Investigation of Dissolution in Diammonium Hydrogen Phosphate Solutions of Gypsum

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Abstract-Gypsum (CaSO₄.2H₂O) is a mineral that is found in large quantities in the Turkey and in the World. In this study, the dissolution of this mineral in the diammonium hydrogen phosphate solutions has been studied. The dissolution and dissolution kinetics of gypsum in diammonium hydrogen phosphate solutions will be useful for evaluating of solid wastes containing gypsum. Parameters such as diammonium hydrogen phosphate concentration, temperature and stirring speed affecting on the dissolution rate of the gypsum in diammonium hydrogen phosphate solutions were investigated. In experimental studies have researched effectiveness of the selected parameters. The dissolution of gypsum were examined in two parts at low and high temperatures. The experimental results were successfully correlated by linear regression using Statistica program. Dissolution curves were evaluated shrinking core models for solidfluid systems. The activation energy was found to be 34.58 kJ/mol and 44.45 kJ/mol for the low and the high temperatures. The dissolution of gypsum was controlled by chemical reaction both low temperatures and high temperatures.

Keywords—Diammonium hydrogen phosphate, Dissolution, Gypsum, Kinetics.

I. INTRODUCTION

GYPSUM and anhydrite occur in arid and dry climatic conditions. It is located in layers. Gypsum layers take place as a result of evaporation of seas and lakes containing high salt. Gypsum is converted to anhydride in more depth. Natural anhydride is anhydrous calcium sulfate ($CaSO_4$).

Precipitation of gypsum and anhydrite depends on the water temperature and salinity. Anhydride occurs instead of gypsum under the conditions above these conditions. When anhydride formation is continued, the precipitation of NaCl begins. Therefore, in many gypsum and anhydrite beds are found some rock salt and iron oxides [1], [2].

Raw gypsum attends paper and cotton textile materials as white paint and filler. It is used to increase the ash content of coal dusts in coal minings. Crude gypsum is used in increasing amounts every day in the cement industry. It facilitates the melting of nickel smelting and is used for fermentation in beer industry [1], [2]. Gypsum and anhydrite in Turkey are available in many areas [3].

Studies done on the dissolution kinetics are summarized below. Reference [4] obtained boric acid and calcium phosphates from colemanite ore by using phosphoric acid. They determined the optimum conditions of dissolution. Reference [5] studied the dissolution of copper ore in aqueous NH₃-(NH₄)₂SO₄ system. They searched the effects of ammonia concentration, ammonium sulphate concentration, the reaction time and the solid-to-liquid ratio. They leached 98.87% of copper in ore. Reference [6] researched the kinetics and mechanism of the dissolution of sphalerite in sodium nitrate solutions with sulphuric acid. According to selected kinetics model, the dissolution of sphalerite controlled by the diffusion film ash. The activation energy was found as 55 kJ/mol. Reference [7] made a study on the dissolution kinetics of bastnaesite in hydrocloric acid solutions. The parameters effected the dissolution were hydrocloric acid concentration, the temperature, the solid-to-liquid ratio and the particle size. The dissolution of bastnaesite controlled by the diffusion film ash, and the activation energy was 59.39 kJ/mol. Reference [8] investigated the dissolution kinetics of Chevreul's salt in hydrocloric acid solutions. The dissolution of Chevreul's salt controlled by the diffusion film ash, and the activation energy was 57 kJ/mol. Reference [9] researched the removal of calcium from waste-waters containing calcium chloride. They were reacting the distilled liquid waste with gaseous reagents. As a result, they precipitated as gypsum (CaSO₄.2H₂O) the calcium. The parameters such as gaseous velocities, the reaction time and the stirring speed were taken as factors affecting the precipitation efficiency. The precipitation yield was %44 under these conditions.

II. EXPERIMENTAL PROCEDURE

A. Dissolution Tests

The dissolution experiments were performed in two different temperature groups. All of the experiments carried out in the ranges of the low temperatures and the high temperatures. First, the experiments were performed in the range of low temperatures. These temperatures was chosen as 25° C, 35° C, 45° C, 55° C. The total time of the experiments was taken as 150 minutes. Samples were taken at 30 min intervals. The water bath temperature was adjusted to 25° C. The prepared mixture was placed into the reactor, and stirring was performed by a magnetic stirrer. The stirring operation was continued throughout the reaction time. Withdrawing the sample from the solution every 30 minutes, the stirring was stopped. The sample was taken and filtered as 3 mL. After the samples are filtered, the amount of sulfate in solution were

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determined [10].

