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Alpha-Alumina Extraction from Al-Ga'ara Bauxite

Abdul-Halim A. K. Mohammed, Ala Ibrahim, and Marwan G. Madhhoor

Chemical Engineering Department - College of Engineering - University of Baghdad - Iraq

Abstract

The bauxite produced from Al-Ga'ara area in Al-Enbar containing 50.4 wt. percentages Al_2O_3 was used for α -alumina production.

For α -alumina production bauxite was mixed with calcium carbonate in a ratio 1:3 and the mixture was burned at temperature range 1150-1350°C, cooled to 500°C; crushed and ground. The powder produced treated then with 6% sodium carbonate solution using different temperature (60-100°C) for fifteen minutes. After filtration, the produced sodium aluminate solution was reacted with 2% of calcium hydroxide at 98°C for thirty minutes. The filtrate was carbonated with purified CO2 gas at 70°C for different times (30-180min) and different flow rates (20-80 cm³/min). Alumina was precipitated as hydrate, then washed with water, dried at 110°C, and calcined at 1200°C. The produced α -AL2O3 was identified by X-ray diffraction.

Keywords: Alumina extraction, bauxite.

Introduction

In 1886-1887 [1] Bayer developed process for preparation pure alumina from bauxite. The Bayer process involves hydrochemical treatment of bauxite with sodium hydroxide at elevated temperature and pressure. On cooling, the aluminate liquor becomes supersaturated and alumina hydrate precipitates.

Variation of lime (Grzymek's method) [2] and limesoda sintering processes are used in treatment of highsilica bauxite for alumina production. In these processes a soluble aluminate is formed and silica is insolubilized by its reaction with calcium hydroxide.

Alumina is extracted from calcium aluminate produced from bauxite and limestone by treatment with Na₂CO₃ [2] by Equations 1 and 2.

$$CaO.Al_2O_3 + Na_2CO_3 \rightarrow 2NaAl_2O_2 + CaCO_3 \tag{1}$$

$$12CaO.Al_{2}O_{3} + 12Na_{2}CO_{3} + 5H_{2}O \rightarrow 14NaAlO_{2} \\ + 12CaCO_{3} + 10NaOH$$
 (2)

Desilication of aluminate solution is taken place by treatment with calcium hydroxide by Equation 3:

$$2Na_2SiO_3 + 2NaAlO_2 + Ca(OH)_2 + 2H_2O \rightarrow CaO.Al_2O_3.2SiO_2$$
 (3)
+ 6NaOH

Alumina hydrate is precipitated by treatment with CO₂ [5] according to Equation 4:

$$2NaAlO_2 + CO_2 + 3H_2O \rightarrow Al_2O_3.3H_2O + Na_2CO_3$$
 (4)

Conversion of alumina trihydrate to monohydrate occurs below 250°C by Equation 5.

$$Al_2O_3.3H_2O \to Al_2O_3.H_2O + 2H_2O$$
 (5)

Alumina monohydrate is converting to γ -alumina in the temperature range of 300 to 900°C by Equation 6.

$$Al_2O_3.H_2O \rightarrow \gamma - Al_2O_3 + H_2O \tag{6}$$

while α-alumina is produced at about 1100 to 1200°C. Sintering and acid-extraction process have been investigated for production of alumina from non-bauxite resources such as kaolin, anorthosite and nepheline ores containing 25 to 30 wt. % Al₂O₃ [3].

Yearly large quantities of alumina are produced for the manufacture of metallic aluminum, abrasive, chemical, refractories and ceramic [4].

Alpha-alumina is one of the most versatile of refractory ceramic oxides and finds used in a wide range of applications [5]. Alumina exists in a number crystalline form. Only one of these forms, α -alumina is thermodynamically stable. When hydrates are heated dehydration occurs and is accompanied structural changes. A series of metastable intermediates are progressively formed, the stable α -Al₂O₃ being the final product of thermal decomposition.

Dehydration and calcination of alumina hydrate also produce change in surface area and density. BET surface area reaches a maximum of about 350 $\rm rn^2$ g-1 at about 200°C and then decreases [6]. Density increases to near the theoretical value of 3.97 g cm-3 for α -alumina when calcined at 1200°C [7].

On the other hand, α-alumina was found in nature under the name corundum. There are several precious stones are forms of Al₂O₃ containing traces of other metal e.g., chromium (Red Ruby), iron or titanium (Blue sapphire) and (Black Emery) is an impure form [8].

