# Drying Process Optimization and Efficiency of Aluminum Fluoride (AlF<sub>3</sub>) Plant

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*Abstract* – Aluminum Fluoride (AlF<sub>3</sub>) is a substantive material used to reduce the melting point temperature of the Aluminum from 1500°C to 600-850°C. The production applies a wet process using raw materials of Fluosilicic Acid and Aluminum Hydroxide with a by-product of Silica Dioxide. In this case, AlF<sub>3</sub> products are expected to have a maximum H<sub>2</sub>O content of 2.6%. However, in the production process, the H<sub>2</sub>O content reached 3% due to the non-optimum drying process. Therefore, the optimization process was carried out by increasing the dry air temperature of the recycle from stage 2 to stage 1 in the drying process, so that the mass transfer of H<sub>2</sub>O from AlF<sub>3</sub> crystals to the air can be maximized. After the optimization, it was found that the H<sub>2</sub>O content has met the quality standard of 1.8%.

Keywords – Aluminum fluoride, Fluosilicic acid, Aluminum hydroxide, Drying process

## I. INTRODUCTION

urrently, the Indonesian government pays more attention to the development and expansion of the metal industry sector in Indonesia to meet both domestic and foreign needs to strengthen and stabilize the economic growth. However, the management of the metal industry has not yet achieved independence, one of which is the aluminum refining industry. The demand for aluminum for the domestic industry currently reaches 600,000-800,000 tons per year, but the amount that can be met by PT Y, the largest Aluminum industry in Asia, is only around 104,000 tons per year. It is estimated that by 2025, the demand will reach 2 million tons/year, especially since PT Y is preparing to increase its production capacity to 400,000 tons/year [1].

It is known that the Aluminum demand is quite high, so that, the fulfillment of aluminum needs must be carried out [2]. However, in the production process, aluminum requires a high melting temperature of around 1200-1500°C. This has an impact on energy consumption and large costs. Therefore, a substantive material is needed to reduce the melting point temperature to around 600-850°C with the addition of AlF<sub>3</sub> compounds so that it can reduce the production costs and the production process runs more efficiently [3]. However, to meet this high demand, only PT X is currently operating to produce Aluminum Fluoride using wet process with Fluosilicic Acid and Aluminium Hydroxide as raw material. The specifications of each raw material can be seen in Table 1. and Table 2.

Optimization and efficiency process at the Aluminum Fluoride plant was carried out, so that, it is expected the production can run optimally and becomes one of the sectors that can boost the development of Indonesian industry, especially in the metal industry sector. In this case, the optimization and efficiency processes were carried out by evaluating the production process that has been running at the Aluminum Fluoride factory of PT X, where the data were collected.

TABLE 1	•
SPECIFICATION OF ALUM	INUM HYDROXIDE
Parameter	Information
Molecular Formula	Al(OH) <sub>3</sub>
Molecular Weight (g/mol)	78
form	Solid, white
Melting Point (°C)	300
Bulk Density (kg/m <sup>3</sup> )	1150
Al(OH) <sub>3</sub> content	Minimum of 98.5%
SiO <sub>2</sub> level	Maximum of 1.5%

TABLE 2.

SPECIFICATIONS OF FLUOSILICIC ACID				
Parameter	Information			
Molecular Formula	$H_2SiF_6$			
Molecular Weight (g/mol)	144.08			
form	Liquid, colorless			
Melting Point (°C)	-30			
Specific Gravity	1.15			
$H_2SiF_6$ . levels	Minimum of 18%			
P <sub>2</sub> O <sub>5</sub> level	Minimum of 0.025%			
H <sub>2</sub> O level	Maximum of 82%			

#### II. OPTIMIZATION METHOD

This optimization process was carried out to increase the level of  $H_2O$  released from AlF<sub>3</sub>.3H<sub>2</sub>O crystals by comparing the calculation of the mass balance and energy balance before and after the optimization process, so that the H<sub>2</sub>O content in the product is in accordance with the predetermined specifications. This drying process consisted of 2 stages. The operating temperature of stage 1 is 200°C, while stage 2 is 500°C. This operating temperature condition produced a product with an H<sub>2</sub>O content of 3%. This optimization process was carried out by increasing the recycle temperature from stage 2 to