B. Dissolution Reactions

The reactions taking place in the solution can be written as:

$$CaSO_{4.2}H_2O_{(s)} \leftrightarrow Ca^{+2}_{(aq)} + SO_{4}^{-2}_{(aq)} + 2H_2O_{(l)}$$

$$\tag{1}$$

$$(NH_4)_2 HPO_4_{(aq)} \leftrightarrow 2NH_4^{+}_{(aq)} + HPO_4^{-2}_{(aq)}$$
(2)

$$CaHPO_{4.}2H_{2}O_{(s)} \rightarrow Ca^{+2}_{(aq)} + HPO_{4}^{-2}_{(aq)} + 2H_{2}O_{(l)}$$
(3)

$$(NH_4)_2 SO_{4(aq)} \rightarrow 2NH_4^+_{(aq)} + SO_4^{-2}_{(aq)}$$
(4)

When gypsum is added to the diammonium hydrogen phosphate solutions, the total reaction taking place in the solution can be written as:

$$\begin{array}{l} CaSO_{4.}2H_{2}O_{(s)}+(NH_{4})_{2}HPO_{4(aq)} \rightarrow CaHPO_{4.}2H_{2}O_{(s)} \\ +(NH_{4})_{2.}SO_{4(aq)} \end{array} \tag{5}$$

The parameters and their values used in the experiments are given in Table I.

	TABLE I
PARAMETERS AND THEIR VALUES	
Parameters	Levels of Parameters
Temperature (°C)	25 35 45 55 65 75 80 85 90
Reaction time (min.)	30 60 90 120 150
Solid/liquid ratio (g.mL ⁻¹)	1/100
Stirring speed (rpm)	650
Concentration (M)	0.5 1 2 3
Particle size (mm)	0.03482

*While the effect of one parameter was studied, the values of the other parameters were kept constant.



III. RESULTS AND DISCUSSION

Fig. 1 Variation of $1-(1-X)^{(1/3)}$ with time for the low temperatures (25-55°C)

In this study, pre-experiments were performed first. In light of these pre-experiments, the parameters affecting the dissolution kinetics of gypsum in diammonium hydrogen phosphate solutions were chosen as the temperature, the reaction time, the stirring rate and the solution concentration. According to the test results, graphs for 3 model were drawn and were calculated R2 values in relation to these models. It was found that the most appropriate model is the surface chemical reaction controlled model. This situation was shown in Fig. 1.



Fig. 2. Variation of 1-(1-X)^(1\3) with time for the high temperatures $(75-90^{\circ}C)$



Fig. 3 Effect of the reaction temperature on the dissolution of gypsum

According to the test results, graphs for 3 models were drawn and were calculated R^2 values in relation to these models. It was found that the most appropriate model is the

surface chemical reaction controlled model. This situation was shown in Fig. 2.

The temperature is a factor of great importance for the leaching kinetics. The effect of reaction temperature was examined at 25, 35, 45, 55, 75, 80, 85, 90°C. The fact that the activation energies are different can probably be attributed to the differences of the chemical reaction mechanism. The effect of reaction temperature was investigated for the temperature range 25-55°C and 75-90°C temperatures. The dissolution curves obtained for 20-55°C are shown in Fig. 3 and 75-90°C are shown in Fig. 3 and Fig. 4 that, the quantity of gypsum dissolved increases with increasing of reaction temperature.





Fig. 5 Effect of the diammonium hydrogen phosphate concentration on the dissolution of gypsum at the low temperature

In general, the dissolution rate increases with increasing of concentration of reagent, but only up to a certain maximum level. The effects of concentration of diammonium hydrogen phosphate solutions was studied by varying to 0.5, 1.0, 2.0 and 3.0 M. The dissolution curves are given in Figs. 5-6. It can be seen from the Figs. 5 and 6 that the dissolution level of the process increases with increasing in the concentration of diammonium hydrogen phosphate solutions.







Fig. 7 Arrhenius plot of the dissolution process for the low temperatures

The Arrhenius plots of ln k versus 1/T were drawn for to found the activation energy of the reaction. Arrhenius plot of the dissolution process for reaction temperatures are shown in Figs. 7, 8. From the slopes of the straight lines of Figs. 7, 8, the activation energy of the reaction is found to be 34.58 kJ/mol and 44.45 kJ/mol. These values indicate the dissolution rate of gypsum is a chemical reaction controlled [8], [11]-[17].



Fig. 8 Arrhenius plot of the dissolution process for the high temperatures

IV. CONCLUSIONS

The aim of the study was to investigate the dissolution kinetics of gypsum in diammonium hydrogen phosphate solutions in a mechanical agitation system. Based on the results obtained in this research, the following conclusions may be drawn:

The dissolution rate of gypsum increased with increasing in the low and the high reaction temperatures.

