An Aspen Model for the atmospheric acid leaching of nickel-copper matte

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Abstract

The note summarized in this paper deals with modeling and simulation of atmospheric leaching process of nickel-copper matte. Computer simulation is a cost effective means of examining how the process is affected by using certain quantities of chemicals, varying flow rates, and varying operating conditions. To understand the nature of the leaching process during the stage of operation, the leaching mechanism of the matte is explained. Through simulation the effects of different parameters on nickel leaching were investigated, mainly operating temperature, and acid concentration. The modeling of this process was performed on a computer simulation package known as Aspen Plus. Under atmospheric pressure 68.8% of nickel, 14.05% of iron, 24.92% of copper, 10.66% of cobalt and 4% of zinc are extracted under suitable conditions of sulfuric acid concentration and reaction temperature.

Key words: Nickel-copper matte, atmospheric leaching process, computer simulation.

1. Introduction

Nickel-copper mattes are conventionally treated by pyrometallurgical method, but later the hydrometallurgical processes have gained an advantage for recovering valuable metals and reducing the environmental impact of industrial activities related to the processing of nickel ore deposits [1]. Nickel has strategic and industrial importance due to its extensive application in foundry, producing stainless steel, alloy, battery, and catalyst [2]. Recently, relatively great attention has been paid to the research on the recovery of nickel from secondary resources: spent catalyst [3], waste battery [4, 5] and tailings [6].

Nickel is commonly found in iron meteorites as the alloys kamacite and taenite. Nickel ores are of two types [7]: (1) Primary sulphide deposits associated with mafic and ultramafic rocks, (2) Nearsurface laterite deposits formed over olivine-rich host rocks following intense weathering. The sulphide ores have been the major source of nickel; however, the lateritic ores have been estimated to constitute about 73% [8] of the known nickel reserves of the world. According to the latest data by the USGS, world nickel reserves amount to 80,000,000 metric tons. From the chart below, one can see that Australia and New Caledonia hold currently 45% of the world's nickel reserves [9].

As a result of the high nickel production costs associated with traditional pyrometallurgical techniques and the depletion of high-grade sulphide ores, renewed interest has been developed concerning the production of the nickel and cobalt by high pressure acid leaching (HPAL) of nickel laterites. Industrial leaching processes generally comprise multiple leaching autoclaves that accommodate 3-phase (solid-liquid-gas) systems that exhibit complicated chemistry and kinetics. Given that leaching processes are principally the location of hydrometallurgical refinery bottlenecks, it is essential that these processes operate optimally to ensure maximum refinery throughput [10]. Understanding how changes in process and operating conditions influence the performance of the leach is of paramount importance in arriving at this state of operation.

2. Nickel processing

Conventionally, two process routes are employed for leaching Ni-Cu matte. The first one involves both atmospheric and pressure leaching in acidic nickel-copper sulphate solution, with oxygen being the oxidizing agent. In this process the matte is first leached under atmospheric conditions followed by a pressure leach step, and in both steps O_2/air is splashed into the leaching vessels. During atmospheric leaching substantial quantities of nickel and cobalt are dissolved while copper and iron are precipitated from the solution, and any platinum group metals (PGM) present remains in the solids. The precipitated copper and any unleached nickel and cobalt are dissolved in the subsequent pressure leach step or, in some cases, fed to a smelter. The other leaching process route employs a pre-leach step, which is essentially a matter repulping step, prior to a pressure leach step.

The repulping process may be considered to be basically a non-oxidative atmospheric leach stage since leaching of the matte starts immediately after it comes into contact with the $CuSO_4$ -H₂SO₄ repulping solution. During the pre-leach stage nickel, iron and cobalt are partially dissolved, while the copper is precipitated. In both of these Ni-Cu matte process route alternatives, nickel and cobalt can be recovered from the solution as metal by electrowinning or hydrogen reduction, or they may be recovered as sulphate crystals. Copper can be recovered as metal by electrowinning from the sulphate solution or by electrorefining if the precipitated copper was produced by the smelting process [11].

This case study deals with the leaching of nickel-copper matte. The principal phases present in the matte are metallic nickel (Ni⁰), millerite (NiS), metallic copper (Cu⁰), chalcocite (Cu₂S), CuS, metallic cobalt (Co⁰), metallic zinc (Zn⁰), metallic iron (Fe⁰) and pyrrhotite (FeS), Pyrite (FeS2). Nickel contributes to approximately 65% of the total particulate feed.

