc+	•	•	1 r on	f ec av	r er y
	we 1 ght	sample total		assay I	st age	total		assay I	stage	total		9229A J	stage	total		assav 1	stage	tot
bull flotation	476.50	25 697 25 692		1.00	73.301	73.301		12.64	83.681			7.61		27.03%		30 10	30.17	
pyrite flotation	676 60	36 491 62.171		0 17	17.501	70.80Z		1.12	10.532			7.44		56.74%		₹6.95	52 511	
zinc enuglier flotation	221 49	11.947 74 107		0.14	4.68%			0.90	2.771			32.07	41 912			17 44	8 117	
final flotation tails	480.40	25.901 100.001		0.06	4.521	100 00X		0.45	3.021	100.002		0.48	1.35%	190.001		°.17	4.751	100.

10-402 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation

test number	FL	D T A	TION	i C	O N D I	1 1	0 N S
sample reference	pH at	ctivator added	Pre-cond:	tioning		or ddition	flotation Qas
•		g/t	gas	time		g/t	•
pull rougher - 30 seconds A	c. ₁	-	31 <i>r</i>	20 min	SIFX (SA)	40	air
bull rougher - 30 seconds B	0.1	-	-	-	-	-	315
bulk rougher - 1 minute A	0.1	-	-	-	SIPX/SAX	5	air
bull rougher - 1 minute 8	°.1	-	-	-	•	-	ā1"
bulk rougher - 2 minutes A	0.1	-	-	-	SIF1 SAX	5	aır
bull rougher - 2 minutes B	0.1	-	-	-	SIF t'SAX	5	air
pyrite rougher - 2 minutes A	9.1	-	nitrogen	4	SIFT 'SAT	Ł	nitrogen
pyrite rougher - 2 minutes B	°.1	-	nitrogen	4 min	SIPY SAX	6	nitrogen
pyrite rougher - I minutes C	9.1	-	nitrogen	4 min	SIFY SAY	b	nitrogen
pyrite rougher - 2 minutes D	9.1	-	nitrogen	4 min	SIFY SAX	6	nitrogen
zinc rougher - 30 seconds A	10.2	630	none	10 a ia	EIFY/SAX	35	air
zinc rougher - IO seconds 8	19.2	-	-	-	-	-	air
zinc rougher - 1 minute A	10.0	-	-	-	-	-	air
zinc rougher - 1 minute 8	10.2	-	-	-	51P1/5AX	5	alr
minc rougher - 1 minutes A	10.1	-	-	-	-	-	air
zinc rougher - 2 minutes B	10.2	-	-	-	SIPY/SAX	5	31"
NOTE:			6F INDIN6	CONDITI	ONS:		
Soda ash used for pH adjustment du	ring bulk fl	lotation	25 m inu	ites. mili	d steel cha	rge. BMS	i mill
Lime used for pr adjustment during	zinc flotal	tion	150 g/t	sulphur	dic :de		

10-402 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation

	SOLIES M	ASS BALANCE		COPFEP N	NSS BALANC	Œ	i	LEAD MASS	BALANCE		1	INC MASS	PALANCE		1	IPON MASS	PALANCE	
	sample	@ass #ull	cop	er	COPS	ier	le	af	ìe	ad	211	ıc	211	nr	151			
	weight	sample total	assay	units	F 00 0 1	rery	2553y	waits	reco	VET Y	355 3y	wasts.	reco	-			17: F e co	
Sample reference			(∑)		stage	total	(Z)		st age	total	(2)		stage	total	(2)	84173	stage	total
bull enuglier 30 seconds A	101-10	5 061 5 061	2 52	0.178	32 B4I	32. 841	28.40	1.438	38. 291	38.201	7.56	0.383	4 679	4 ***			_	
bull rinigher 10 coronds B	37.70	1.872 5 952	1 84	0 035	8.94Z	41.782	17.60	0.370	9.072	48.072	B. 36	0.363	4.532	4 512	20 20	1.023	4, 331	4, 17
bulk rougher I minute A	64.70	3.241 10.191	1 12	0.034	9,742	51.12%	15.10	0.489	13.001	61.022	10.30		1.072	6 412	22 29	0.41*	1.772	6.10
Bulk rougher I minute A	97 90	4.402 14 502	0 84	0 037	9.52%	٠٥.632	0.35	0.367	7.761	70 792	10.30	0 334	3.952	10 362	25.50	0.026	3.502	7.60
bull regirer 2 minutes A	Ive au	5.451 20 041	0.47	0 026	8.602	67.231	5.15	0.281	7.462	79.25%		0.453	5.372	15.732	29 70	1.397	5.532	15.13
Pulk rougher 2 minutes B	41.30	2.072 22.117	9.45	0.007	2.492	49.63X	3.70	0.077			11.20	9.611	7.241	22.•71	30.60	1.66	7.062	?? 19
There is a second of the secon	41.30	2.0/2 22.110	**. 43	0.007	2.492	87.834	3./#	W. U//	2.031	80.281	11.10	0.230	2.722	25.692	31.70	0.656	2.772	34. ℃
pyrate ronc. In scazenger 1	16 79	0 821 22.931	9 14	0.001	0.201	69.921	1.38	0.011	9.301	89.581	42.40	0.348	4.122	29 B12	17 60	0.111	0.4°Z	25 43
pyrate comr. In scavenger 2	24.29	1 212 24 143	0 15	0.002	0.47%	79 3°Z	1.74	0.024	0.621	81.21%	39.30	0.476	5 647	35.452	17.10	0.207		
pyrite conc. In scavenger 3	43.30	2.172 26.312	0.21	0.005	1.17%	71.56%	2.39	0.052	1.38I	87.581	21.87	0.473	5.602	41.051			0.091	26. 11
pyrite final rour (Zn scav tails)	513.00	25.73% 52.03%	9.17	9.044	11.267	87. 82I	1.22	0.314	0.342	90.922	1.64	0.424		_	26.50	0.575	2.432	28 74
pyrite ruhr concentrate	597.60	29.921	0.17	••••	13.191		1.34	••••	10.642	·V. 724	5.74	V. 422	5.001 20.362	44.051	37.50 35.22	9 647	40.832 44.602	69.57
zinc emugher - time 30 seconds	103.20	5.172 57 202	0 16	0.006	2		0.79								23.22		44.604	
TING CONGLET TIME TO SECONDS	46 60	2.311 59.511	0.13	0.003	2.132 0. '81	04.95% 03.73%	0.79	0.041	1.081	*2.01Z	41.50	2.144	25.417	71.462	10 20	0.527	2.232	71.00
• • • • • • • • • • • • • • • • • • • •								0.018	0.492	92.50%	42.60	0.994	11.702	03.242	10.00	0.252	1.072	72 86
Zanc i migher tine Lainnte	34.99	1.702 41.242	0.17	0.003	0.742	86.472	1.02	●.017	0.4£Z	92 .9 61	41.00	0.698	8.271	91.512	13 30	0 226	0.96I	73.82
zinc rougher - time 1 dinute	28 50	1.432 67.662	0 20	0.003	0.772	07.21%	1.52	●.022	0.502	93.531	25.30	0.361	4.282	95.792	10 5	0.278	1.197	75.00
zanc religher - tame 7 maintes	21.09	1 072 63.757	ŷ 19	9.002	0.577	87.742	1.57	0.017	0.46Z	93 901	17.90	0.195	2.321	90.112	20.80	0.227	0.967	75 %
zinc rougher - time 2 minutes	33.50	1.682 65.432	0.16	0.003	0.691	80.431	1.33	0.022	0.59Z	74.581	4.30	0.073	9.972	78. 982	27.50	0.461	1.957	77.91
final tails	690.40	34.572 100.001	0.13	0.045	11.572	100.001	0.59	0.204	5, 427	100.001	0.25	9.084	4 430	*** ***				
reconstituted feed	1997.20			0.380	100.00I		••••	3.764	100.00Z	100.004	V. 23	0.439	100.002	100.00I	15.10	5.220 23.431	22.0°2 100.002	100.00
TEST SURBARY.	samle												21111111			23.031	100.091	
IE31 (Manual)	me styt	mass pull sample total		Capper	recov			lead	recov	•	•	2100	FECEV	ery		1790	FPCOV	er y
	me gut	Sample Colai		assay 1	st age	total		assay I	stage	total		assay I	stage	total		assay l	st age	tctal
bulk flotation	441.60	22.112 22.113		1.22	69.672	49 43%		13 67	80.28Z	80.282		7.00	75.692	25.691		26.68	24.962	24 60
pyrite final concentrate	513 80	25 73% 47.84%		0.17	11 76%	80.88I		1.22	8.34Z	00.62%		1.64	5.001	30.492		37.50		24.95
zinc from pyrite float	83.80	4.20%		0.18	1.932			2.06	2.301			30.90	15.361	30.074			49.831	65.79
zanc rampher flotation	234.10	11.72%		0.16	4.92%			0.78	3.07%			37.40	52.052			21.20	3.782	
tinc rombined concentrate	317.90	15.92% 63.757		0.17	6 B6Z	87.74Z		1.27		93.991		35.74				12.09	4.301	
final finiation tails (incl last zinc con)	723.90	36.251 100.001		0.13		100,00Z		0.62					67.422	78.112		15.10	10.172	75.96
	. 234 10	20.120 164.461		9.13	11.105			₩.02	0. VI Z	100.00I		0.44	1.872	100.001		15.47	24.04Z	100.00

10-403 Bulk copper, lead flotation in air, pyrite flotation in nitrogen with zinc reverse cleaning, and zinc flotation in air

	FL	0 T A	T I O N	C	0 N D I	T I	0 N S
sample reference	рН	activator added	Pre-cond1	tioning	collect type a	or Iddition	flotation 1 Qas
		g·t	gas	time	,,,	g/t	
bull rougher - 30 seconds A	9.1	-	air	20 min	SIFX/SAX	40	alr
bulk rougher - 30 seconds B	°.1	-	-	-	-	-	317
bulk rougher - 1 minute A	9.1	-	-	_	SIFY 'SAX	5	air
buik rougher - 1 sinute B	9.1	-	-	-	-	-	air
bulk rougher - 2 minutes A	9.1	-	•	-	SIPY /SAX	5	air
bulk rougher - 2 minutes B	°.1	-	-	-	SIP1/SA1	5	air
pvrite rougher - 2 minutes A	8.9	•	nitrogen	4 #10	SIPX/SAX	6	nitrogen
pyrite rougher - 2 minutes B	8.9	-	nitrogen	4 #13	SIPY/SAX	6	nitrogen
pyrite rougher - 2 minutes C	8. c	-	nitrogen	4 min	SIPX 'SAY	6	nitrogen
pyrite rougher - 2 minutes D	8.9	•	nitrogen	4 1 111	SIPY 'SAX	6	nitrogen
zinc rougher - 30 seconds A	10.2	6 30	none	10 min	SIFX 'SAX	25	aır
zinc rougher - 10 seconds B	10.2	-	-	-	-	_	air
zinc rougher - 1 minute A	10.2	-	-	-	-	-	air
zinc rougher - ! minute B	10.2	•	-	-	SIP1/SAX	5	alf
zinc rougher - I minutes A	10.2	-	-	-	-	-	air
zinc rougher - 2 minutes B	19.2	-	-	-	SIPY/SAX	5	ālf
zinc reverse flotation - 30 seconds	10.5	10e	air	5 81.	SIPX/SAX	S.	317
zinc reverse +lotation = 1 minute	10.5	-	-	-	-	-	air
zinc reverse flotation - 2 minutes	10.5	-	-	-	-	-	air
NOTE:			6FINDIN6	CONDITIO	NS:		
Soda ash used for pH adjustment during	j buli f	lotation	25 exnut	es. mild	steel char	ge, BM	eill .
Lime used for pH adjustment during bir	nc flota	tion	150 g't	sulpher	dio≈i de		

10-403 Bulk copper, lead flotation in air, pyrite flotation in nitrogen with zinc reverse cleaning, and zinc flotation in air