Figure 1. Location of Aluminum Fluoride Factory

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 $250^{\circ}$ C using steam in the heater so that more H<sub>2</sub>O was released at stage 1. Therefore, the mass of H<sub>2</sub>O released from AlF<sub>3</sub>.3H<sub>2</sub>O crystals can increase. This further lightened the drying load on stage 2. Furthermore, the recycle air temperature was increased to 250°C so that the output temperature of AlF<sub>3</sub>.0.5H<sub>2</sub>O became 300°C.

## III. RESULTS AND DISCUSSION

## **Process existing**

## 3.1 Process Description

In the production process, Aluminum Fluoride is made using raw materials of Fluosilicic Acid and Aluminum Hydroxide. The specifications for the Aluminum Fluoride product can be seen in Table 3 according to the specification that defined by PT X, refers to SNI 06-2603-1992. In this case, the quality of the product must be maintained. From the PT. X data [3], it is known that there was a discrepancy between the H<sub>2</sub>O content of the product and the qualified specifications. Based on the material balance calculation, there was also an imbalance between the production capacity of AlF<sub>3</sub> which should be 1708.3 kg/hour but the existing process produce 1728.3 kg/hour capacity.

The process of making Aluminum Fluoride in this factory applies a wet process which consists of the following several stages:

- 1) Preparation Unit
- 2) Reaction Unit
- 3) Separation of SiO<sub>2</sub> Unit
- 4) Crystallization Unit
- 5) Separation of AlF<sub>3</sub>.3H<sub>2</sub>O Unit
- 6) Drying Unit
- 7) Cooling and Packaging Unit

In general, the process in the manufacture of Aluminum Fluoride is shown in Figure 2.

## A. Preparation Unit

 $H_2SiF_6$  from the Phosphoric Acid Factory from PT X is stored in storage tanks. Then, it is pumped to the second storage tank (preheated tank). In this second storage tank,  $H_2SiF_6$  is preheated in a Heat Exchanger before being reacted with Al(OH)<sub>3</sub>. The temperature in the Heat Exchanger is further maintained in the range of 78°C-80°C. When the temperature of  $H_2SiF_6$  has reached 75°C-78°C, it will be sent to the reactor. However, if the temperature has not reached 75°C, it will be returned to the preheated tank to be flowed back to the Heat Exchanger so that the temperature before the reaction is reached. Al(OH)<sub>3</sub> from the storage warehouse with a maximum  $H_2O$  content of

	TABLE 3.
Specific <sub>A</sub>	TION OF ALUMINUM FLUORIDE

Parameter	Information
Molecular Formula	AlF <sub>3</sub>
Molecular Weight (g/mol)	83.98
form	Powder, white
Melting Point (°C)	1290
Bulk Density (kg/m <sup>3</sup> )	770
AlF <sub>3</sub> levels	Minimum of 97%
SiO <sub>2</sub> level	Minimum of 0.2%
P <sub>2</sub> O <sub>5</sub> level	Minimum of 0.02%
Fe <sub>2</sub> O <sub>3</sub> content	Maximum of 0.07%
H <sub>2</sub> O level	Maximum of 2.8%

1.3% is then sent to the  $Al(OH)_3$  Silo using a bucket elevator. To avoid dust from escaping from the silo, the equipment is installed with an exhaust fan and filtered by a filter from the  $Al(OH)_3$  silo.

