

# Hydrogen Bonding Adsorption Principle in Caustic Solution of Aluminium Hydroxide

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## Abstract

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In this paper, the concept of hydrogen bonding adsorption was investigated for  $\text{Al}(\text{OH})_3\text{-NaOH-H}_2\text{O-S}$  (solid) system. Aluminium hydroxide transfer between solid and liquid phase by hydrogen bonding adsorption and desorption, only with hydrogen bond breaking or formation of aluminium hydroxide crystal. When the  $\text{Al}(\text{OH})_3\text{-NaOH-H}_2\text{O-S}$  (solid) system achieves equilibrium, this is due to hydrogen bonding adsorption. Equilibrium equation of the system was obtained,  $\text{CE} = 90 \text{ S}^{-0.1} \text{ N}^{1.5} \cdot \exp(-18522.9/(\text{R} \cdot \text{T}))$ . No chemical reaction occurs when aluminium hydroxide dissolves in caustic solution process or crystallises in a seeded precipitation process. In the liquid phase, alumina is in the form of  $\text{Al}(\text{OH})_3$ , rather than  $\text{Al}(\text{OH})_4^-$ . In this concept, supersaturation does not exist in sodium aluminate solution.

**Keywords:** Bayer process, Solubility, Hydrogen bonding, Phase equilibrium, Seed precipitation.

## 1. Introduction

Bayer process was born more than one hundred years, The basic principle of Bayer process is established on the  $\text{Na}_2\text{O-Al}_2\text{O}_3\text{-H}_2\text{O}$  system, according to the generally accepted theory, alumina exists in form of  $\text{Al}(\text{OH})_4^-$  or two poly-aluminate ions in caustic solution. Alumina concentration in caustic solution depends on the solubility, where seeded precipitation process is a chemical reaction, which eventually stops when alumina solubility is reached. However, according to this theory, the super saturation phenomenon is difficult to explain, even it is thought that spent liquid is supersaturated. Researchers have attributed this to the complex structure of  $\text{Al}(\text{OH})_4^-$  in caustic solution, still there is no reasonable results supporting this up to now.

Researching on the basic principles of the Bayer process is important to the alumina industry, also it is of great significance to improve the quality of the products. In this paper, the principle of hydrogen bonding adsorption was adopted on  $\text{Al}(\text{OH})_3\text{-NaOH-H}_2\text{O-S}$  (solid) system.

## 2. Experiment

A series of experiments were conducted to study various aspects of the crystallisation phenomena, such as the effect of temperature, solid concentration and caustic concentration.

### 2.1 Experimental Materials

Aluminium hydroxide: grade AH-1 GB/T 4294-2010, specific surface area:  $0.104 \text{ m}^2/\text{g}$ . Caustic: chemical pure.

**Table 1. Size distribution of aluminium hydroxide.**

Size, $\mu\text{m}$	10	15	20	30	40	50	60	80	120	150	200
%	0.91	3.53	5.80	8.29	11.16	16.69	24.83	44.57	77.91	91.18	99.22

## 2.2 Experimental Instrument and Analysis

A 5 L stainless steel container was used, with stirring, automatic temperature control of  $\pm 0.1$  °C. Laser particle size was analyzed using a Mastersizer 2000 instrument.

## 2.3 Experimental Operation

### 2.3.1 The Relationship Between the End Alumina Concentration and Temperature

Sodium aluminate solution 5 L, caustic concentration  $\text{Na}_2\text{O}$  100 g/L, alumina concentration  $\text{Al}_2\text{O}_3$  80 g/L, heated to 60 °C. Dried aluminium hydroxide 3 kg was added to the sodium aluminate solution, stirring. After 50 hours, the slurry was separated, analysis alumina concentration, sampling once every 5 hours. Three of difference of alumina concentration is less than 0.5 g/L, that reach the end, three results mean as the end of alumina concentration. Then cooling to 55, 50 and 45 °C, repeated sampling.

### 2.3.2 Relationship Between the End Alumina Concentration and the Amount of Solid Aluminium Hydroxide

Sodium aluminate solution 5 L, caustic concentration  $\text{Na}_2\text{O}$  100 g/L, alumina concentration,  $\text{Al}_2\text{O}_3$  80g/L, temperature 50 °C. Respectively, adding dried aluminium hydroxide 2, 3, 4 and 5 kg. Sampling was the same as described in 2.3.1.