	SOLIDS IN	ISS BALAM	CE.		COFPER M	SS BALANC	E	t	EAD MASS B	AL MICE		21	INC MASS B	ALANCE		11	RON MASS B	MAKE	
	sample.	8455 P	ul I	сорр	er	copp	er	lea	4	lea	đ	2100	:	:10	c	110		110	M
	ee ight	sample	total	assay	units	recov		assay	units.	FEEDV	ery	assay	wasts	recov	ery	8558y	units	r ec gv	ery
Sample référence				(Z)		stage	tatel	(2)		st age	total	(2)		stage	total	(2)		stage	total
Bull rougher - 2 minutes (A)	97.00	4 862	4.861	2.62	0.127	32 471	32.472		1.320	34.532	34 532	5.48	●.276	3.241	3. 242	18.90	0.919	3.851	3.852
bull rougher - 2 minutes (B)	180.00	9.032	13.892	1.10	0.099	25 2°Z	57.762	14.10	1.273	33.102	67.632	10.50	0.748	11.122	14.372	25.20	2.275	9.532	13.301
bulk scazenger - 2 minutes (A)	27 00		15 751	0.60	0.008	7.072	57.831	8.46	0.117	3.052	70 681	10.70	0.145	1.791	16.072	26.30	0.356	1.492	14.957
Bulk scavenger - 2 minutes (B)	77.50	3 892	19.132	9 46	0.018	4.552	64.39%	4.70	0.183	4.751	75.431	7.15	0.356	4.171	20.241	32.20	1.252	5.242	20.122
bull scavenger - 2 minites (C)	154.70	7.77	24.901	0 27	0.021	5.342	69.731	2.59	0.201	5.232	80 662	7.04	0.547	6.422	26.46%	35.10	2.727	11.432	31.55%
Bulk sravenger - 2 minutes (D)	237.10	11.892	38.792	0.15	0.018	4.542	74.27%	1.57	0.187	4.851	85.511	5.72	0.680	7. 9 81	34.642	35.70	4.245	17.792	49.341
zanc rougher - tame 30 seronds	99.69	4 991	43,792	0.18	0.009	2.291	76.561		0.056	1.452	E6.971	39.70	1.783	23.271	57.912	11.20	0.559	2.342	51 697
zanc emigher time I minute	101 10	5 072	48.86%	0.22	0 911	2.84%	79.40%		0.066	1.731	68.6°Z	38.40	1.947	22.051	80.762	13.20	0.669	2.501	54 472
zzne rangher - time 2 minutes A	74 70	₹ 75 ፤	52.601	0.26	0.010	2.48 1	e: 001		0.061	1.697	90.292	29.20	1.094	12.842	43.401	18.20	0.482	2.561	57.342
zanr complex time 2 minutes &	37 20	1 872	54. 47 <u>2</u>	0.24	9.004	1.147	83.03Z		0.033	0.85X	91.151	11.70	0.218	2.562	74.16Z	24.80	0.463	1.942	
Zanc Travenger - time 30 seconds	77.59		55.60I	0.27	0.003	0.781	83.891	,	0.018	0.472	71.617	4.11	0.049	0.812	96.972	28.90	0.376	1.372	
Zinc stavenger - time I minute	23.70		56. 792	9.76	0 003	0.792			9.020	0.521	92.13Z	3.9L	9.046	●. 55 z	97.512	30.20	0.359	1.501	
final tails	861 70	41 212	100.001	0.14	0.061	15.412	100.90 1	€.70	0.303	7.872	100.001	0.47	0.212	2.497	100.00I	20.90	9.932	37.052	100.002
reinistituted fred	1994.00				0.393	100.00Z			3.844	100.001			8.521	100.002			23.864	100.001	
TEST SIMMARY:	sample	0 855	•		copper	recov	•		lead	recov			SINC	recov	•		110n	recov	
	weight	sample	total		assay I	stage	total		assay I	stage	total		assay I	st age	total		assay l	st age	total
holb (fotation	277.00	13.872	13.892		1.63	57.761	57.761	:	19.72	67.632	67.631		0.81	14.372	14.372		22.99	13.392	13.307
pyrate finiation	496 50	24, 902	30.79Z		0.74	16.517			2.76	17.88Z	05.512		6.94	20.272	34.642		34.46	35.938	49.343
zinc rougher flutation	312.60		54, 472		0.22	E. 751			1.38	5.632			33.44	41.52%			15.14	9.94Z	
Zinc Gravenger fintation	46.20		56.792		0.74	1.567			1.64	●.991			4.98	1.751			29.57	2.871	
final fintation tails	861.7♦	43.211	100.002		0.14	15.412	100.001	Į.	0.70	7.072	100.001		9 49	2.492	100.00Z		20.90	37.851	100.007

10-404 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.

(Mc Gill procedure)

		F L O	T 4 T I	0 N	C 0	N D I	T I O N
Sample reference	pH	activator	Pre-condi	tioning		ector	flotation
sample reverence		added g/t	gas	time	type	addition g/t	gas
bulk rougher - time 2 minutes	8.5	-	215	10 a in	EIF,	25	
bulk scavenger - time 2 minutes	8.5		-	- 10. min	SIFX	15	alr álr
pyrite rougher - time 2 minutes A	8.5	-	nitrogen	5 min	SIFX	6	nıtrogen
pyrite rougher - time 2 minutes B	8.5		nıtrogen	4 810	SIFX	6	nitrogen
pyrite rougher - time 2 minutes C	8.5	-	nitrogen	4 010	SIFX	6	nitrogen
pyrite rougher - time 2 minutes D	8.5	-	nitrogen		SIFT	6	nıtrogen
zinc rougher - time 30 seconds	10.5	1200	-	5 min	SIFi	50	alr
zinc rougher - time 1 minute	19.5	-	-	•	-	-	alf
zinc rougher - time 2 minutes	10.5	-	-	-	-	-	317
zinc rougher - time 2 minutes	10.5	-	-	-	-	-	air
zinc scavenger - time 0.5 minutes	10.5	-	-	-	SIPY	30	alr
zinc scavenger - time 1 minute	10.5	-	-	-	-	•	air
			SRINDING	CONDITION	45:		
			25 minut	es. oild	steel c	harge. BMS	•ili
			Tinc Sul	phate, 13	20 art		

10-404 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.

	SOLIDS N	iss balam	Œ		COFFER MA	SS BALANC	E	l	EAD MASS I	AL MICE		1	INC MASS &	PLANCE		1	40N MASS B	N. MEE	
	saeple	8455 B	oi i	C 009	er	¢qo ⊅	27	lea	đ	lei	ıd	210	τ	218	ĸ	iro	10	110	: 0
	₩21 ght	sample	total	9659A	enits	recov	er y	3553y	units.	FECOY	er y	25527	units	recov	er y	8558y	units	recov	PF y
sample reference	(gr ans)			(2)		stage	total	(2)		stage	total	(1)		stage	total	(2)		stage	tetal
bulk rougher 2 minutes	58 44	5 451	5.451	3.39	0.185	57 971	57.971	17.20	0.937	37.061	37.061	6 80	9.371	3,082	3.092	25 00	1 '63	5.558	5.55
bulk sravenger - 2 minutes	50.74	4.732	10 167	0.75	0 035	11,142	69. 112	7.34	0.442	17.48%	54.541	1.80	0.464	4.98I	B. *6Z	27.00	1 270	5.202	10.751
pyrite conc. In scavenger t	14.50	1.762	11.542	9.25	6 003	1.471	70. [7]	5.42	0.074	2.412	57.452	32.89	0.446	4.702	13.752	22.30	0.303	1.232	11.48
prite conc. In scavenger 2	0.09	9.067	11 542	0.64	9 000	0 00%	70.172	0.00	000	0.001	57,452	0.00	0.000	0.001	13.751	0.00	0.000	0.002	11.59
greate final conc (In scar tails)	410.49	39 281	47.82%	0.11	0.042	13 212	83.38X	1.50	0.574	22.791	89.152	4.97	1.902	29.432	34. 18Z	38.10	14.584	50.362	71.34
pyrite refu concentrate	424.98	70 64Z		●.11		14 20%		1.63		25.411		5.92		25.221		37.54		60.59I	
gans cougher tame 30 Seconds	30.22	7 921	52.641	0.18	0 905	1.5%	84.972	1.09	0.031	1.211	81.371	58.40	1.646	17. 281	51.862	7.14	0.258	1.052	72.35
gane rougher time I minute	30.72	2 871	55 502	0.20	u wif	1 502	66.77%	1.76	0.036	1.432	82 84I	49.70	1.410	15.142	67 002	9.71	0.278	1.132	73.5
zauc enquirer - tame 2 minutes	42.45	3.961	59.46Z	0.17	9 ang	2. 362	87.132	1.46	0.056	2.291	85.08 %	35.70	1.413	15.182	82 172	12.60	0.499	2.032	
gam scalenger - tame 30 seconds	20.30	1 842	61.367	9.29	0 004	1.197	90.322		0.033	1.291	86. 18%	31 40	9.595	6.782	89.561	16 70	●.32●	1.301	76.8
zinc scalenger - time t minute	16.74	1 511	62.871	0.21	ያሳብ ቀ	1.001	91.321	1.74	0.029	1.161	87.542	25.90	0.392	4.217	92.772	20.60	0.312	1.27%	78.1
garir scavenger - time 2 minutes	22.59	2.112	64.98 Z	0.15	0.003	0.991	72.311	1.03	0.039	1.521	8 9.061	12.00	0.253	2.721	95. 491	22.40	0.472	1.922	80.0
final tails	375.50	35 021	100 001	0.07	0.025	7.6°I	100.00I	0.79	0.277	10.941	100.001	1.20	0.420	4.512	100.002	14.00	4.003	19.96Z	100.0
reconstituted feed	1072.18				0.319	100.001			2.529	100.002			7.312	100.061			24.569	160.001	
TEST STIMMARY	sample	#255	pul l		Capper	recov	rery		Jead	recov	ery		ZINC	*eco	ery		1100	recov	/ 2 7 y
	we: ght	sa op l c	totai		2552y Z	stage	total		assay I	stage	total		assay Z	stage	total		assay I	st age	total
bult flatation	107.18	10.161	10.181		2 16	69.112	69. 111	ŀ	13.55	54.542	54 542		8.19	0.96Z	8.962		25.93	10.75%	19.7
p,rite flutation	424.98	1 362	11.547		0.25	1.072	70.171	L	5.42	2.912	57.45%		32.80	4.70%	13.751		22.39	1.232	11.9
zinc emigher flotation	103.39	47 921	59.461		0 13	10.961	87.137	l	1.46	27.632	85.08%		13.30	48,421	02.172		32.59	63.571	75.5
zinc cralenger flutation time 0 1.5 min	59.13	5 511	44.98Z		0.18	3.10%	92.31	Z	1.83	3.78%	8°. (%Z		22.48	13.312	95.49I		20.07	4.402	€ ∩.(
final flotation tails	375.50	35, 021	100 002		0.07	7.672	100.00	Z	0.79	10.941	100.00I		1.29	4.512	100.002		14.00	17.961	100.0

10-501 Bulk copper, lead flotation in air, pyrite flotation in argon and zinc flotation in air.

FLOTATION CONDITIONS

sample reference	ρH	activator added	Pre-con	ditioning	coll type	ector addition	flotation gas
		g/t	gas	time		g/t	•
bull rougher - time 2 minutes	8.5	-	air	10 min	SIPX	50	air
bulk scavenger - time 2 minutes	8.5	-	-	-	SIPX	30	air
pyrite rougher - time 2 minutes A	8.5	-	argon	2 min	SIPX	20	argon
pyrite rougher - time 2 minutes B	8.5	-	argon	2 min	SIPX	20	argon
pyrite rougher - time 2 minutes C	8.5	-	argon	2 ein	SIPX	5	argon
pyrite rougher - time 2 minutes D	8.5	-	argon	2 a ın	SIPX	5	argon
zinc rougher - time 30 seconds	10.0	1200	-	5 a in	SIPX	50	air
zinc rougher - time 1 minute	10.9	-	-	-	-	-	air
zinc rougher - time 2 minutes	10.0	-	-	-	-	-	air
zinc scavenger - time 0.5 minutes	10.0	-	•	-	SIPX	30	air
zinc scavenger - time ! minute	10.0	-	-	-	-	-	air
zinc reverse rougher - 1 minute	10.0	200	none	5 e in	SIPX	8	alf
zinc reverse scavenger - 1 minute	10.0	-	-	•	-	-	217

GRINDING CONDITIONS:

30 minutes, mild steel charge

Zinc Sulphate, 1200 g/t

10-501 Bulk copper, lead flotation in air, pyrite flotation in argon and zinc flotation in air.

11. APPENDIX 3; MATTABI ORE TESTWORK

EFFECT OF COLLECTOR AND SULPHITE ADDITION.

- 11-101 Bulk copper, lead flotation in air with 40 g/tonne collector and 3 kg/tonne sodium sulphite, the zinc flotation in air.
- 11-102 Bulk copper, lead flotation in air with 40 g/tonne collector and 1.5 kg/tonne sodium sulphite, then zinc flotation in air.
- 11-103 Bulk copper, lead flotation in air with 5 g/tonne collector and 1.5 kg/tonne sodium sulphite, then zinc flotation in air.
- 11-104 Bulk copper, lead flotation in air with 5 g/tonne collector and 1.0 kg/tonne sodium sulphite, then zinc flotation in air.

NITROGEN FLOTATION

- 11-201 Bulk copper, lead flotation in nitrogen with 5 g/tonne collector and 1.0 kg/tonne sodium sulphite, then zinc flotation in air.
- 11-202 Bulk copper, lead flotation in air with 5 g/tonne collector and 1.1 kg/tonne zinc sulphate, then zinc flotation in air.
- 11-203 Bulk copper, lead, pyrite flotation in air following conditioning with 5 g/tonne collector and 1.1 kg/tonne

zinc sulphate, then zinc flotation in air.

- 11-204 Bulk copper, lead, pyrite flotation in nitrogen following conditioning with 5 g/tonne collector and 1.1 kg/tonne zinc sulphate, then zinc flotation in nitrogen.
- 11-205 Bulk copper, lead, pyrite flotation in nitrogen following conditioning with 5 g/tonne collector and 1.1 kg/tonne zinc sulphate, then zinc flotation in air.

COLLECTORLESS FLOTATION

11-301 Collectorless flotation followed by zinc flotation.