3. Leaching mechanism based on process chemistry

Based on the literature data for Ni-Cu matte leached in acidic solution, $CuSO_4$ -H₂SO₄-O₂, and the characterization of the leaching liquor and residues [11-13], the principal leach chemistry is the leaching of metals with simultaneous precipitation of aqueous copper via the cementation process and the atmospheric pressure oxidative dissolution in the sulfuric acid. The actual reacting system is a much more complex series-parallel reaction network involving, among other characteristics, metals oxidation and the progressive oxidation of sulfides. Through cementation process, metallic nickel is being leached according to reaction (1). The dissolution of metallic iron is believed to proceed according the well-known reaction of cementation of copper by metallic iron [14], reaction (2).

$$Ni^{o} + CuSO_{4} \rightarrow Cu^{o} + NiSO_{4}$$

$$Fe^{o} + CuSO_{4} \rightarrow Cu^{o} + FeSO_{4}$$

$$(1)$$

$$(2)$$

For the leaching of metals by direct acid attack; in the absence of an oxidant, such as air or oxygen, some of the nickel, cobalt and copper from the alloys are believed to be leached by the sulfuric acid according to the following reactions:

$Ni-Cu+2H_2SO_4 \rightarrow NiSO_4+CuSO_4+2H_2$	(3)
$Ni-Fe+2H_2SO_4 \rightarrow NiSO_4+FeSO_4+2H_2$	(4)
$Ni-Co + 2H_2SO_4 \rightarrow NiSO_4 + CoSO_4 + 2H_2$	(5)

The other possible reactions are the leaching of NiS, FeS, CuS and Cu_2S by the sulphuric acid, reactions (6) through to (9).

$NiS + H_2SO_4 \rightarrow NiSO_4 + H_2S$	(6)
$FeS + H_2SO_4 \rightarrow FeSO_4 + H_2S$	(7)
$CuS + H_2SO_4 \rightarrow CuSO_4 + H_2S$	(8)
$Cu_2S + H_2SO_4 \rightarrow Cu_2SO_4 + H_2S$	(9)

4. Nickel refinery process model and Aspen modeling

Simulating the Ni-Cu matte leaching process on an Aspen Plus simulation package serves as a useful tool to predict the behavior of the process. It is a cost effective means of examining how the process is affected by using certain quantities of chemicals, varying flow rates, and varying operating

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conditions. Simulators are often used at the laboratory and pilot plant stages of plant design, and it is no different for the process. This simulation acts as a guideline as to how this process will react under actual conditions. Aspen Plus makes it possible to determine the effects of non-ideal conditions on Ni-Cu matte leaching and serves as a guide for the trends that are obtained when analyzing various sensitivities.

For nickel leaching process, autoclave is the basic unit operation. An autoclave contains mechanical agitators and the compartments that are separated by weir gates. Basically, the leaching is performed in a compartment where after its content is conveyed into next compartment and so forth. For the present study case, the process model comprises three central unit operations: leaching reactors (autoclave compartments), tank and thickener. Tank was modeled as perfect mixer, whereby the output rate is equal to the sum of the input rates; the tank receives the dissolved metals. Thickeners were modeled as perfect splitters; it mixes all its feed streams then splits the resulting mixture into two or more streams according to substream specifications. Two unit operations and splitter model to separate the minerals no dissolved from the dissolved minerals. Within the leaching reactor the equilibrium data and the kinetics are not known, the atomic and mass balances are specified; the operating conditions are 90°C and 1 atm. Here a leaching reactor can be understood as one compartment of the autoclave.

5. Results and discussion

The Aspen simulation flow sheet is illustrated in figure 1. The feed stream 1, matte, enters at a flow rate of 20 kg/h, while feed stream 2 enters at a flow rate of 40 l/h; feed composition is summarized in Table 1. Atm-LCH is a leaching reactor. To model an autoclave of three compartments, three leaching reactors are arranged in series. The output of the atm-LCH1 is split into two streams, 4 and 5, the minerals not leached are conveyed into atm-LCH2 for further leaching through stream 5. For the last leaching reactor, it is fed through stream 9; streams 12 and 13 are the output. Streams 2, 6, and 10 are sulphuric acid streams added to atm-LCH1, atm-LCH2 and atm-LCH3, respectively. Streams 4, 8, and 12, mineral dissolved, are conveyed into the tank prior further processing of recovering dissolved minerals. To gain a deep insight in the operation and effect of changes in operating conditions, sensitivity analyses were performed; the thereafter discussions and remarks are produced from the first compartment, atm-LCH1.



Figure 1. Aspen simulation flowsheet

Leaching rate of the matte

From simulation results, the sulfides are leached to a high extent than their respective metals. The most suitable leaching conditions are as follows: sulfuric acid concentration of 10 M, and reaction temperature of 70 $^{\circ}$ C. On the other hand, all minerals in the matte are leached but to different rates; for

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each component of matte as a standalone, 42.83% of metallic nickel, 76.6% of nickel sulfide, 7.75% of metallic iron, 18.2% of iron sulfide (pyrrhotite), 11.33% of Pyrite, 20.5% of metallic copper, 26.31 of copper sulfide, 26.31 of chalcocite, 10.66% of metallic cobalt and 4% of metallic zinc can be extracted through an autoclave of three compartments under the thereof operating temperature and acid concentration.

6. Conclusions

(1) In the sulfuric acid leaching of Ni-Co concentrate, the effects of sulfuric acid concentration on the nickel extraction efficiency are remarkable; the effect of reaction temperature is not mild. Extraction efficiency of nickel and other minerals in the matte is improved by increasing both sulfuric acid concentration and reaction temperature to certain optimal values, 10 M and 70 $^{\circ}$ C, respectively.

(2) Considering the components of matte based on element, 68.8% of nickel, 14.05% of iron, 24.92% of copper, 10.66% of cobalt and 4% of zinc can be extracted under suitable conditions of sulfuric acid concentration and reaction temperature.

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