## B. Reaction Unit

The reaction between  $H_2SiF_6$  and  $Al(OH)_3$  occurs in the reactor at a temperature of 90°C with a maximum temperature of 99°C for 13 minutes. The reactor is equipped with an agitator so that there is no precipitation from the remaining of the reaction products. The product formed of the reaction is exothermic. The amount of  $Al(OH)_3$  required for the reaction is calculated according to the concentration of  $H_2SiF_6$  entering the reactor. The following reaction occur in the reactor:

$$H_2SiF_{6(l)}+2Al(OH)_{3(s)} \rightarrow 2AlF_{3(aq)}+SiO_{2(s)}+4H_2O_{(l)}$$



Figure 2. Block Diagram of Aluminum Fluoride Manufacturing Process using Wet Process



**Figure 3.** Flowchart of AlF<sub>3</sub> Production Process

#### C. Separation of $SiO_2$ Unit

In order to obtain good product quality, the silica contained in the reaction product must be separated immediately and the moisture content of the silica must also be decreased. After the reaction, it flows into the centrifuge. In the centrifuge, AlF<sub>3</sub> will be separated from SiO<sub>2</sub>. This separation is done based on the density. There is a significant difference in density between AlF<sub>3</sub>, H<sub>2</sub>O, and SiO<sub>2</sub>, where the highest density is in SiO<sub>2</sub> which is 2,650 kg/m<sup>3</sup>, while the density of AlF<sub>3</sub> and H<sub>2</sub>O is 770 and 1000 kg/m<sup>3</sup>.

# D. Crystallization Unit

In the crystallization process, the slurry from the centrifuge will be crystallized. The crystallizer is a batch that lasts for 4 hours so that 4 crystallizers are used in parallel. Before entering the crystallizer, the slurry will be accommodated in the AlF<sub>3</sub> distributor to be flowed into the crystallizer through the rotary valve. The function of the AlF<sub>3</sub> distributor is to regulate the purpose of the slurry flow so that there is no overflow/spill on the crystallizer. In this process, the slurry is in direct contact with the steam from the bottom of the crystallizer. The steam used is saturated steam (wet steam). With increasing solid porosity, it is expected that the product will have low bulk density. In addition, the crystallizer is equipped with an agitator or stirrer so that the product does not harden during heating. Based on the solubility curve [4], at a temperature of 90°C the concentration crystals formed during crystallization is 140 mol/m<sup>3</sup>. After the process is complete, it will be put in a holding tank.

## E. Separation of AlF<sub>3</sub>.3H<sub>2</sub>O Unit

This process aims to separate  $AlF_3.3H_2O$  using saturated solution (mother liquor) which will later be recycled back to the crystallizer. The separation is carried out using a centrifugal type centrifuge so that the separation is based on density. Solid  $AlF_3.3H_2O$  will be separated from mother liquor at the operating temperature of 80°C. Mother liquor contains unreacted raw material and some uncrystallized  $AlF_3$ . Mother liquor will be further put in the vessel since some of it will be recycled to the crystallizer, while the other part is purged to waste water treatment. Recycle serves as an initiator of crystal formation in the crystallizer. The separated crystals will enter the hopper and then flow to the drying stage.

# F. Drying Unit

In this process, the reduction of water content in AlF<sub>3</sub> is divided into 2 stages [5], namely the first stage is to reduce the water content of AlF<sub>3</sub>.3H<sub>2</sub>O to AlF<sub>3</sub>. H<sub>2</sub>O and the second stage is to reduce the water content in AlF<sub>3</sub>. H<sub>2</sub>O to become AlF<sub>3</sub>. This process is carried out on a rotary dryer. At this stage, the water content of the AlF<sub>3</sub> crystals will also be reduced in both free and hydrate forms to obtain the desired product specifications. The moisture content will be reduced to a maximum of 2.6%w. This process uses hot dry water with a temperature of 600°C which is heated using a burner as a water vapor carrier. The operating condition temperature in the first stage is 250°C, while in the second stage is 600°C. This drying process takes place in direct contact with countercurrent flow in the rotary dryer.

$$AlF_{3.}3H_{2}O \rightarrow AlF_{3.} 0.5 H_{2}O + 2.5 H_{2}O$$
$$AlF_{3.} 0.5 H_{2}O \rightarrow AlF_{3} + 0.5 H_{2}O$$

## G. Cooling and Packing Unit

The AlF<sub>3</sub> solid that comes out of the rotary dryer has a temperature of about 500°C. So it must be refrigerated before the packaging. Meanwhile, the cooling process uses a rotary cooler type, so that the cooling water does not directly contact the product. The product temperature after cooling process is 40°C. The product is packaged in sacks/bags for 1 ton. Therefore, a weight tool is used in the packaging process.