	SOLIBS N	ASS BALAM	CE		COPPER M	ASS BALAN	Œ	U	EAD MASS D	ALANCE		21	INC MASS B	RLANCE		10	MOR MASS &	WWEE	
	500016	0055 p	ul]	cop	ter	(np j	er	lea	4	lea	d	2100		219	ť	110		178	69
	⊯erght	sample	totai	assay	units	reco	/Pf y	8558V	units	recev	er y	4558y	units	1 0COV	er y	a552y	mats	FECOV	v er y
sample reference				(2)		stage	total	(2)		stage	total	(2)		stage	total	(2)		stage	total
hulf ravaller - 30 seconds	114.66	11.582	11.581	2.44	0.304	46.102	44.10Z	1.40	0.195	27. 86 1	27.86%	3.80	0.440	4.232	4.232	47.63	4. 936	17.372	17.391
bulk sca enger - L minutes	59.51	6.017	17.597	4.12	0.246	37 342	83.44Z	3.34	0.202	20.921	54.792	7.70	0.433	4.162	8. 40Z	33.30	2.006	7.071	24.461
ult scarenger II - 2.5 minutes	29.21	2.851	20.441	1.60	0.046	4.871	70.312	3.35	0.075	13.67%	70.441	13.40	0.382	3.471	12.072	24.71	0.704	2.402	26.74
inc rougher - time 1 minute	107.75	11.082	31.522	0 12	0.013	7.011	•2.31Z	0.3i	0.034	4.921	75.302	55.22	6-121	50.911	70 7 82	7 63	0.844	2.982	29 92
anc raugher - tame 2 minutes A	44.71		36.061	0.70	0.013		94.232	0.67	0.030	4.351	79.731	46.45	2.116	20.361	91.35%	10 90	9.474	1.74%	31.64
anc rougher - time 2 minutes 8	9.09	0.00Z	36.962	0.09	9 900	0.001	94.232	P. 90	0.000	0.001	79.732	0.00	9.000	0.001	91.35%	0 00	0.000	6.001	31.6
ranc scalpinger - tabo 30 sprouds	24.94	7.572	.w. 581	0.30	9 000	1 141	55 37 2	1.76	0.047	7.072	86. 80Z	21.56	0.543	5.231	96.571	24.24	0.612	2.161	33.01
INC SCAVEIGN - TIMP T MINUTE	0.09	0.001	30.581	0.00	0.000	0.00Z	95.372	0.00	0.000	0.001	86.801	0.00	0.000	0.00Z	96,571	0.00	0.000	0.001	33 8
IRC SCALENGER - time 2 aimutes	0.00	0.00Z	30.501	0.00	0.000	0.001	95.372	0.00	0.000	0.001	86.80I	0.90	0.000	0.001	96.571	0.00	0.000	9.00Z	33 8
limat tails	698.20	61.47Z	100.00I	9.05	0.03t	4.63%	100.001	0.15	0.072	13.202	100.002	9.50	0.354	3.431	100,001	30.57	10.707	M.191	100.0
reconstituted feed	770.17				0.663 -	100.001			0.498	100.001			10.391	100.001			20.300	100.001	ı
IEST SUMMARY:	sample	0055	ui i		Copper	recov	er y		lead	recou	rery		SINC	recev	er y		tron	reces	-
	wes ght	sample	total		assay I	st age	total		assay 1	stage	total		assay I	stage	total		assay Z	stage	teta
bulk flot stron	207.38	20.442	29.442		2.93	70.312	90.312		2.41	70.461	70.461		6.14	12.071	17.07%		37.41	26 992	26.5
zinc roweler flotation	154.67	15.67	36 06I		0.17	3.921	94.232		0.41	9.272	79.731		52.73	79.271			0.58	4,72%	
zinc scalenger flotation time 8-1.5 min	24.44		30.502		●.30	1.14%			1.76	7.072			21.56	5.232			24.29	2. 161	
final flutation tails	608.Z0	61.422	100.00Z		0.05	4.632	199.001		0.15	13. 20Z	100.00I		1.58	3, 432	199,001		30.57	56.171	Z 100.0

11-101 Bulk copper, lead flotation in air with 40 g/tonne collector and 3kg/tonne sodium sulphite, then zinc flotation in air.

FLOTATION CONDITIONS

	ρН	activator	Pre-con	ditioning	colle	ector	flotation
sample reference		added g/t	gas	time	type	addition g/t	gas
bulk rougher - 30 seconds	9.2	-	31r	3 a in	plant#	50	alf
bulk scavenger - 1 sinute	9.2	-	•	-	-	-	air
bulk scavenger - 2.5 minutes	9.2	-	-	-	-	•	alf
zinc rougher - 1 minute	10.5	1100	none	5 ain	YEX	30	air
zinc rougher - 2 minutes	10.5	-	-	-	-	-	alr
zinc scavenger - 1 minute	10.5	-	-	-	KEX	20	air

GRINDING CONDITIONS:

20 minutes, mild steel charge

2000 g/t sodium sulphite

NOTE:

Plant xanthate contains 50% Ethyl manthate and

50% dithiophosphate (Cyanamid 241)

11-101 Bulk copper, lead flotation in air with 40 g/tonne collector and 3kg/tonne sodium sulphite, then zinc flotation in air.

	50L185 N	NSS BALAN	CE		COPPER N	ISS MLAN	Œ	ı	EAD MASS I	MLANCE		1	INC MASS I	MLANCE		1	IRON HASS B	ALMEE	
	samole	8355 \$		cop		C 001		lea		ie		218	c	215	ı c	170	m	174	**
sample reference	uesght (grans)	sample	total	assay (Z)	waits	recei stage	•	essay	unit5	recon stage	rer y total	853y (2)	wasts	recov stage	erv total	455 4 y (1)	enits	stage recen	ver v total
bull rougher 30 seconds	294.93	29.331	29,332	1.03	0.302	46.10Z	46.101	1.00	0.293	41.872	41.072	4.40	1.291	12 662	12.461	30.03	11.389	39.751	39,751
bull scarenger - I aimutes	179.95	17.902		1.31	0.234	35.772	01.00X	1.36	0.243	34.742	76 612	4.66	0.834	0.162	20.842	39.30	7.033	24.541	
bulk scarenger 11 2.5 acoutes	48.87	4.861	52.092	1.57	0.076	11.642	73.522	1.73	0.984	12.001	88.417	6.30	0.306	3. 00 I	23.842	37.97	1.845		70.731
zinc ringher tree 1 arnute	113.00	11.247	63.322	0.13	0.015	2.232	95.75Z	9.16	0.018	2.571	91 18Z	55.82	4.273	61.532	65.372	7.55	0.840	2 961	73.491
Zinc rougher - time 2 cinutes &	28.19	2.00Z	66.132	0.32	0.00*	1.372	77.12%	0.44	0.013	1.841	93.021	44.34	1.243	12.193	97.56Z	11.24	0.315	1.107	
zinc emglier tipe 2 ainutes B	0 90	0.001	66.137	0.00	9.000	9.002	97.121	0.00	0.000	0.00I	93.02I	0 04	6 300	A AAT	.56	v 10	6 000	0 (-12	74.79
Zinc Stavenger - fine 30 seconds	48 79	4.85%	70 981	0.15	0 007	1 112	98.231	●. 29	0.014	2 01T	95.032	3.21	0.156	1.532	90.092	37 75	1.932	* ***	81.19
ZIRC Scavenger - time & minute	0 00	9 00Z	70 48 %	0 00	0 000	0.091	90.232	₿ ûn	0.000	9 90I	95.03Z	0.00	0.000	0.001	99.041	0.00	0.000	0.602	91.16
zinc scavenger - time 2 minutes	0.00	9 00Z	70.981	0.00	9.000	0.001	98.231	0.00	U. 444	3.55%	95.032	0.00	0.000	0.001	99.092	9.00	0.000	9.001	81.18
final faits	291.00	29.021	100.001	0.94	0.012	1.77%	100.001	0.12	0.035	4.972	100.672	0.22	0.073	0.91Z	100.00Z	18.58	5.392	19, 821	100.00
reconstituted feed	1405.53				●.455	100.00I			0.701	100.00Z			10.195	100.001			29.455	100.001	
TEST SIMMART:	sa o ple	8455	pul !		copper	recov	er y		lead	recov	er y		21 MC	recov	er y		17.00	/ eco	~ 97 y
	weight	sample	total		assay l	stage	total		assay l	stage	total		assay 1	stage	total		1 yezes	stage	total
bulk fintation	\$23.75	52.97Z	52.9*1		1,18	93.522	93.52%		1.17	80.612	88.412		4.67	23.84%	73.841		30. 91	70.732	70.73
Zinc emigher fletation	141.19	14.042	66.132		0.17	3.601	97.121		●.22	4.411	93.021		53.53	73.721	97.56Z		8.29	4.061	74.7
zinc scavenger flotation	48,79	4.052	79.982		0.15	1.117	98. 23X		0.29	7.011	95.031		3.21	1.532	99.09%		37.75	4. 392	01.10
final fintation tails	291.00	29.022	100.00Z		9.04	1.771	100.00I		0.12	4.972	100.00I		♦.32	0.91Z	100.00T		18.50	18.821	100.00

11-102 Bulk copper, lead flotation in air with 40 g/tonne collector and 1.5kg/tonne sodium sulphite, then zinc flotation in air

	- 49		_				-
sample reference	рн а	added	Pre-con	ditioning	colle type	ector addition	flotation
		g/t	gas	tise	cype	g/t	gas
bulk rougher - 30 seconds	9.5	-	air	3 a in	planti	46	air
bulk scavenger – i minute	9.5	•	•	-	-	-	alf
bull scavenger - 2.5 minutes	9.5	-	-	-	-	•	air
zinz rougher - 1 minute	.5	1100	none	5 s in	KEX	30	atr
zinc rougher - 2 minutes	10.5	-	-	•	-	-	air
zins scavenger – 2 minutes	10.5	-	-	-	KEX	29	air
			6RINDIN	E CONDITIO	NS:		
			20 min	utes. mild	steel c	harge	
			1600 g	/t sodium	sulphite		
	N	OTE:					
	P)	lant xanti	hate cont	ains 50% E	thyl ×an	thate and	
		507 dithi	nnhacabət	e (Cyana s i	d 24!)		

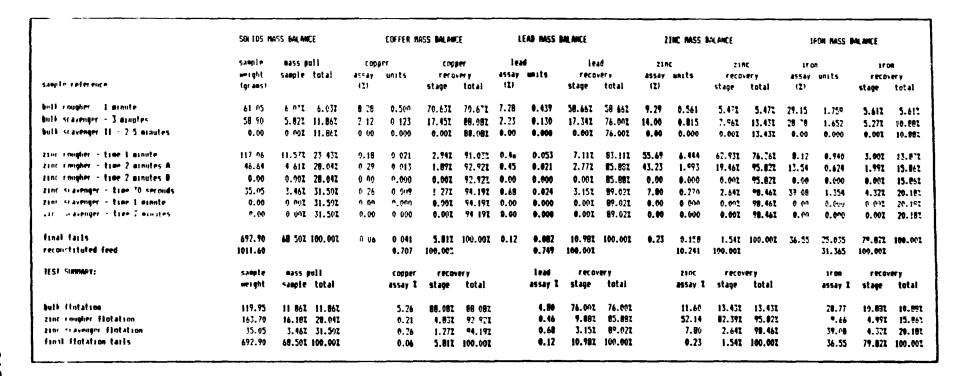
11-102 Bulk copper, lead flotation in air with 40 g/tonne collector and 1.5kg/tonne sodium sulphite, then zinc flotation in air

	SOLIDS N	455 PALAN	CE		EUFPER IN	155 BALANC	E	Ĩ	EAD MASS I	MLMICE		1	INC MASS B	al amee		1	PON NASS I	AL MICE	
	130010	0355	ull	cop) PT	сорд	er	100	M	lea	ıd	218	c	210	C	iro	•	111	•
	wes glit	sample	total	85584	units	rece	PF Y	assay	un i ts	recov	er y	assay	muts	recov	er y	8558Y	units	reco	v e c y
sample reference	(grans)			(2)		stage	'otal	(2)		stage	total	(1)		stage	total	(2)		stage	total
hist rougher 1 minute	25 43	7 47%	2 421	7.00	0 218	31.421	31.42%	B 58	0.209	27.251	27 252	9.57	0.731	2.30%	2.301	26.99	0 653	7.152	2.152
bull sravenger - 3 minutes	40 37	3 84I	6.262	7.04	0.270	39.022	70 44Z	5.66	0.217	20.541	55 791	12.92	0.4%	4,932	7.24%	25.99	9 638	3.792	
bull scarenger II - I minutes	0.00	0.001	4.242	0.00	0.000	0.00E	70.441	0 00	0.000	0.00Z	55 79%	6.00	0.000	0.00I	7.242	0.00	0.000	0.00I	5.432
zin' rougher - time minute	107.16	10.19Z	16.45Z	0.38	0.039	5.592	76 93Z	1.04	9.106	13.922	69.712	52 87	5.391	53 621	60.862	1.64	0.883	2.911	0.342
zinc impher - time 2 alautes A	46.52	4.422	70. 981	0.53	9.023	3 382	77.412	0.83	0.037	4.02%	74.54%	47.91	2.120	21 092	81.942	11.42	0.505	1.661	10.001
zinc reigher - time 2 minutes D	0.00	0.00I	20. 06 %	0.00	0.000	0.001	79.412	0.00	0.000	0.00I	74.54Z	0.00	0.000	0.002	81.941	0.00	0.000	0 00Z	10.002
zant stavenger - time 30 seconds	71.65	6 82Z	27.692	0.82	0.056	8.071	87.482	1.36	0.073	12.17%	86 71Z	23.24	1.584	15.75%	97.70%	26.71	1.020	5.992	
zanc scavenger - time minute	0.00	0.001	27.69%	0.00	0.000	0.001	e7.48%	0.00	0.000	0.001	86.711	0.00	0.000	0.001	97.701	0.00	0.000	0.00%	16.002
zinc sravenger - time 2 minutes	0.00	9.001	27.691	0.00	9.000	9.001	87.481	0.00	0.000	0.002	86.712	0.00	0.000	0.001	97.702	0.00	0.000	0.002	16.00Z
final tails	760.20	72.312	100.00Z	♦.12	0 097	12 521	100.001	0.14	0.101	13.29%	100.001	0.32	6.231	2.301	100.001	35.29	25.518	84.001	100.00I
reconstituted feed	1051.33				0.493	100.001			0.762	100.00I			10.054	100.001			30.377	100.00I	
TEST SIMMARY:	samle	0255	eul l		CDBPET		ery		lead	recev	ery		2100	recov	er y		11 DR	1900	very
	weight	sample	total		assay I	stage	total		assay l	stage	total		assay I	stage	total		assay 1	stage	total
bulk flotation	65.80	6 761	6.267		7.80	70.442	76.442		6.79	55.791	55.791		11.63	7.242	7.242		26.30	5.432	5 431
zinc rougher flotation	153.68	14.621	20.082		0.43	0.972	79.412		0.78	10.74%			51.38	74.71%			9 50	4.572	
ZINC SCAVENGER Flotation	71.45	6.871	27.692		0.82	3.07%	87.48%		1.34	12.172			23.24		97.70%		26.71	5.992	
final lintation tails	760.20	72.311	100.00Z		0.12	12.521	100.00Z		0, 14	13.291	100.001		●.32	2.301	100.00Z		35.2 9	84. 9 0%	100.001

11-103 Bulk copper,lead flotation in air using 5 g/tonne collector and 1.5kg/tonne sulphite depression of pyrite, then zinc flotation in air.