The dissolution rate of gypsum increased with increasing of diammonium hydrogen phosphate concentrations in the low and the high reaction temperatures.

The dissolution process follows a shrinking core model with the heterogeneous chemical reaction as the rate controlling step.

The activation energy of the reaction is found to be 34.58 kJ/mol and 44.45 kJ/mol for the low and the high temperatures.

Reaction rate expressions of dissolution of gypsum at the low temperatures and the high temperatures controlled by chemical reaction are as follows, respectively.

$$\left[1 - \left(1 - X_B\right)^{1/3}\right] = k_{1.}e^{-5159.5/T}.t$$
(6)

$$\left[1 - \left(1 - X_B\right)^{1/3}\right] = k_{2.}e^{-5346.8/T}.t$$
(7)

where k_1 and k_2 are constants depending on the diammonium hydrogen phosphate solution concentration, the solid/liquid ratio, the stirring speed and the particle size.

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REFERENCES

- Anonim, "Türkiye'nin Doğal Kaynakları" http://www.marbleport.com. pp. 1-10.
- [2] Anonim, "Jips Bugünü ve Geleceği, Metal Teknoloji Platformu Oluşturma Çalıştayı", Tübitak, Ankara, 2003, pp. 1-74.
- [3] Anonim, "Türkiye Sanayi Sektörünün Değerlendirilmesi", Türkiye Cumhuriyeti Bilim Sanayi ve Teknoloji Bakanlığı, Ankara, 2011, pp. 1-55.
- [4] M. Yeşilyurt, S. Çolak, T. Çalban and Y. Genel, "Determination of the optimum conditions for the dissolution of colemanite in H3PO4 solutions", Ind. Eng. Chem. Res., vol.44, 2005, pp. 3761-3765.
- [5] T. Çalban, S. Çolak and M. Yeşilyurt, "Optimization of Leaching of Copper from Oxidized Copper Ore in NH3-(NH4)2SO4 Medium", Chem. Eng. Comm., vol. 192, 2005, pp. 1515-1524.
- [6] M. Sokic, B. Markovic, V. Matkovic, D. Zivkovic, N. Strbac and J. Stojanovic, "Kinetics and Mechanism of Sphalerite Leaching by Sodium Nitrate in Sulphuric Acid Solution", Journal of Mining and Metallurgy, vol. 48 (2), 2012, pp. 185-195.
- [7] B. Xue, Y. Shao-hua, L. Yao and W. Wen-yuan, "Leaching Kinetics of Bastnaesite Concentrate in HCl Solution", Trans. Nonferrous Met. Soc. of China, vol. 21, 2011, pp. 2306-2310.
- [8] T. Çalban, B. Kaynarca, S. Kuşlu, S. Çolak, "Leaching kinetics of Chevreul's salt in hydrochloric acid solutions", Journal of Ind. and Eng. Chem., vol. 20, 2014, pp. 1141-1147.
- [9] T. Çalban, E. Kavcı, "Removal of Calcium from Soda Liquid Waste Containing Calcium Chloride", Energy Sources: Part A, vol. 32, 2010, pp. 407-418.
- [10] H. Gülensoy, "Kompleksometrinin Temelleri ve Kompleksometrik Titrasyonlar", Fatih Yayınevi, İstanbul, 1984.
- [11] E. Kavcı, T. Çalban, S. Kuşlu, S. Çolak, "Leaching kinetics of ulexite in sodium hydrogen sulphate solutions", Journal of Industrial an Engineering Chemistry, vol. 20, 2014, 2625.
- [12] O. Levenspiel, "Chemical Reaction Engineering", Wiley, 2nd edition: NewYork, 1972.
- [13] F. Habashi, "Kinetics of Metallurgical Processes", 1999.
- [14] A. Ekmekyapar, E. Aktaş, A. Künkül, N. Demirkıran, "Investigation of leaching kinetics of copper from malachite ore in ammonium nitrate solutions", Metallurgical and Materials Transactions B, vol. 43B, 2012, p. 764.
- [15] B. Dönmez, F. Demir, O. Lacin, "Leaching kinetics of calcined magnesite in acetic acid solutions", Journal of Industrial an Engineering Chemistry, vol. 15, 2009, p. 865.
- [16] S. Kuşlu, F. Ç. Dişli, S. Çolak, "Leaching kinetics of ulexite in borax pentahydrate solutions saturated with carbon dioxide", Journal of Industrial an Engineering Chemistry, vol. 16, 2010, p. 673.
- [17] R. Guliyev, S. Kuşlu, T. Çalban, S. Çolak, "Leaching kinetics of colemanite in potassium hydrogen sulphate solutions", Journal of Industrial and Engineering Chemistry, vol. 18, 2012, p. 38.