# 3.2 Material Balance

Based on the calculations results from the material balance of the Aluminum Fluoride plant, it was found that in order to obtain a production capacity of 12,600 tons/year (1708.33 kg/hour), it requires  $H_2SiF_6$  and AlOH<sub>3</sub> of 8346.588 kg/hour and 1652.37 kg/hour, respectively in each production [6][7].

## **Process Optimization**

# 3.3 Optimization

Product quality is important in a production process. AlF<sub>3</sub> products produced at this factory are expected to be in accordance with the specifications set by the factory, which include having a purity of minimum 94%, SiO<sub>2</sub> of maximum 0.2%, H<sub>2</sub>O of maximum 0.26%, and LOI of maximum 0.85%. In practice, there are still problems related to product quality, in which the water content in AlF<sub>3</sub> products exceeds the predetermined standard (3% of H<sub>2</sub>O). This mismatch of water content can be caused by the non-optimal drying process in the rotary dryer. In the production process, anhydrous AlF<sub>3</sub> compound is obtained from its hydrate form in the form of AlF<sub>3.3</sub>H<sub>2</sub>O



Figure 4. Process Flow Diagram of Drying Unit Before (left) and After (right) Optimization

(trihydrate) by going through a drying process. The drying process is carried out by direct contact with AlF<sub>3</sub>.3H<sub>2</sub>O and dry air using a counter current flow type.

It is known that there is a mismatch between the  $H_2O$  content in the manufactured AlF<sub>3</sub> product and the product quality standards. Figure 3. shows a flow diagram of the production process where there is a drying unit. The mismatch of  $H_2O$  levels in the product can be caused by the non-optimal release of  $H_2O$  from AlF<sub>3</sub>.3H<sub>2</sub>O crystals in the rotary dryer during the drying process. Therefore, it is necessary to optimize the drying process to obtain  $H_2O$  content in accordance with the product specifications.

Based on the data from PT X's Aluminum Fluoride unit [3], it is known that there were problems that occur, namely the quality of the product where the H<sub>2</sub>O content is not in accordance with the product specifications. In this case, it was found that the H<sub>2</sub>O content in the product reaches 3%. This can be caused by the drying process, which is not optimal, so that there is H<sub>2</sub>O content in the AlF<sub>3</sub> crystals that is not evaporated. The calculation of material balance can be seen in Figure 5.

The dry air temperature condition that is fed to the rotary dryer after being heated in the burner is 500°C (stage 2) and the recycled dry air resulting from stage 2 is fed to stage 1 with a temperature of 200°C. At these operating temperature conditions, the total mass flow of the product obtained was 1728.83 kg/hour, which indicates that it exceeded the expected capacity of 1708.33 kg/hour (based on the calculation of the production capacity of 41

tons/day). This occurred due to the presence of excess water content in the product. The mass of  $H_2O$  that is evaporated from AlF<sub>3</sub>.3H<sub>2</sub>O was 1169.21 kg/hour, which requires 6507.5 kg/hour of dry air to evaporate the H<sub>2</sub>O.

Based on Figure 6. it can be seen that at stage 1, streams 11, 11a, 15, and 16 are AlF<sub>3</sub>.3H<sub>2</sub>O crystals, AlF<sub>3</sub>.0.5H<sub>2</sub>O crystals, recycle air, and exhaust air, respectively. In stage 1 with an operating temperature of  $250^{\circ}$ C after the optimization (increasing temperature using heater [8]), the mass of H<sub>2</sub>O removed was 897.159 kg/hour. In stage 2, the mass of H<sub>2</sub>O removed was 291.51 kg/hour. With a stage 1 operating temperature of  $250^{\circ}$ C, dry air mass of 5536.47 kg/hour is required with a humidity of 0.0011 kg H<sub>2</sub>O/kg Dry Air, it can reduce the H<sub>2</sub>O content of 1188.21 kg/hour so that the H<sub>2</sub>O content in the product is 30.75 kg/hour