	FL	O T A	T] 0	N C	0 N D	I I I	0 N S
ample reference	pH	added		nditioning	coll type	ector addition	flotation gas
		g/t	gas	tise		g/t	
ulk rougher - 1 minute	9.5	-	air	3 min	plant#	5	air
ulk scavenger - 3 minutes	9.5	-	-	-	-	-	317
inc rougher - 1 minute	10.5	1100	none	5 e in	KEX	40	air
inc rougher - 2 minutes	10.5	-	-	-	•	-	air
inc scavenger - 2 minutes	10.5	-	-	-	KEX	20	alr
			GRINDII	V6 CONDITIO	NS:		
			20 11	nutes, øild	steel c	harge	
			1600 (/t sodium	sulphite		
	N	OTE:					
	ρ	lant xanti	hate conf	tains 50% E	thyl xan	thate and	
		507 4.46.	anhaanh s	te (Cyanamı	4 2411		

11-103 Bulk copper, lead flotation in air using 5 g/tonne collector and 1.5kg/tonne sulphite depression of pyrite, then zinc flotation in air.



11-104 Bulk copper, lead flotation in air using 5 g/tonne collector and 1.0kg/tonne sulphite depression of pyrite, then zinc flotation in air.

FLOTATION

GRINDING CONDITIONS:

20 minutes, mild steel charge

C O N D I T I O N S

1000 g/t sodium sulphite

NOTE:

Plant xanthate contains 50% Ethyl xanthate and

50% dithiophosphate (Cyanamid 241)

11-104 Bulk copper, lead flotation in air using 5 g/tonne collector and 1.0kg/tonne sulphite depression of pyrite, then zinc flotation in air.

	SOLIDS W	ASS BALAM	CE		COPPER N	ISS BALANC	Ē	t	EAD MASS (ALMICE		1	INC MASS I	AL MICE		1	RON MASS 1	ALMICE	
	sample	0455 p	ull	cop	per	copy	er	lea	4	lea	ıd	ZIR	c	218	c	170	10	170	08
	weight	sample	tetal	assay	units	recov	er y	assay	units	recev	rery	assay	units	recov	er y	4553Y	unsts	reco	YEFY
Sample reference	(graes)			(2)		stage	total	(1)		stage	total	(2)		stage	total	(1)		stage	total
bulk reigher Lasnufe	97.71	9.721	9.72I	2.64	0.257	37.072	29.072	2.44	0.257	42.522	42.52%	5.39	0.524	5.117	5.112	38.09	3.703	11.94Z	11.903
bulk scavenger - 3 nautes	A1 30	5117	15 832	1.50	0.096	14.692	53.762	1.50	0.094	15.782	59.502	7.09	0.433	4.232	9. 34Z	37.02	2.310	7.451	19.392
bulk scavenger 11 - 3 minutes	9.00	9.001	15.832	0.00	0.000	0.00I	53.762	0.00	0.000	0.001	50.501	0.00	9.000	0.001	7.341	0.00	0,000	0.001	17.392
zinc rougher - tion 1 ainute	52.70	5 252	21.08I	0.44	0.049	7.512	61.272	0.94	0.047	0. 18Z	66.481	47.61	7.405	25.431	34.772	9.34	0.470	1.502	20.972
zanc resigher - tame 2 minutes &	25.36	2.922	24.002	1.02	0.030	4.542	65.812	1.02	0.030	4.942	71.612	43.53	1.272	12.412	47.182	11.10	0.324	1.051	22.021
zanc rougher - take 2 minutes B	0.00	0.002	24.00%	0.00	0.000	0.00Z	45.812	0.00	0.000	0.001	71.612	0.00	0.000	0.001	47.182	0.90	0.000	0.001	22.021
zinc sravenger - fine 2 minutes	148.43	14.771	38.771	0 42	0.062	7.442	75.251	0.42	0 062	10.28Z	81.89%	34.27	5.061	47.4CI	°6.591	22.70	3.352	10.812	32.039
ZINC SCAVENGER - time 2 minutes	20.36	2 032	40.792	0.72	0.015	2.721	77.47%	0.72	0.015	2.421	04.31Z	7.32	0.148	1.451	79.031	34.04	0.730	2.361	35.191
zinc scazenger - time 2 minutes	0.00	0.002	40.771	0.00	e. 000	0.007	77.471	0.00	0.000	0.001	g. 31I	0.00	0,000	0.001	78.031	0.00	0.000	9.061	35.197
final fails	595.10	57. 212	100.00I	♦.25	9.148	22.531	100,001	0.16	0.075	15.672	100.00Z	0.34	0. 201	1.971	100.001	33.94	20.075	64.613	100.00
reconstituted feed	1005.12				0.657	100.001			0.404	100.007			10.244	100.001			31.005	100.001	
TEST SHIMARY:	sample	8255 (ol 1		copper	recov	er y		lead	r ecev	ery		ZINC	recov	ery		1700	reco	v er y
	wes ght	sample	total		assay l	st age	total		assay I	stage	total		assay 1	stage	total		assay I	stage	total
bulk finiation	157.07	15.032	15.83Z		2.23	53.761	53.742		2.23	50.501	50.50L		4.05	7.341	7.341		37.99	14.391	17.37
zinc rougher flotation	82.14	6.172	24.00Z		€.97	12.051	65. 8 12		0.97	13.117	71.41Z		47.44	37.842	47.18Z		9.97	2.432	22.02
ging sia enger flotation	168.79	16.79%	49.792		0.46	11.66Z	77.47%		0.46	12.692	84.311		31.62	50.851	78.032		24.31	13.172	35.19
final flotation tails	595.10	59.212	100.001		0.25	22.531	100.00I		9.16	15.492	100.00Z		0.34	1.971	100.00Z		33,74	44.817	100.00

11-201 Bulk copper, lead flotation in nitrogen with 5 g/t collector and 1.0 kg/tonne sulphite depression of pyrite, then zinc flotation in air.



Sample reference	pH a	ectivator added	Pre-cond:	tioning	coll: type	ector addition	flotation
		g/t	gas	time	cype	g/t	gas
bulk rougher ~ 1 minute	9.5	-	nitrogen	20 min	plant#	5	nıtrogen
bulk scavenger - 3 minutes	9.5	•	•	-	•	-	nitrogen
zinc rougher - ! minute	10.5	1190	none	5 ein	KEX	35	alr
zinc rougher - 2 minutes	10.5	-	-	-	•	•	alf
zinc scavenger - 2 minutes A	10.5	-	-	-	KEX	20	217
zinc scavenger - 2 minutes 8	10.5	-	-	-	-	-	air
			6RINDIN6	CONDITIO	NS:		,
			29 minut	tes. oild	steel c	harge	
			1000 g/(sodiu s	sulphite	ı	
	W	OTE:					
	P:	lant manti	hate contai	ins 50% E	thyl xan	thate and	
	í	50% dithii	ophosphate	(Cyanaes	4 2811		

11-201 Bulk copper, lead flotation in nitrogen with 5 g/t collector and 1.0 kg/tonne sulphite depression of pyrite, then zinc flotation in air.

AA	SOLIDS IN	ISS BALANC	.E		EUTPER IM	ISS BALANC	E	i	EAD IMSS I	MLMEE		2	INC MASS B	AL MICE		1	ROW MASS B	AL ANCE	
test number 77	sample	8455 Pt	111	cop	jer	Capp	er	lea	ıd	lea	ıd	219	t	210	ĸ	100		170	m
sample reference	weight (graes)	sample	total	4558Y	units	reco	er y total	assay (Z)	units	recon stage	total	2553Y (Z)	m ite	recar stage	ery total	assay (I)	emits	recev stage	rery total
built rangker Sinate	143 95	17 952	13,951	2,39	0 334	40.771	48.772	1.48	0.207	27.812	27 811	3 62	9.421	4.172	4.17Z	41.78	5 830	19.54E	19.542
buth scarenger - 3 minutes	152.47	14 802	28.74%	1.35	9 700	29 197	77 712	1 62	0 740	32.291	60.10Z	4.71	0.497	4. 90Z	11.072	40.39	5.978	20.031	39.572
bulk scorenger 11 - 3 minutes	0.00	0.002	28 762	0 00	0 000	0.001	77.911	0.00	0.000	0.00I	40.101	0.00	0.000	0.001	11.07%	0.00	0.000	0.001	39.572
rior insigher time 2 minutes A	115.20	11 17%	39.922	0.80	0.987	13.052	90.96Z	1.65	0.184	24.821	84. 472	28.92	3.230	31,97%	43.04I	21.46	2.732	7. 15I	40.73
zanc rinigher time 2 minutes 9	72.19	7.001	46.922	0.25	0.01?	2.562	73.521	0.42	0.027	3. 96Z	66.66I	30.28	2.119	20.781	64.022	22.79	1.595	5.35%	54.07
zinc emiglier - time 2 minutes C	9.90	0.001	44.92%	0.00	0 000	0.00Z	93.522	0.00	0.000	0.001	89. 881	1.00	0.000	0.001	64.022	0.00	0.000	0.001	54.07
zinc scarenger - time Z minutes à	172.00	18.672		0.10	0 017	2.73%	76.252	0.17	0.036	4.78%	73.66%	16.16	3.020	29.902	93.922	35.11	6.562	21.091	74.07
zinc starpiger - fine 2 minutes 0	92.26	7.97%	73.592	0.09	0.007	1.052	97.302	0.14	0.013	1.721	95.381	5.95	0.474	4.702	78.61Z	38.74	3.089	19.351	86.42
zanc scavenger - time 2 minutes C	0.00	0.002	73.592	0.00	0.000	0.001	97.30E	0.00	0.000	9.00I	95.381	0.00	9.000	0.001	98.412	0.00	0.000	0.00I	86.42
final tools	272.46	26.417	100.00I	0.07	0.018	2.701	190.00Z	0.13	0.034	4.621	100.001	0.53	0.140	1.392	100.001	15.34	4.052	13.591	100.00
reconstituted feed	1031.53				9.485	100.00I			0.743	100.002			10.107	100.00Z			29.838	100.00I	
TEST SIMMARY:	sample	8955	oul 1		copper	recov	егу		lead	FECEN	erγ		SINC	recov	ery		1706	recov	rer y
	weight	samle	total		assay l	st age	total		assay I	stage	total		assay I	si age	total		assay I	stage	total
bult finiation	2%.62	28.76%	20.762		1.85	77.912	77.912		1.55	40.102	60.10Z		3.89	11.072	11.07%		41.06	39.571	39.57
zene raugher flatation	187.39	19.172	46.92I		9.59	15.412	93.521		1.10	28.77%			27.44	52.95%	64.02%		23.87	14.501	54 07
Zinc Stavenger (totation	275.06	26.672	73.592		0.10	3.7 8 Z			0.18	6.50I			13.11		70.412		36.20	32.352	86 42
final fintation tasks	272.46	26.412	100.001		0.07	2.701	100.00Z		0.13	4.62%	100.001		0.53	1.397	100.001		15.34	13.501	100.00

11-202 Bulk copper, lead, pyrite flotation in air following conditioning in nitrogen with 5 g/t collector and zinc sulphate depression of sphalerite. Zinc flotation in air.

F., .	activator added	Pre-condi	tioning	colle type	ector addition	flotatio gas
	g/t	gas	time	.,,,,	g/t	yas
9.5	-	nitrogen	15 sec	plant#	5	air
9.5	-	-	-	•	-	air
10.5	1100	nene	5 min	KEX	35	a1 <i>1</i>
10.5	-	-	-	-	-	air
10.5	•	-	-	KEX	20	air
10.5	•	-	•	-	-	21"
		SPINDING	CONDITIO	MS:		
		20 minut	es, eild	i steel c	harge	
		1100 g/t	ZINC SU	ılphate		
N	OTE:					
P	lant xanti	hate contai	.ns 5 0% e	ethyl xan	thate and	
	9.5 10.5 10.5 10.5	9.5 ~ 9.5 ~ 10.5 1100 10.5 ~ 10.5 ~ 10.5 ~	9/t gas 9.5 - nitrogen 9.5 10.5 1100 nene 10.5 10.5 5PINDING 20 minut 1100 g/t	9.5 - nitrogen 15 sec 9.5	9.5 - nitrogen 15 sec plant8 9.5	9.5 - nitrogen 15 sec plant* 5 9.5 10.5 1100 none 5 min KEY 35 10.5 KEX 20 10.5 SRINDING CONDITIONS: 20 minutes, mild steel charge 1100 g/t zinc sulphate

11-202 Bulk copper,lead, pyrite flotation in air following conditioning in nitrogen with 5 g/t collector and zinc sulphate depression of sphalerite. Zinc flotation in air.