Crystal structure of corundum has very low electrical conductivity, thus alumina is used commonly for high-quality electrical applications where low dielectric loss and high resistivity are needed [9].

Experimental Work

Materials

The bauxite produced by Al-Ga'ara area in Al-Enbar containing 50.4 wt. % Al₂O₃ as shown in Table 1.

Table 1 Chemical analysis of bauxite

Compound	Wt.%
Al ₂ O ₃	50.4
SiO ₂	30.04
CaO	0.72
Fe ₂ O ₃	1.17
TiO ₂	1.98
Na ₂ O	0.27
K ₂ O	0.08
MgO	0.67
L.O.I	14.67

The calcium carbonate produced by Al-Kilo 160 area in Al-Enbar containing 53.85 wt. % CaO was mixed with bauxite. Sodium carbonate (99.98 % purity) supplied from BDH. Calcium hydroxide (98 %) supplied from BDH. Carbon dioxide (99.99 %) was supplied from Baghdad Company.

Alumina Extraction

Bauxite particle and limestone with dimension of less than 90 μm had been mixed in 1:3 ratio, then the mixture was burned in electrical furnace at different temperature ranging from 1150 to 1350°C for six hours. Gradually,

the temperature reached to 500°C (cooling stage) the mixture is taken out. Dicalcium silicate (C₂Si), calcium aluminates 12CaO.7Al₂O₃ (C₁₂A₇) and CaO.Al₂O₃ (CA) are formed, as shown in Table 2 [10].

Table 2 Chemical composition (wt. %) of Al-Ga'ara bauxite and limestone mixture (mixing ratio 1:3) after calcinations at 1350°C

Compound	Wt. %
C ₁₂ A ₇	15.17
CA	3.84
C₂Si ·	80.29
Total	99.3

After burning, the mixture was crushed to fine powder. Alumina is extracted from the disintegrated powder by extraction with 6% Na₂CO₃ solution at different temperature ranging from 60 to 100°C for 15 minutes.

Sodium carbonate treatment was repeated three times. The calcium aluminates are converted to soluble aluminates according to Equations 1 and 2. The insoluble which contained C₂Si and CaCO₃ are removed by vacuum filtration at 60 mmHg.

Desilication of alumina solution is affected by treatment with 2 % calcium hydroxide at 98°C for thirty minutes according to Equation 5. The desilication process was repeated three times. The filtrate (aluminates solution) is then carbonated with CO gas 70°C for different times ranging from 30 to 210 minutes. Alumina hydrate precipitated according to Equation 4.

Alumina is then washed with water for removal of sodium salt, dried at 110° C, then crushed and calcined at 1200° C for 6 hours. The produced α -Alumina was identified by X-ray diffraction.

Results and Discussion

Effect of calcination temperature of bauxite and limestone mixture on the percentage extraction and yield of Al(OH)₃

Figures 1 and 2 present the effect of calcinations temperature in the range of 1150-1350°C on the percentage extraction of A1(OH)₃ based on Al₂O₃ in the raw material and on bauxite.

Figures and show that the increasing in the calcinations temperature increases the percentage extraction of Al(OH)₃, yields based on bauxite and Al₂O₃ in bauxite. The higher percentage extraction and yield obtained at 1350°C. The increasing in calcinations temperature resulted in a good transformation of bauxite-limestone mixtures to dicalciurn silicate CS and calcium aluminates CaO.Al₂O₃ (CA) and 12CaO.7Al₂O₃ (C₁₂A₇). The calcination temperature 1350°C gave 85.44 wt. % extraction of Al(OH)₃ from bauxite. This result is in agreement with Grzyrnek's work [11].

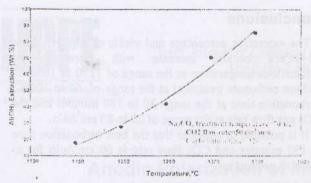


Fig. 1 Effect of calcinations temperature on the Al(OH)₃ extraction from bauxite

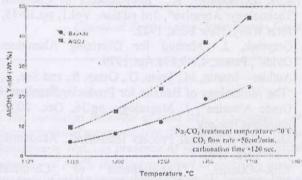


Fig. 2 Effect of calcinations temperature on the Al(OH)₃ yield based on bauxite and Al₂O₃ in bauxite

Effect of sodium carbonate treatment on the percentage extraction and yield of Al(OH)₃

Figures 3 and 4 show the effect of Na₂CO₃ treatment temperatures ranging from 60 to 100°C on the percentage of Al(OH)₃ extraction and the yields of Al(OH)₃, based on bauxite and Al₂O₃ in bauxite.