Figure 5. Mass Balance of Stage 1 and 2 Rotary Dryer Before Optimization (Problem Condition)

<11> 	st	AGE 1	<11a> → <15>	<11a> 	STA	GE 2	<17> 
Input (k	g/jam)	Output (kg/jam)		Input Output		put	
Stream	<11>	Stream <	11a>	Stream <	11a>	Stream	n <17>
A1F3	10,515	A1F3	10,515	A1F3	10,515	A1F3	1674,17
A1F3.3H2O	2733,14	A1F3.0.5H2O	1841,9	A1F3.0.5H2O	1841,9	H <sub>2</sub> O	30,750
H <sub>2</sub> O	149,928	H <sub>2</sub> O	144,01	H <sub>2</sub> O	144,01		
Stream <15>		Stream <16>		Stream <13> Stream <15>		n <15>	
Dry Air	5536,5	Dry Air	5536,5	Dry Air	5536,5	Dry Air	5536,5
H <sub>2</sub> O	1188,7	H <sub>2</sub> O	1194,6	H <sub>2</sub> O	5,92	H <sub>2</sub> O	1188,7
Humidity	0,215	Humidity	0,216	Humidity	0,011	Humidity	0,215
$H_2C$	) removed :	= 897,159 kg/jar	n	H2O re	moved = 2	.91,509 kg/ja	am
Total	9622.14	Total	9622.14	Total	8433.47	Total	8433.47

Figure 6. Mass Balance Stage 1 and 2 Rotary Dryer Expected Condition (After Optimization)

(1.8%wt; specification standard). This data is obtained based on material and energy balance calculation [9][10].

## 3.4 Effect of the Optimization Process

The optimization process carried out affected both the production process and the economic analysis. After the optimization, the final product of  $AIF_3$  has met the standard and material balance can be achieved at a production capacity of 1708.3 kg/hour. Therefore, based on the evaluation and calculations carried out, a comparison of conditions before and after the optimization can be obtained, both in terms of the production process (material balance) and evaluation of the economic analysis as shown in Table 4.

 TABLE 4.

 COMPARISON OF CONDITIONS BEFORE AND AFTER OPTIMIZATION

Parameter	Before Optimization	After Optimization	Unit
Mass Flow	1728.3	1708.33	kg/hour
%H <sub>2</sub> O at <17>	3	1.8	%wt
H <sub>2</sub> O mass <17>	51.25	30.75	kg/hour
Mass of AlF <sub>3</sub> <17>	1674.17	1674.17	kg/hour
Mass of SiO <sub>2</sub> <17>	3.4167	3.4167	kg/hour
Dry Air Mass	6507.5	5536.47	kg/hour
Humidity<13>	0.00107	0.00107	kg
Humidity<15>	0.1797	0.215	H <sub>2</sub> O/kg
Humidity<16>	0.181	0.216	Dry Air
H <sub>2</sub> O Removed	1169.21	1188.21	kg/hour
IRR	37.82	36.97	%
POT	3.29	3.36	year
BEP	34	35	%

## 3.5 Economic Analysis

Economic analysis was carried out to determine the feasibility of a factory. In the case of this economic analysis an evaluation was carried out by considering several things as follows:

- 1) Internal Rate of Return (IRR)
- 2) Payout Time (POT)
- 3) Break Event Points (BEP)

The calculation of the economic analysis was carried out using the discounted cash flow method, in which the cash flow value was projected at the present time. The followings are the assumptions used in the calculation.