	SOLIDS IN	ASS BALAH	Œ		COPPER N	ASS BALANC	Œ	L	EAD MASS B	ALANCE		21	INC MASS B	RLANCE		1	ROW MASS IN	NAMEE	
	sample	0455 P	wll	copi	per	copy	jer	lea	4	lea	d	2100		210	c	Iro	m	110	36
	wei ght	sample	t ot al	3 55 3 y	units	reco	ver y	85589	mits	recov	er y	assay	mits	FECOV	ery	25527	mits	recov	rêt y
sample reference	(graes)			(2)		stage	total	(Z)		stage	total	(2)		stage	tatal	(2)		stage	total
bull rougher - I minute	213 30	70.96Z	20.961	2.20	0.461	65.772	65 771	1.67	0.354	46.037	46.831	3.40	0.755	7.321	7.322	41.38	8.674	20.222	28.221
bull scavenger - 3 ainutes	48.64	4.78%	25.74Z	2 69	0.128	18 27%	84 04Z	2.52	0 120	15.921	62.751	7.88	0.472	4.581	11.901	35.15	1.680	5.472	33.692
bull scavenger [] - 3 minutes	0.00	0.001	25.741	0.00	0.000	0.001	84.04I	9.00	0.000	0.001	42.751	0.00	9.000	0.001	11.701	0.00	0.000	0.001	33.497
zanc rougher - tame 3 minute	133.33	13.102	30.642	0.45	0.059	8.412	77. 45X	1.30	0.181	23.902	B6.652	41.04	5.377	52.141	64.041	16.97	2.224	7.231	40.72
zinc rougher - time 2 minutes A	53.71	5.281	44.122	0.21	0.011	1 591	94.032	0.41	0.022	2.86%	87.511	36.26	1.714	18.561	82.40%	19.00	1.003	3.761	44.19
zinc rougher time 2 minutes B	0.00	0.002	44.122	0.00	0.000	0.00Z	94.03Z	0.00	0.000	0.002	07.511	0.00	9.000	0.001	82.602	0.00	0.000	0.001	44.19
einc scavenger - time 2 minutes A	67.85	4.19X	50.301	6.09	0.004		94 821	0.22	0.014	1.89Z	71.30%	21.20	1.309	12.791	95.292	30.37	1.076	6.10I	50.29
finc scavenger - time 2 minutes B	31.25		53 37%	0.12	0 004	0 537		0.32	0.010	1.301	92.602	10.95	0.334	3.261	98.551	34.23	1.051	3.422	53.71
zinc schenger - time 2 minutes C	●.00		53.371	0.00	9.000	0.00Z	-	0.00	0.000	0.00I	92.60Z	9.00	9.009	0.001	78.55 Z	0.00	0.900	0.001	53.71
final tarts	474.50	46.432	190.00Z	0.07	0.033	4.66Z	190.00I	0.12	0.056	7.402	100. 00 I	0.32	0.14*	1.452	100.001	30.51	14.227	44.202	100.00
reconstituted feed	1017.58				0.701	100.00I			€.757	100.001			10.313	100.001			30.734	100.001	
1EST SUMMARY:	saopi e	8855 (pol t		copper	FECU	ery		lead	reces	rery		2180	recev	er y		1700	recov	very
	weight	saaple	total		assay 2	stage	total		assay 1	stage	total		assay I	stage	total		assay l	stage	total
bulk flutstron	261.94	25.74%	25.741		2.29	84.04Z	84.042		1.84	62.751	62.751		4.77	11.702	11.901		40.22	33.492	33.6
zinc rougher flotation	187.04	10.381	44.12%		6.30	9.002	74.032		1.10	26.761	87.511		39.47	70.70Z	82. Ant		17.55	10.501	
zinc scavenger flotation	94.10	9.257	53 37Z		0.10	1.322	95.34Z		0.75	3. 10 1			17.80	15.761			31.65	7.521	
final flotation tails	474.50	46.631	100.00I		0.07	4.661	100.00I		€. 12	7.40I	100.00Z		0.32	1.451	100.00Z		30.51	46.242	100.0

11-203 Bulk copper, lead, pyrite flotation in air following conditioning with 5 g/t collector and zinc sulphate depression of sphalerite. Zinc flotation in air.

	FL	O T A	1 0	N C	0 N D	1 7 1	0 N S
sample reference	pH		Pre-con	ditioning		ector	flotation
pampie reference		added g/t	gas	time	type	addition g/t	gas
bulk rougher - 1 minute	9.5	-	alr	3 e in	plant#	5	air
bulk scavenger - 3 minutes	9.5	-	-	-	-	-	air
zinc rougher - ! minute	10.5	1100	1016	5 min	KEX	35	alr
zinc rougher - 2 minutes	10.5	-	•	-	-	-	alr
zinc scavenger - 2 minutes A	10.5	-	-	-	KEX	20	alf
zinc scavenger - 2 minutes B	10.5	-	-	-	-	•	air
			6PINDIN	6 CONDITIO	INS:		
			20 ein	utes. mild	stee ¹ c	harge	
			1100 g	/t zinc su	ilphate		
	N	10TE:					
	P	lant xant	hate cont	ains 50% e	ethyl xan	thate and	
		50% dithi					

11-203 Bulk copper,lead, pyrite flotation in air following conditioning with 5 g/t collector and zinc sulphate depression of sphalerite. Zinc flotation in air.

	SOLIDS IN	ASS BALAN	CE		COFFER IN	ISS BALANC	£	1	EAD MASS I	DAL MICE		2	INC MASS I	MLANCE		I	RON MASS (MLANCE	
	Sample	0355 P	uli	CODI	per	C 00 J	187	le:	ıd	lea	ad	218	ĸ	210	ıc	110	a	111	04
	weight	sample	total	8558 ¥	enits	FECEN	er y		maits	F 00 01	ver y	2553Y	units	reco	er y	3552 y	units	reco	ver y
sample reference	(graes)			(1)		stage	total	(1)		stage	total	(2)		stage	total	(1)		stage	tetal
bull rougher 1 minute	463.30	42.421	42.421	0.45	0.171	20.571	20.571	●.56	9.239	34.44Z	34,442	1.61	0.483	4.772	6.72%	45.22	19.101	64.632	44.432
bolb scavenger - 3 ainutes	150 36	13.772	56. 1BZ	1.18	0.162	24.32 Z	52.8°Z	1.34	0.184	26.751	41.191	3.83	0.527	5. 192	11.912	41.23	5.676	19.132	83.762
bult rravenger 11 - 2.5 minutes	9.00	9.00I	54.182	0.08	0.000	0.00I	52.091	●.00	0.000	0.002	41.192	0.00	0.000	9.961	11.912	9.00	0.000	0.001	83.762
zinc cougher - time 1 minute	78.47	7.031	65.227	2.45	0.739	35.833	88.72%	2.23	0.201	29.217	90.40Z	37.28	3.368	33.142	45.051	15.12	1.366	4.602	00.34 2
231H rougher - time 2 minites A	27.64	7.711	67.932	0.72	0.020	2.921	91.651	0.54	0.015	2.12X	92.532	43.13	1.170	11.52%	56.571	11.47	0.317	1.071	87. 43Z
zinc cougher - time 7 minutes B	0.90	0.00Z	67.931	0.00	0.000	9.00I	91.451	0.00	0.000	0.002	92.531	0.00	0.000	0.00Z	54.571	0.00	0.000	0.001	87.432
zinc scavenger - time 2 minutes a	72 32	6.621	74.552	0.42	0.028	4.16Z	75.811	0.24	0 016	2 102	24.832	50.58	3.347	32.962	89.53Z	9.14	0.405	2.042	91.47%
2101 4-Pager - time 2 minutes b	0 22	0.001	74.55%	0 00	0 000	0.001	95 817		0.900	0.001	94 831	0.00	0.000	0.001	87.532	0.00	9.000	001	91 472
zinc stavenger - time 2 minutes C	9.00	9.001	74.551	0.00	0.000	0.001	95. 81 Z	●.00	0.000	9.00Z	94 831	0.00	0.000	0.001	89.532	0.00	0.000	0.001	91.472
finat tails	211.47	25.451	100.001	0.11	0.028	4.192	100.00I	0.14	0.034	5.172	100,001	4.18	1.064	10.471	100.00I	9.95	2.532	6.532	100.001
reconstituted feed	1097.26				0.468	100.60L			0.490	100.00Z			10.141	100.00Z			29.474	100.002	
1651 SHMMARY:	sample	B 855	pull		Copper	FPCOV	er y		lead	reco	rer y		zinc	recov	er y		1708	rece	very
	weight	samle	total		assay I	stage	total		assay I	stage	total		assay I	stage	total		assay I	stage	total
hilb flitatatenn	613.66	56. 182	56. 18Z		0.43	52. 89 %	52.892		0.75	61.192	61.172		7.15	11.71%	11.712		41.24	83.762	83.741
zanc rougher flotation	129. 31	11.752	67.932		2.20	38.761	91.651		1.84	31.34Z			38.63	44.46Z	56.572		14.32	5.471	87. 432
zinc scavenger fintation	72.32	6.627	74.55Z		0.42	4.162	95.817		0.24	2.30Z			50.58	32.961	87.5 31		7.14	7.04Z	71.472
final flotation tails	277.9?	25.451	100.001		0.11	4.171	100.007		0.14	5.172	100.00Z		4.18	10,472	100.00I		9.95	8.532	100.001

11-204 Bulk copper,lead,pyrite flotation in nitrogen following nitrogen conditioning, 5 g/t collector and zinc sulphate depression. Zinc flotation in nitrogen.

sample reference	pH a	ictivator added	Pre-condi	tioning	coll type	ector addition	flotatio qas
		g/t	gas	tiee	.,,,	g/t	943
oulk rougher - 1 minute	9.5	_	nitrogen	20 min	-	-	nıtrogen
bulk scavenger + 3 minutes	9.5	-	-	-	-	•	nitrogen
nnc rougher - 1 minute	10.5	1100	none	5 018	KEY	35	air
zinc rougher - 2 minutes	10.5	-	-	-	-	-	air
zinc scavenger – 2 minutes	10.5	-	-	-	KEX	20	air
			6P IND ING	CONDITIO	NS:		
			20 minut	tes, arld	stee!	charge	
			1100 g/1	t zinc su	lphate		
	N	OTE:					
	P	lant vant	hate contai	ins 50% e	thyl xa	nthate and	
		507 4.46.	ophosphate	(fyanan)	d 2411		

11-204 Bulk copper,lead,pyrite flotation in nitrogen following nitrogen conditioning, 5 g/t collector and zinc sulphate depression. Zinc flotation in nitrogen.

	SOLIDS IN	ASS BALANCE		COPPER N	ASS BALAM	Œ	ŧ	EAD MASS I	ALANCE		Z	INC MASS B	ALANCE		1	POW MASS B	ALANCE	
	5amole	mass pull	c	opper	£0 9	per	lea	4	lea	•	210	c	\$10	c	110	a	170	30
	##1glit	sample tot	al assa	y whits	reco	127 /	8558y	units	recov	ery	4552y	units	recev	ery	425 8 V	units.	FECOV	ref y
Sample reference	(gr aes)		(2)		stage	total	(Z)		stage	total	(2)		stage	total	(2)		stage	total
bull rinigher 39 seconds	176 55	17.731 17.	.731 1.7	3 0.307	45.221	45.221	1.14	0. 202	28.742	28 741	3.21	0.569	5.551	5.552	40.46	7.173	24.992	24, 792
hill irasenger - 1 minutes	48 02	6 83% 24.	.56Z 3.0	3 0.207	30.51X	75.732	2.18	0.147	21.10%	40 922	7.78	0.531	5.187	10.732	35.44	2.421	0.432	33.42
oulb scavenger II - 7.5 minutes	0.00	0.001 24.	.56Z 0.0	0 0.000	0.001	75.732	€.00	0.000	8.00I	49.92%	0.00	0.000	0.001	10.731	0.00	0.000	0.00Z	33.42
tanc ciecust																		
introopher time ininite	168.50	16.72% 41.	.492 0.7	0 0.118	17.462	•3.192	1.45	0.245	34.892	84.612	47.76	0.081	78.82%	87. 551	11.76	1.990	4.932	40.33
anr rougher - tame 2 manites A	21.94	2.201 43	681 0 4	2 0.009	1.362	94 562	0.84	0.017	2.432	87.44Z	35.27	0.777	7.50%	97.132	10.24	0.402	1.492	41.75
incompler tree 2 minutes 0	0.00	0.001 43	681 0.0	0.000	0.002	94.56%	0.00	9.000	0.00Z	87.441	0.00	0.000	0.002	97.132	0.00	8.00 0	0.002	41 7
enc scavenger - time 30 seconds	108 68	10 911 54	601 0.0	6 0.007	1.291	95.951	0.28	●.031	4.351	°1.78%	1.54	0.169	1.442	98. 771	44.24	4.870	16.877	58.57
inc scavenger - time 1 minete	23 42	2 352 56	952 0.1	0.002	0.352	74.192	0.26	0.006	0.872	92.651	1.17	0.028	0.27%	99.032	43.16	1.015	3.542	62.1
zinc sravenger - time 2 minutes	9,00	0.001 56.	.951 0.0	0 0.000	0.001	96-191	9.00	0.000	0.002	92. 451	0.00	0.000	0.001	77.031	0.00	0.000	0.001	47.1
inal tails .	428.70	43.05Z 100.	.001 0 (å 0.02å	3,812	100.00I	0.12	0.052	7.351	100.00Z	0.23	0.009	0.971	100.001	25.27	10.877	37.072	100.0
reconstituted feed	795.81			0.678	100.001			0. 703	100.001			10.254	100.00I			28.706	100,001	
EST SIRMARY.	sample	mass puli		Copper	recov	er y		lead	recov	ery		21 AC	recov	ery		IFOR	FECO	very
	weight	sample to	tal	assay I	stage	total		assay I	stage	total		assay l	stage	total		2553y I	stage	tota
ull Untation	244.57	74.567 24	.561	2.09	75 732	75.731		1.43	49.922	47.721		4.48	10.732	10.731		39.04	33.421	32.6
incrompter flotation	190.44	17.121 43	189	0.67	10.831	94.561		1.39	37.521	87.442		46.37	86.39Z	•7. 13X		12.51	8.332	41.
ing scavenger flotation	137.10	13.271 56	.952	9 08	1.632			0.28	5. 221	92.65Z		1.47	1.911	?7.031		44.05	20.351	67.1
final flotation tails	420.70	43.05Z 100		0.06		100.002		0.12	7, 352	100.00Z		0 23	0.972	100.00I		25.27	37.892	100.