### 2.3.3 The Relationship between the End Alumina Concentration and Caustic Concentration

Sodium aluminate solution 5 L, caustic concentration  $\text{Na}_2\text{O}$  120, 110, 100, 90 g/L, alumina concentration  $\text{Al}_2\text{O}_3$  90 g/L, temperature: 50 °C. 3 kg dried aluminium hydroxide was added to the sodium aluminate solution, stirring. Sampling was the same as described in 2.3.1.

## 3. Results and Discussion

Alumina concentration at the end under the different conditions are shown in Table 2.

**Table 2. Alumina concentration at the end under different conditions.**

No.	N, g/L	W, g/L	T, °C	C, g/L
1	100	600	60	74.2
2	100	600	55	67.5
3	100	600	50	60.6
4	100	600	45	54.5
5	100	1000	50	57.3
6	100	800	50	58.6
7	100	400	50	62.2
8	120	600	50	79.1
9	110	600	50	69.3
10	90	600	50	51.2

N: caustic ( $\text{Na}_2\text{O}$ ) concentration; W: solid content; T: temperature; C: end alumina ( $\text{Al}_2\text{O}_3$ ) concentration,

## 6. Reference

1. Li Xiao-bin et al., Research progress in theory and technology of gibbsite precipitation from sodium aluminate solution, *Chinese Journal of Nonferrous Metals*, 2011, Vol.21, No10
2. Meng Ling-li, Study on dissolution kinetics of the aluminium hydroxide in the caustic soda, *Light Metals*, 2012, Vol.9 (in Chinese).
3. Huang Zhen-qian and Hu Nan-yuan, *Physical Chemistry*. Changsha: Central South University Press, 1991:258-285
4. Bi Shi-wen and Yu Hai-yan, *Alumina production process*, Beijing: Chemical Industry Press, 2006, 78-90
5. Zhang Ying et al., Preparation and Characterization of  $\text{Al}(\text{OH})_3$  Coarse Particles by Decomposition of Caustic Aluminate Solution with Methanol, *The Chinese Journal of Process Engineering*, 2009, Vol.9, 300-305
6. S. Scott Fogler, *Elements of Chemical Reaction Engineering*, Beijing: Chemical Industry Press, 2006, 80-141
7. C. Misra and E.T. White, *Chemical Engineering Progress Symposium*, 1970, 67(11):53
8. C. Misra and E.T. White, *Journal of Crystal Growth*, 1971, 8, 172
9. Bi Shi-wen, Xue Hong and Yang Yi-hong, Kinetics of decomposition of Bayer sodium aluminate solution, *The Chinese Journal of Nonferrous Metals*, 1998, 8(1), 131-134
10. Li Xiao-bin et al., Kinetics of carbonation decomposition of sodium aluminate solution. *Chinese Journal of Nonferrous Metals*, 2004, 14(5):850-853
11. Li Xiao-bin et al., Thermodynamics of carbonization of aluminate solution, *Chinese Journal of Nonferrous Metals*, 2003, 13(4):1005-1010
12. Fang Jing-dong, Wu Su-fang and Wang Zhang-mao, The studies of decomposition reaction of sodium aluminate solution, *Journal of Chemical Engineering of Chinese Universities*, 2002, 16(1):33- 37
13. I. V. Davidov et al., Optimization of temperature condition of the liquor decomposition process, *Light Metals*, 1976, 1:173
14. S. Veessler and R. Boistelle, Growth kinetics of hydrargillite  $\text{Al}(\text{OH})_3$  from caustic soda solution, *Crystal Growth*, 1994, 142:177
15. Li Jie, Study on the structure character and decomposition mechanisms of supersaturation of sodium aluminate solutions, *Changsha, Central South University*, 2002
16. P. Smith and G. Woods, The measurement of very low growth rates during the induction period in aluminum hydroxide growth from Bayer liquors, *TMS Light Metals*, 1993:113-117
17. C. Misra, Solubility of aluminum hydroxide (hydrargillite) in sodium hydroxide solutions, *Chemistry and Industry*, 1970, (20):157
18. Li Jia-zheng, Sun Jiang-xiao and Shi Zuo-qing, Hydrogen Binding Adsorption and its Prospect, *Ion Exchange and Adsorption*, 2001, 17(6):561-566
19. Huang Jian-gou et al., A thermodynamic study on adsorption of phenols onto D301 resin. *Ion Exchange and Adsorption*, 2003, 19(1):37-42
20. Chen Xiao-hu et al., The mathematic model of the pregnant aluminate precipitation, *Journal of Guizhou University Technology*, 2002, 31(5):22-24