- 1) Capital, consists of 30% of own capital and 70% of loan capital
- 2) Bank interest is 10% per year
- 3) Inflation rate is 3.5% per year
- 4) The construction period is 2 years, with the first year using 50% of own capital and 50% of loan capital. As for the second year using the remaining capital owned.
- 5) Loan repayment is within 10 years.
- 6) Production capacity at
  - a) First year = 60%
  - b) Second year = 80%
  - c) Third year
- 7) Income tax of:
  - a) Up to IDR 50,000,000 = 10%
  - b) IDR 50,000,000 IDR 100,000,000 = 15%

= 100%

c) More than IDR 100,000,000 = 30%

Based on these data, calculations were made to obtain the IRR, POT, and BEP values. IRR is defined as a certain interest rate on conditions when all revenues will still cover the entire amount of capital issued. Based on the calculation, the IRR value is 37.82%. The IRR value obtained is greater than the interest from bank loans. Furthermore, the POT value is known through the calculation of accumulated cashflow and the payback period, which is 3.29 years (before tax) and 4.06 years (after tax).

BEP is the break-even point when the total production costs are equal to the sales results, so that the total production capacity at that point can be known. BEP calculation involves several parameters including Fixed Cost (FC), Semi Variable Cost (SVC), Variable Cost (VC), and Total Sales (S). Based on the calculation, it can be obtained that the BEP value is 34% at a production capacity of 432,159,814 kg/year.

The optimization process on the drying unit has an influence on the economic analysis, including the IRR, POT, and BEP values. The optimization process was carried out by adding a heater to the drying unit to increase the recycle temperature from stage 2 so that it will reduce

 TABLE 5.

 Fix Cost, Variable Cost, Semi Variable Cost, and Total Sales

No	Information	Amount (Rp)	
1	Fix Cost (FC)	IDR3,286	
	- Depreciation	IDR2,629	
	<ul> <li>Property tax</li> </ul>	IDR394	
	- Insurance	IDR263	
2	Variable Cots (VC)	IDR5,892	
	- Raw material	IDR4,450	
	- Utilities	IDR1,258	
	- Royalties	IDR184	
3	Semi Variable Cost (SVC)	IDR9,225	
	<ul> <li>Employee salary</li> </ul>	IDR628	
	- Supervision	IDR94	
	- Maintenance & Repair	IDR1,840	
	- Operating Supplies	IDR276	
	- Laboratory	IDR276	
	- General Expenses	IDR4,318	
	- Plant Overhead Cost	IDR1,793	
4	Total Sales (S)	IDR30,000	

the IRR value to 36.97%. The payback period (POT) increased to 3.36 years (before tax) and 4.14 years (after tax). The optimization process will also increase the BEP value to 35% at a capacity of 439,232,519 kg/year.

## IV. CONCLUSION

Optimization and efficiency are carried out in the drying process by changing the operating conditions of the recycle air temperature from stage 2 to 250°C using a heater. Therefore, the H<sub>2</sub>O content in the AlF<sub>3</sub> product does not exceed the predetermined standard (<2.6%). Based on the calculation results, the percentage of H<sub>2</sub>O contained in the product is obtained. In this way, the factory can reduce off-specification products so as not to repeat the drying process (saving cost and production time). The optimization of the process carried out has an influence on the economic analysis of the factory, where before the optimization, the IRR value is 37.82%; POT of 3.29 years (before tax) and 4.06 years (after tax); and BEP of 34%. After the optimization, the IRR value is 36.97% and POT is 3.36 years (before tax) and 4.14 years (after tax); and BEP of 35%. From the economic analysis, the existing process give better value than after optimization but not specific, the IRR value increased 0.85%: POT only increased 0.07 years before tax and 0.08 years after tax; and BEP increased only 1%. However, the optimization still need to be done even the H<sub>2</sub>O content is only excess 0.2% from the specification because when the H<sub>2</sub>O content of the AlF<sub>3</sub> product does not meet specification, it will affect the whole process, including production capacity and the qualified standard of the product, product is not in anhydrous form but is still in hydrate form.

# SUPPLEMENTARY MATERIAL

*Process Flow Diagrams* (PFD) of Aluminum Fluoride Plant from Fluosilicic Acid and Aluminum Hydroxide can be seen in Figure 7 (before the optimization) and Figure 8 (after the optimization).

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Figure 7. Process Flow Diagram Aluminum Fluoride Plant (Before Optimization)



Figure 8. Process Flow Diagram Aluminum Fluoride Plant (After Optimization)