11-205 Bulk copper,lead,pyrite flotation in air following 5 g/t collector addition and zinc sulphate depression. Then zinc flotation in air.

9.5	activator added g/t	Pre-cor	nditioning time	coll type	lector addition q/t	flotation gas
		gas	time	суре		gas
					3	
	-	alr	3 a in	-	-	alr
9.5	•	-	-	-	-	alf
10.5	1100	none	5 410	YFY	75	air
10.5	•	-	-	-	-	air
10.5	-	_	-	KEY	20	alr
10.5	-	-	-	-	-	air
		6RINDIN	6 CONDITION	NS:		
		20 mm	utes. eild	steel c	harge	
		1100 g	/t zinc sul	lohate		
N	OTE:					
P	lant xant	hate cont	ains 50% el	thyl xan	ithate and	
	502 dithi	ophosphat	e (Cvanami	d 241)		
	10.5 10.5 10.5	10.5 1100 10.5 - 10.5 - 10.5 - NOTE:	10.5 1100 none 10.5 10.5 10.5 6RINDIN 2.0 min 1100 g	10.5 1100 none 5 min 10.5 10.5 10.5 6RINDING CONDITION 20 minutes, mild 1100 g/t zinc su NOTE: Plant xanthate contains 50% e	10.5 1100 none 5 min KEX 10.5 KEX 10.5 KEX 10.5 KEX 10.5 1	10.5 1100 none 5 min KEX 35 10.5 KEX 20 10.5 KEX 20 10.5

11-205 Bulk copper, lead, pyrite flotation in air following 5 g/t collector addition and zinc sulphate depression. Then zinc flotation in air.

	SOL 195 #/	nss balam	Œ		COFFEP MA	55 BALANC	£	Ĺ	EAD MASS I	AL MICE		1	INC MASS B	AL AMCE		i	POW PASS 8	AL MICE	
	5400 i n	0355 P	ul!	copp) t r	(09)	er	lea	ŧ	ìea	đ	210	τ	211	c	110	•	iro	30
	weight	sa e ple	total	4553Y	units	recov	ery	2552y	units	recov	er y	assay	units.	recov	er y	assay	units	7000	185)
sample reference	(graes)			(2)		stage	total	(2)		stage	total	(2)		stage	totai	(1)		stage	total
collectorless rougher - I minute	5 8 50	5 651	5 651	7 52	0 425	67.68I	67.682	3.91	0 221	29.197	?a.1az	12.48	0.705	4.882	4 8 62	27 57	1.557	5.717	5.217
collectories scavenger - 3 minutes	36 76	3.55%	a 502	2.97	0 105	16.801	84 47Z	3.31	0.117	15.531	44 771	14.40	0.511	4.991	11.671	20.00	1.025	3.432	8.647
collectorless scavenger II - 2 5 minutes	0.00	0 002	7.201	0.00	0 000	0 001	84.472	●.00	0.000	0.00I	44.721	0.00	0.000	0.002	11.872	0.00	0.000	100.0	8.447
rinc cougher - time I minute	131 77	12 727	21.922	0 31	0.039	6.787	90 762	2.20	0.280	34.991	B1 711	45.43	5.780	56 432	68 312	13.22	1.682	5.632	14.27
zior rougher - time 2 minutes A	44 B1	4 131	26 251	0.30	719.0	2.07%	92 832	9.70	0.030	4.002	85 71Z	35.66	1.543	15.06Z	83 372	16.79	0 726	2.431	16.71
zanc rougher - tame 2 manutes B	0 00	0 001	26.25%	0 00	9.000	0.001	72.837	0.00	0.000	0.00Z	85.712	0.00	0.000	0.001	03 372	0.00	0.000	0.002	16.71
zenc scavenger - time 2 minutes A	60 70	5 871	30.117	0 16	0 0∪≎	1 591	94 121	0 38	9.422	2.05%	5: MAI	25 9 0	1 514	14.792	98.152	26 13	1.574	5 132	21.84
rinc scavenger - time 7 ex stee B	14 57	1 402	15 EJ4	7	1 (11	9 38Z	94.70%	0.43	0.906	0.Bez	87.462	3.53	0.04	0.4 8 %	98 64Z	37 93	0.51	1.742	23.56
zinc kia enger - time 2 minutes C	0.00	0.001	33.521	0.00	0.000	0.001	94.701	0.00	0.000	0.00Z	60 491	9.09	0.00*		e8 142	÷		0.001	23.58
final tails	688.69	46.487	190.092	0.05	0.933	5 301	100.001	0.12	0.000		100.001	0.21	0.140	1 361	100.002	34.34	22.630	76.421	100.00
reconstituted feed	1035.75				0.628	100.001			0.757	100.00Z			10.242	100.001			29.073	100.001	
IESE SIMMARY-	sample	#8 95 (el l		copper	recov	ery		lead	recov	er y		2100	recov	ery		17 OR	recov	rery
	weight	sample	total		assay I	stage	total		assay I	stage	totai		assay I	stage	total		assay I	stage	tota
collectoriess flotation	95.26	9.203	9.201		5.76	84. ; " į	84 472		3.68	44.721	44 721		13.22	11 872	11.872		28.08	8.641	0.6
zine rougher flotation	176.58	17.05%	26.251		0.31	8.351	92.831		1.12	41.00Z			42.95	71 502			14 13	8.061	16.7
zinc scasenger flotation	75 1	7 272	33 52%		0.16	1.882	74.793	:	0.39	3.742			21.51		°8.641		28-23		23 5
final flotation tails	688.60	66.482	100 00Z		0.05	5.302	100.002		0.12	10.54Z	100.001		0.21	1.362	100.00I		34.34	76.422	100.0

11-301 Collectorless flotation followed by zinc flotation.

N	
w	

	F L	0 T A	T 1 0	N C	0 N I	1 1 1	0 N S
Sample reference	ρН	activator added	Pre-con	ditioning	coll type	ector addition	flotation Qas
		g/t	gas	tine	сурс	g/t	yas
collectorless rougher - 1 minute	9.5	-	aır	10 min	-	-	a)r
collectorless scavenger - 3 minutes	۹.5	-	-	-	-	-	air
zinc rougher - 1 minute	10.5	1100	none	5 a in	KEX	35	aır
zinc rougher - 2 minutes	10.5	-	-	-	-	-	air
cinc scavenger - 2 minutes A	10.5	-	-	-	KEX	20	air
zinc scavenger - 2 minutes B	10.5	-	-	-	-		air
			GRINDIN	6 CONDITIO	NS:		
			20 min	utes, eild	steel o	tharge	
			1100 g	/t zinc su	lphate		
	N	OTE:					
	P	lant xant!	nate cont	ains 50% e	thyl xar	ithate and	
		50% dithi	nnhoenhat	e (Fuanae)	d 241)		

11-301 Collectorless flotation followed by zinc flotation.

12. APPENDIX 4: KIDD CREEK TESTWORK

STANDARD AND NITROGEN FLOTATION

- 12-101 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation in air.
- 12-102 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.
- 12-103 Bulk copper, lead flotation in air, pyrite flotation in gas containing 95% nitrogen and 5% oxygen, and zinc flotation in air.
- 12-104 Bulk copper, lead flotation in air, pyrite flotation in gas containing 90% nitrogen and 10% oxygen, and zinc flotation in air.

	SOLIDS M	ASS PALAN	LE		COPPER M	ISS MILANC	Œ	(EAD MASC	BALANCE		1	INC MASS	BALANCE			RON MASS	PALANCE	
	5.100 le	0055 p	ol1	cop	er	נייף	ier	10	ud	le	H	zin	K	210	nc	ire	10	ir	
	weight	sa op i e	total	455ay	units	1600	•		units	r ece			wats	reco	1 27 y		mits	F9081	487 Y
sample reference				(2)		stage	tctal	(2)		stage	total	(2)		stage	total	(2)		st age	tetal
copper/lead rgfir - 2 minute A	53.30	5.971	5.971	1.51	0.517	63.852	63.851	7.12	0.544	32.962	32.961	13.40	0.800	12.1 Æ	17.131	24.69	1.448	7.182	7.191
copper/lead right - 2 ainute D	30.30	3.392	7. 36Z	3.27	0.111	13.832	77. 69 Z	8.62	0.293	17.712	50.662	14.40	₽.563	8.342	20.672	23.50	0,797	3.902	11.072
copper/lead rain - 2 minute C	26.30	2.951	12.312	1.52	0.045	5.50X	83.272	7.30	0.215	13.021	43.401	18.40	0.542	5.222	20.00Z	24.90	0.733	3.581	14.467
copper/lead right - 2 minute 9	31.00	3.472	15.78Z	0.74	0.033	4.0/%	87.341	3.20	0.114	4.872	70.572	14.70	0.510	7.742	3e.621	27.30	0.746	4.631	17.272
pyrite rangher - 2 minutes A	15.90	1.70%	17.561	0.86	0.015	1.912	87.251	2.44	0.043	2.632	73.20 1	15.50	●.276	4.102	49.817	24.80	0.442	2.152	21.451
pyrite rinigher - 2 aimites B	11.00	1. 722	18.88Z	0.53	0.007	0.872	90.122	2.37	0.031	1.70Z	75.102	11.00	0.156	2.361	43.17%	27.60	0.365	1.781	23.231
pyrate resigner - 2 minutes C	19.60	1.191	20.071	0.40	0.005	0.591	90.711	1.94	0.023	1.372	76.492	11.60	0.140	2.121	45. 20%	29.60	0.351	1.721	24, 951
pyrate rangher - 2 arautes 8	10.50	1.187	21.25%	0.27	€.003	0.432	11.142	1.47	0.017	1.052	77.542	9.59	0.113	1.712	47.00Z	32.00	0.374	1.847	26.79
zinc raugher - 30 seconds	22 80	2.551	23.001	0.15	0.004	0.491	71.621	0.40	0.010	0.622	78.16Z	47.40	1.215	10.432	65.432	9.31	9.238	1.16%	27.95
21M rougher - 1 ainute	29.90	2.342	26.14Z	0.19	9,904	0.531	92.14%	9.57	9.013	0.017	78.97%	40.20	0.741	14.272	79. 701	11.20	0.262	1.201	27.231
zinc rougher 0 - 30 serouds	31.40	3.541	29.681	0.14	0.005	0.621	92.762	0.02	0.029	1.761	8 0.721	23.00	0.814	12.34 z	92.04Z	26.90	0.746	4.641	33.071
zinc rougher B - † minute	54.30	4.09Z	35.74%	0.11	0.007	0.032	93.592	0.00	0.047	2.952	83.67Z	3.04	0. 185	2.801	74.841	33.40	2.031	9.932	43, 801
final tails	573 40	64.242	100.001	0.00	9.051	6.412	100.001	9.42	0.270	14.332	100.00Z	0.53	0.340	5. 16Z	100.001	17.90	11.479	56.30%	100.007
reconstitued feed	892.90				●. 902	100.00I			1.452	100.00Z			6.596	100,001			20.460	100.002	
TES! SUMMAPY:	sample	8855 (Pal l		Copper	recov	ery		lead	recev	er y		2180	recov	ery		1708	F@C@V	ery
	merant	samle	total		assay I	stage	total		assay I	stage	total		assay I	stage	total		assay I	stage	total
copper/lead flotation	140.70	15.781	15.781		4.41	67,341	87.341		7.39	70.571	70.571		15.31	36.622	36.622		25.01	19.292	19.29
pyrife flotation	48.80	_	21.25%		0.54	3.501	91.142		2.11	6.97%	77. 541		12.53	10.302	47.603		20.07	7.501	
zamc rougher flotation	43.70		26.142		0.16	1.001			0.48	1.432	78.972		44.66	32.472	77.702		10.21	2.441	
zinc rougher flotation 8	85.70		35.74%		0.12	1.452			0.01	4.701			10.38	15. 142			30.97	14.561	
final flotation tails	573.40	64.242	100.00Z		0.00	4.412	100.00Z		0.42	16.33Z	100.00Z		0.53	5. 142	100.00Z		17.90	56.201	100.00