Figures 3 and 4 indicate that the percentage of Al(OH)₃ extraction and the yields increase with Na₂CO₃ treatment temperatures increasing.

It is possible to consider that the optimum sodium carbonate treatment temperature is 90°C, because higher temperature gives slight effect on Al(OH)₃ extraction, yields of Al(OH)₃ based on bauxite and Al₂O₃ in bauxite.

There is a slight difference in the percentage of Al(OH)₃ extraction from Ga'ara bauxite compared with the Grzymek work [12]. This may be due to the high SiO₂ (30 wt. %) content of Iraqi bauxite compared with that studied by Grzymek.

Effect of carbonation time and CO₂ flow rate on the percentage extraction and yield of Al(OH)₃

Figures 5 and 6 present the effect of carbonation time in the range of 30 to 180 minutes on the percentage of Al(OH)₃ extraction and yields of Al(OH)₃ based on bauxite and Al₂O₃ in bauxite.

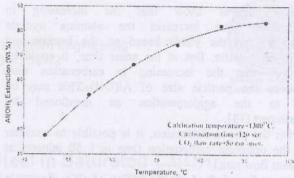


Fig. 3 Effect of Na₂CO₃ treatment temperature on the Al(OH)₃ extraction from bauxite

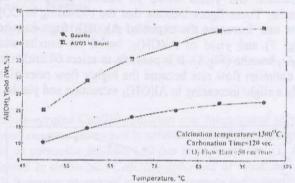


Fig. 4 Effect of Na₂CO₃ treatment temperature on the Al(OH)₃ yield based on bauxite and Al₂O₃ in bauxite

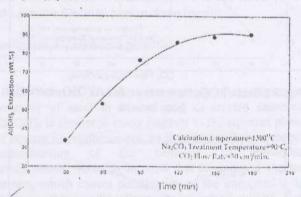


Fig. 5 Effect of calcinations time on the Al(OH)₃ extraction from bauxite

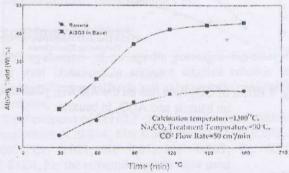


Fig. 6 Effect of calcinations time on the Al(OH)₃ yield based on bauxite and Al₂O₃ in bauxite

These figures show that the increasing in carbonation time increases the alumina hydrate extraction and the yields based on the bauxite and the Al₂O₃ bauxite, But at the same time, it could be observed that the increasing in carbonation time increases the particle size of Al(OH)₃. This may be due to the agglomeration as mentioned by Tschampert [13].

From the above discussion, it is possible to consider that the optimum carbonation time is 1 50 minutes at constant flow rate.

Figures 7 and 8 show the effect of CO₂ flow rate ranging from 20 to 80 cm³/min on the percentage of extraction and yields based on bauxite and Al₂O₃ in bauxite. These figure indicate that the increasing in CO₂ flow rate increases the extracted Al(OH)₃ from bauxite (Fig. 7) and yield of Al(OH)₃, based on bauxite and Al₂O₃ bauxite (Fig.8). It is possible to select 60 crn3/min as optimum flow rate because the higher flow rates give only a slight increasing in Al(OH)₃ extraction and yields.

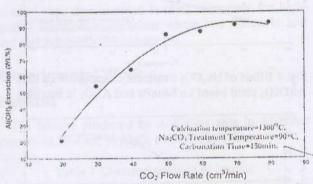


Fig. 7 Effect of CO₂ flow rate on the Al(OH)₃ extraction from bauxite

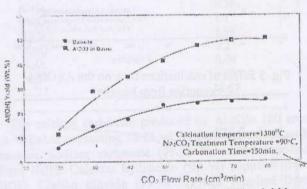


Fig. 8 Effect of CO₂ flow rate on the Al(OH)₃ yield based on bauxite and Al₂O₃ in bauxite

Conclusions

The extraction percentage and yields of Al(OH)₃ from Al-Ga'ara bauxite increase with increasing of calcinations temperature at the range of 1150 to 1350°C, sodium carbonate treatment at the range of 60 to 100°C, carbonation time at the range 30 to 180 minutes and the CO₂ gas flow rate at the range of 20 to 80 cm³/min.

It is possible to consider that the best carbonation time is 150 minutes and CO_2 flow rate is 60 cm³/min for α -alumina production.

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