12-101 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation in air.

	pH	activator	Pre-cor	nditioning		ector	flotation			
sample reference		added g · t	a 3 c	tiee	t,pe	addition	gas			
		y t	gas	[] @E		g./t		DISSOLVED OXYGEN LEVELS: ppm.	BEG INNING	END
Copper lead rougher - 2 minute A	10.2	-	air	10 min	SIF:	5	air	copper/lead rougher - 2 minute A	7.5	10.1
Topper lead rougher - 2 minute B	10.2	-	-	-	SIFI	5	air	copper/lead rougher - 2 minute B	7.3	10.
coper lead rougher - 2 minute C	16.2	-	-	-	SIF	5	air	copper/lead rougher - 2 minute C	6.5	9.
copper lead rougher - 2 minute D	10.2	-	-	-	SIPI	5	alf	copper/lead rougher - 1 minute D	6.3	7,
:.rite rougher - time 2 minutes A	6.5	-	air	5 a in	SIP:	5	air	pyrite rougher - time 2 minutes A	7.8	~. ;
ovrite rougher - time I minutes B	°.5		air	2 min	SIP	5	air	pyrite rougher - time I minutes B	7.8	∹.
Cirite rougher - time I minutes C	Ģ.E	-	air	2 min	SIFX	5	air	pyrite rougher - 1.me 2 minutes C	7.8	7.
purite rougher - time 2 minutes D	۰.5	-	air	2 min	SIFY	5	air	pyrite rougher - time 2 minutes D	7.8	7.
inc rougher - time 30 seconds	11.0		-	5 e in	-	-	alr	zinc rougher - time 10 seconds	7.0	7.
rinc rougher - time 1 minute	11.0	-	-	-	-	-	alr	zinc rougher - time 1 minute	7-1	٦.
cinc rougher B - time 30 seconds	11.0		-	-	SIF	50	air	zinc rougher 8 - time 30 seconds	7.1	٦.
zinc rougher 8 - time 1 minute	!1.0	-	-	-	-	•	air	zinc rougher 8 - time 1 minute	7.1	٦.
			GRINDI	N6 CONDITIO	NS:					
				n6 CONDITIO nutes, øild		charge				

12-101 Bulk copper, lead flotation in air, pyrite flotation in air and zinc flotation in air.

	SOCIES	SOLIDS MASS BALANCE	딾		COPPER IM	COPPER RASS BALANCE	juj	184	LEAD MASS BALANCE			21MC	ZINC MASS BALANCE			100	POR MASS BALANCE	1	
Sample reference	samele merght (grans)	sample total	total	copper 8558y W (2)	# # # # # # # # # # # # # # # # # # #	cooper recovery stage to	er tetal	lead assay units (2)		lead recovery stage to	3	23.00 2553y WH (2)	a in the contract of the contr	zinc recovery stage to	3	1788 2554 W	waits	recovery stage tot	iotal
Copper flesd rafte - 2 annute A copper flesd rafte - 2 annute B copper flesd rafte - 2 annute C	35 30 20.80 13.20 12.80	3.762 2.212 1.402 1.363	3.762 5.972 7.372 0.742	9.36 6.52 3.12 1.91	0.52 0.144 0.044 0.026	51.552 21.162 6.431 3.013	51 551 72.711 79.141 82.951	6.00 16.20 11.10 10.10	6.225 6.226 0.156 6.138	13.931 13.952 9.641 0.501	13.932 27.061 37.521 46.022	10.00 11.20 17.70	6.376 6.319 6.249 6.759	5.85 1.87 1.83 1.83	5.851 10.741 10.612	25.00 23.30 22.40	0.939 0.516 0.315 0.327	5.131 2.017 1.721 1.702	5.131 7.942 9.661 11.442
Pyrife compler - Z aimelen g. pyrife compler - Z aimelen B. pyrife compler - Z aimelen C. pyrife compler - Z aimelen D.	26.50 34.00 51.70 12.00	2 872 3 672 5.522 8 772	11.562 15.172 20.702 29.422	9.80 0.24 0.12	0.073 0.009 0.007	3,212 1 273 0 972 1,591	96.261 87.541 86.511	11.38 2.25 1.05	0.31° 0.081 0.058	19.691 5.631 3.581 4.371	65.772 70.752 74.531	11.10 4.62 3.23 3.59	6.313 6.174 6.174 6.313	4.871 2.712 2.781 4.862	23.511 24.221 29.601 33.671	2 3 8 8 2 2 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3	6.891 1.158 1.822 2.810	4.372 6.327 •.951 15.332	15.012 22.131 32.061 07.412
zinc resubber 30 seconds zinc resubber 8 30 seconds zinc resubber 8 30 seconds zinc resubber 8 Lanute	13.78 15.28 15.58 13.10	4 4/1 1.831 1.652 1.392	34.092 35.922 37.572 39.972	6.19 6.23	9.008 9.003 0.003	1.231 9 512 9.602 9.453	91.28X 91.79X 92.39X	6.50 6.74 6.88	0.023 0.010 0.012 0.012	1.441 0.611 0.751	99.151 99.761 91.511 01.271	5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	2.322 6.818 9.407 9.122	36.132 12.732 7.572 1.902	70.011 B2.741 90.312 92.212	16.74 15.10 15.20	6.408 6.249 6.249	2.231 1.021 1.341 2.641	5.641 52.641 54.701
final trils reconstituted tend	573.60 939.80	61.032 100.002	200.00	8.	0.662	7.161	100.001	•.47	1.610	17.732	100.001	0.82	6.58 1.63	7.791	100.001	13.46	0.301 10.322	45.301	100.001
IEST STYMBRE	saple meicht	eass pull sample total	12 TE TE	-	COpper assay 1	stage t	total	ŕ	lead assay 1	recovery	r tetal	•	71RC 25587 Z	L	r, total		17 OA 85 SAV 12	recovery stage to	r tatal
coprer tead flotation parity flotation zint rougher flutation zint rougher flutation final flutation tails	82.16 194.40 61.16 28.60 573.60	29 691 6 501 3 041	8.742 29.423 35.922 38.972 109.002		6.48 0.18 0.24 0.08	12.952 7.092 1.741 1 052	72.951 90.042 91.791 92.841		2.52 2.54 6.51 6.47	44. e21 32. e81 2. e51 1. 512 17. 731	46. 921 78. 701 80. 761 82. 271 1900. 002		7. 2. 4. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5.	15.247 15.247 4.472 7.7°2	18.642 33.871 82.742 92.211		24.00 31.86 9.15 24.31 13.60	11.441 35.971 3.251 4.941 45.301	11.442 47.411 50.661 54.701

12-102 Bulk copper, lead flotation in air, pyrite flotation in

nitrogen and zinc flotation in air.

	pH	activator	Fre-condi	tioning	coll	ector	flotation			
sample reference	·	added		•	t, pe	addition	gas			
		g t	gas	time		g ' t				
								DISSOLVED DXYGEN LEVELS: ppm.	BEGINNING	END
copper/lead rougher - 2 minute A	10.2		alf	16 min	SIFY	5	alr	Copper lead rougher - 2 minute A	÷	?.
copper lead rougher - 2 minute B	10.2		-	-	SIPT	5	a.r	copper/lead rougher - I minute B	3.8	9.
copper'leas rougher - 2 minute C	10.2		-	-	SIPX	5	air	copper/lead rougher - 2 minute C	5.5	10.
copper/lead rougher - 2 minute D	10.2	-	-	-	SIPX	5	31r	copper/lead rougher - 2 minute D	5.4	۰.
pyrite rougher - time 2 minutes A	9.5	-	nitrogen	5 min	SIFX	5	nitrogen	pyrite rougher - time 2 minutes A	ი.5	0.
pyrite rougher - time 2 minutes B	٥.5	-	nitragen	2 110	SIPY	5	nitrogen	pyrite rougher - time 2 minutes &	0.5	ç.;
pyrite rougher - time 2 minutes E	9.5		nitrogen	2 610	SIPX	5	nitrogen	pyrite rougher - time 2 minutes C	2.5	0.5
pyrite rougher - time 2 minutes D	0.5	-	nitrogen	្ធី 🖦 បា	SIPY	5	nitrogen	pyrite rougher - time 2 minutes D	0.5	0.
zinc rougher - time 30 seconds	11.0		-	5 a in	-	-	air	zinc rougher - time 30 seconds	7.1	7.
zinc rougher - time 1 minute	11.0	-	-	-	-	-	315	zinc rougher - time i minute	7.3	7.
zinc rougher B - time 30 seconds	11.0		-	-	SIPX	25	air	Zinc rougher B - time 30 seconds	7.6	7.
zinc rougher 8 - time 1 minute	11.9	•	-	-	-	-	air	zinc rougher B - time 1 minute	7.4	7.
			6RINDIN6	CONDITIO	INS:					
			30 minu	tes, silo	steel	charge				

12-102 Bulk copper, lead flotation in air, pyrite flotation in nitrogen and zinc flotation in air.

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	SOLIOS N	ASS BALAN	CE		COPPER MA	SS BALANC	.E	(LEAD MASS I	MLANCE		ı	INC MASS	MLANCE		1	RON MASS 1	ML MICE	
	sample	0255 P	ui l	c op j	er	copp	er	lea	ud	300	ad	218	C	216	ıc	ire	M	ir	0e
	weight	saople	total	2552y	units	FPC BY	er y	-	unsts	rece			units	recev	ery	assay	units	rece	very
sample reference	(grans)			(2)		stage	tota!	(2)		stage	total	(2)		stage	total	(2)		stage	total
copper/lead rate - 2 ainute A	24 50	2 782	2.782	8.04	e 274	27.892	27.892	11.00	0.304	10.312	10.311	15.50	0.431	6.29%	4.201	24.60	1.681	3.291	3.291
comper flead rehr - 2 minute #	29.60	3.112	5.872	11.09	0.342	42.421	70.572	8.62	0.247	14.912	33.221	11.20	0.348	5.08X	11.30Z	25.60	0.795	3.032	7.122
copper lead rate - 2 minute C	17.70	2.071	7.952	1.24	0 026	3.201	73.721	8.39	0.173	10. 3BZ	43.612	18.40	0.380	5.551	16.932	25.40	0.525	2.531	9.651
copperited tyle - 2 minute 9	11.40	1.701	7. 152	0.74	0.011	1.401	75.121	5.67	9.448	4.062	47.472	17.70	0.212	3.092	20.021	28.20	0.337	1.421	11.202
pyrite cougher - 2 minutes A	16.50	1.732	10.061	3.31	0 957	7.151	82.271	9,22	0.160	1.561	57.22%	19.40	0.319	4.652	24.602	22.80	0.305	1.902	13. 182
pyrate rougher - 2 ainutes 0	21.50	2 262	13.142	1.77	0 040	4.98Z	87.25%	8 . 26	0.186	11.162	40.302	17.00	0.447	4.522	31.202	23.50	0.530	2.552	15.732
pyrite cougher - 2 minutes C	44.70	4 672	17.832	0.10	0.005	0.572	87 872	0.90	0.042	2.572	70.912	4.05	8 . 284	4.142	35.342	36 20	1.698	6. 1 0 T	23.911
pyrite rougher - 2 minutes 8	48 50	5 092	22.922	0 25	0.013	1.59%	89.42%	2.28	0.116	6 951	77.051	8.34	0.424	4. 202	41.542	38 89	1.975	7.512	33. 422
pyrite rougher - 2 minutes E	62.50	8.66I	31.5 01	0.11	0.010	1.191	90.612	9.44	€.055	3.32I	61.172	5.00	0.440	6.421	47.762	42.60	3.480	17.761	51.181
zanc einigher - 30 seconds	30.50	4.04Z	35.421	0.14	0.006	6.712	71.322	0.40	0.016	0.971		50.70	2.948	20.91%	77.00%	6.42	0.349	1.642	
Zinc complex - I minute	21 10	2.711	37.831	0.17	0.004	0.472	91.791		0.012	0.672	82.83I	41.20	0.912	13.371	91.70Z	12.60	9.279	1.342	
Zinc zungher 0 - 30 seconds	11 69	1.7?2	39.05Z	0 23	0.003	0.351	72.132	9.86	0.410	0.432	83 452	19.10	0-233	√3.49Z	94.60Z	24.10	0.293	1,411	
23 nc 4 migher 3 - 1 minute	18.20	1.912	49.961	0.21	9.004	9.501	92.631	●.47	0.017	0.991	84.452	4.23	9.081	1.181	95.782	61.00	0.783	3.771	59.342
final fails	562 60	59.04Z	100.002	0.10	0 959	7.372	100.002	9.44	9.260		100.001	8.47	€.28°	4.721	100.001	14.30	8,443	40.667	100.001
recon lituted feed	952 90				0.002	100.00I			1.471	100.002			4.048	100.002			20.766	100,00%	
TEST SIMMARY:	sample	8255	pui]		C Opper	recov	rer y		lead	FRCEN	rery		2180	recov	ery		1700	FPCDY	ver y
	weight	saepl e	total		assay 1	st age	total		assay I	stage	total		assay 1	st age	total		assay I	st age	total
copper/lead flotation	0 7. 2 9	9.152	•.151		4.50	75.121	75.122		8.70	47.672			14.98	20.021	20.021		75.59	11.202	11.202
preste flatation	213.70		31.502		0.51	15. 47Z			2.25	33.502			4.57	27. 94 Z	47.96Z		20.50	39.90Z	
Zam emigher fintation	59.60		37.832		0.15	1.182			0.44	1-462			47.34		91.201		*.90	2.98%	
zinc rougher flutation #	29 86		40.962		0.22	0.851			0.87	1.628			10.07		95.782		39.42		50.341
final flotation tails	562.40	59.041	100.002		0.10	7.371	100.001	l .	0.44	15.552	100.00Z		0.49	4.222	100.002		14.30	40.662	100.002

12-103 Bulk copper, lead flotation in air, pyrite flotation in gas containing 95% nitrogen, 5% oxygen, and zinc flotation in air.

		added		itioning	type	ector addition	flotation qas			
		g/t	gas	time	•	g/t	•	DISSOLVED OXYGEN LEVELS: ppm.	BEGINNING	END
opper'iesd rougher - 2 minute A	10.2	_		16	6 1 D v	£			-	J.15
opper lead rougher - 2 minute B	10.2	_	air -	16 min	SIPX SIPX	5	alr	copper/lead rougher - 2 minute A	6.6	10.0
apper/lead rougher - 2 minute C	10.2			_	SIFX	5	31F	copper lead rougher - 2 minute &	6.8	10.1
opper/lead rougher - 2 minute D	10.2	-	•	•		5	air	copper/lead rougher - 2 minute C	6.3	10.6
opper lead rougher - 2 minute v	10.2	•	-	•	SIPY	5	alr	copper/lead rougher - 2 minute D	7.2	10.7
yrite rougher - time 2 minutes A	9.5	-	95% N2	5 min	SIPX	5	951 N2	Pyrile rougher - time 2 minutes A	7.5	2.9
yrite rougher - time 2 minutes B	۰.5	-	95% N2	2 a in	SIPX	5	95% N2	pyrite rougher - time 2 minutes B	2.6	2.5
yrite rougher - time 2 minutes C	9.5	-	95% N2	I min	SIPX	5	951 N2	pyrite rougher - time 2 minutes C	2.5	2.5
yrite rougher - time 2 minutes D	9.5	-	95% N2	2 min	SIFX	5	95% N2	pyrite rougher - time 2 minutes D	2.5	2.5
inc rougher - time 30 seconds	11.0		-	5 a in	SIPX	25	air	zinc rougher - time 30 seconds	8.3	9.6
inc rougher - time 1 minute	11.0	•	-	-	-	-	air	zinc rougher - time 1 minute	8.5	9.5
inc rougher B - time 30 seconds	11.0		-	-	SIPX	25	air	zinc rougher B - time 30 seconds	8.2	9.1
inc rougher B - time 1 minute	11.0	-	-	-	-	•	217	zinc rougher B - time 1 minute	8.3	9.5
			GRINDING	CONDITIO	N5:					
			30 a ini	ites, mild	i steel (harge				

12-103 Bulk copper, lead flotation in air, pyrite flotation in gas containing 95% nitrogen, 5% oxygen, and zinc flotation in air.

	SQL I DS N	ASS DALAN	Œ		COPPER M	ASS BALANI	CE	ι	EAD MASS I	MLMCE		1	INC MASS I	MLANCE		1	PON MASS B	ALANCE	
	5acole	8455 #	uli	Copp	jer	C 09(per	lea	4	lea	d	218	c	219	c	176	ıa.	170	J4
	merdyf	Sample	tetai	assay	units	rece:	ver y	8558V	mits	reces	er v	25527	units	recov	er y	8558y	units	FPCOV	ret y
sample reference	(grans)			(2)		stage	total	(1)		stage	total	(2)		stage	total	(2)		stage	total
copper/lead robr - 2 minute A	32 80	3.501	3.50E	7.47	9.339	45. •22	45.92%	6.77	0.237	14.46Z	14.401	10.20	0.357	5.342	5.342	26.10	0.713	4.40Z	4,402
copper/lead right = 2 minute B	28.26	3.01Z	6.51I	6.43	0.174	26.252	72.17%	9.02	0.272	16.621	31.102	15.80	0.474	7.122	12.46%	22.90	0.600	3.332	7.738
copper/lead rahr - 2 minute E	22.87	2.442	8. 75:	7.51	0.061	6. 791	00.461	11.70	6.246	17.751	49.85I	20.46	0.498	7.442	19.901	22.00	0.556	2.481	10.412
copper/fead right = 2 minute 8	10.11	1.932	10.591	1.33	0.024	3.4 8 Z	83.742	5.62	0.107	4.64Z	55.491	10.40	0.355	5.321	25.271	23.00	9.460	2.221	12.631
pyrite raugher - 2 minutes A	14.51	1.552	12.432	1.06	0.016	2.22%	84.171	6.67	9.103	6.312	61.991	19.10	0.2%	4.422	29.442	24.00	0.371	1.792	14, 423
pyrate raugher 2 minutes 8	15.00	1.492	14.132	0.65	0.011	. I tes	87.662	5.04	0.077	4.051	67.84Z	17.30	0.293	4.382	34.02%	24.60	9.417	2.012	16.432
pyrite rougher - 2 minutes C	12.80	1.30X	15.502	9.47	0.006	0.882	89.532	3.84	0.057	3.232	71 072	15.70	9 216	3.232	37 251	33 80	0.382	1.847	12 272
pyrite raugher 7 oinutes B	■.50	9.712	16.417	9. 79	0.004	0.481	87.011	2.46	0.022	1.361	72.442	14.40	0.132	1.981	39.232	26.50	0.240	1.162	10.432
zinc romeher - 30 seconds	45.13	4.812	21 221	0.13	0.006	0.852	87.861	0.45	0.031	1.911	74 352	40.10	2.316	34.432	73.052	10.10	0.486	2.341	21.771
zinc rooter ainute	20.09	2.221	23.44Z	9.15	0.003	0.45Z	90.312	0.70	0.070	1.221	75.572	37.00	0.037	12.541	86.40 2	15.00	0.333	1.402	23.301
zinc rumpher B - 10 seconds	15.20	1.632	25.072	0.18	0.003	0.40Z	90.711	1.00	0.018	1.067	76.64%	22.70	●.373	5.501	71.70Z	22.00	0.359	1.731	25.117
zanc rougher D - I aanute	34.80	3.732	29.00I	0.12	0.005	0.44Z	91.352	0.07	0.034	2.091	79.731	4.26	0.167	2.501	94. 46 1	37.60	1.476	7.122	32.221
final tails	665.60	71.001	100.00Z	0.09	0.064		100.002	0.47	0,340		100.001	€.52	9.349		100.001	17.80	14.059		100.001
reconstituted feed	937.43				0.738	190.00Z			1.436	100.001			6.48 7	100.00Z			20.743	100.00I	
TEST SHIPMRY:	sample	0055	ret t		Copper	recov	er y		lead	7000	er y		2190	recev	•		100	recev	•
	ne) âyf	Saepi e	total		4554y Z	st age	total		assay I	st age	total		assay 1	stage	total		essay I	stage	total
copper/lead flotation	102.04		10.271		5.40	8 3. 94Z			8,34	55.492			15.49	25.221			24.07	12.632	
pyrite flutation	51.78		16.412		1.4	5.072			5.02	16.751			14.76	14.017			25.54	4.01	-
zinc romaher flatation	65. •3		23.442		9.14	1.301	70.311		0.73	3. 132	75.571		44.85	47.173			11.45	3.951	
zinc rougher flotation	52.90	5.542	70.091		0.14	1.042	91.352		0.93	3.162	78.732		9.73	6.00I	94.40I		33.02	1.04	
final fintation tails	645.60	71.00Z	100.00Z		0.00	8.452	100.001		0,49	21.272	100.001		0.52	5.522	100.001		17.80	67.78%	100.00

12-104 Bulk copper, lead flotation in air, pyrite flotation in gas containing 90% nitrogen and 10% oxygen, and zinc flotation in air.

			f _ 3	- 1 7	¥ []	: :	1 2 1	- : G v	:		
::	an:s re-granca	pH	EC11 E137 EC220	5-9-5182	ָּרור::::	:::1 1,55	aciir acciii:	flotation gas			
1			: t	315	11.36			žes			
			•	•				;	DISSOLVED OXYGEN LEMELS: ppa.	BESI, INTHE	END
::	ocer leso rougher - 2 ainute 4	10.3	-	217	1" 310	SIPY	5	air	cooper·lead rougher - 2 ainute A	5.3	٥٫٥
	oparilazo rougher - I ainuta 2	10.2	-	-	-	Sisa	٤	ŝις	copper/lead rougher - 2 minute B	6.4	
•	operileso rougher - 2 ainute C	10.2	-	•	-	SIFY	=	317	copper (lead rougher - 2 samute C	6.1 6.0	10.0 10.9
25	ober laad rougher - 1 minuta 0	10.2	•	-	-	ĞÎŁA	:	317	cooper/lead rougher - 2 sinute D	5.0	10.5
av	rice rougher - time 2 minutes A	9.5	-	e01 42	5 ain	SIPY	5	<u>6</u> 02 #5	pyrite rougher - tipe 2 ainutes A	5.7	e -
ργ	rita rougher - time 2 minutes B	0.5	-	007 42	2 310	S!SĂ	5	DUZ 315	pyrite rougher - time 2 minutes 8	5.7	5. E 7
27	rite rougher - time 2 minutes 8	0.5	-	oux 45	2 ain	SIPT	5	00% 112	P/fits rougher - time 2 minutes C	5.7	E *
ру	rite rougher - tide 2 dinutes 9	٥.5	-	±04. 45	2 310	SISA	5	907 42	cyrite rougher - time 2 minutes D	5.7	5.7
21	no rougher - time 30 seconds	11.0	1200	-	5 min	SIPX	25	air	Zinc rougher - time 10 seconds	9.5	a. 3
72	no rougher - time 1 minute	11.9	-	-	-	-	-	air	zinc rougher - tipe 1 minute	9.1	0.7
2:	no rougher B - time 30 seconds	11.9	•	-	•	SIPT	77	air	ling rougher 9 - time TO seconds	9.1	ລຸລ
21	no rougher 8 - time 1 minute	11.0	•	•	•	-	-	air	zinc rougher 8 - time 1 minute	8.2	10.1
				SRINDIN G	CONDITION	HS:					
				30 minu	tes. mild	steel o	thar ge				
				Zinc Su	lohate. 1	200 g/t					

12-104 Bulk copper, lead flotation in air, pyrite flotation in gas containing 90% nitrogen and 10% oxygen, and zinc flotation in air.

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13. APPENDIX 5

DISSOLVED OXYGEN CONSUMPTION IN A FLOTATION COLUMN

It is possible to make a rough estimate of the oxygen consumption in a flotation column based on the results reported in this thesis.

Basis:

10 metre column with 5 cm inside diameter. Total volume capacity of 0.02 m³. Bubbles rise typically at a rate of 2 cm per second, so the retention time for the gas is 500 seconds. The pulp retention time is assumed at 20 minutes, giving a feed rate of 0.8 litres per minute. At a pulp density of 1.25 kg per litre, the mass feed rate is 1 kilogram per minute.

Oxygen balance

Oxygen available:

The retention time of the air is 500 seconds, and the gas hold-up, 20% of 0.02 m3 (0.004 m3). From this, the gas feed rate can be calculated at 0.48 litres per minute which, at a density of 1.293grams per litre (at S.T.P.) is 0.6grams per minute. This contains 19% oxygen, and represents an oxygen feed rate of 0.12grams per minute.

Oxygen consumption (from mineral oxygen demand alone):

Figure 4.22 plcts the drop in oxygen content in nitrogenated water, and Kidd Creek pulp. The oxygen level in the pulp drops faster than in the water, because of the oxygen demand from the mineral. Difference between the slopes of the two plots (in the 241)

first twenty seconds, i.e. before the oxygen levels are depleted and the two lines converge again) represents the rate of consumption of oxygen through mineral oxidation:

Drop of oxygen level in water 1.8 ppm in 20 seconds.

Drop of oxygen level in pulp 7.0 ppm in 20 seconds.

oxygen demand during the 20 second period 5.2 ppm.

This matches the oxygen demand measurements previously made on Canadian Shield ores (94). Their work showed that as the pulp ages its oxygen demand drops. A profile from their work which matches Kidd Creek ore is given in Figure 13.1.

ppm DISSOLVED OXYGEN

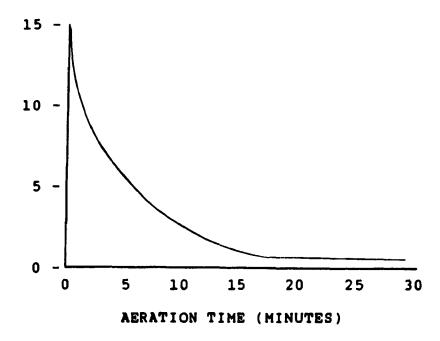


FIGURE 13.1 - Approximate oxygen demand profile for Kidd Creek ore.

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From Figure 13.1, the total oxygen demand for the pulp during a 20 minute residence time can be estimated by linear approximation of the integration to 20 minutes as 90 ppm.

The feed rate to the column is 1kg per minute, and this has an oxygen consumption rate of 90 ppm. So the oxygen demand from the pulp will be 0.09 grams per minute.

SUMMARY:

Oxygen supply from air: 0.12grams per minute

Oxygen requirement for oxidising the pulp: 0.09grams per minute

Oxygen remaining in the air at the top: 0.03grams per minute,

which is a quarter of the oxygen content of the air entering the

column. The gas would therefore be 95% nitrogen and 5% oxygen.

Pyrite is activated at such levels of oxygen content.