## **###PROOF OF SUBMISSION**

#### ESPR-D-19-11554 - ESPR: Submission Confirmation for Aluminum leaching from water treatment sludge and kinetic study - [EMID:eb6cd561ecb60b08] (External) Index ×

Environmental Science and Pollution Research <em@editorialmanager.com> to me • Wed, Dec 25, 2019, 10:30 AM 🔥 🕤 🚦

Dear Dr. Putra,

Your submission entitled "Aluminum leaching from water treatment sludge and kinetic study" has been received by Environmental Science and Pollution Research

The submission id is: ESPR-D-19-11554 Please refer to this number in any future correspondence.

You will be able to check on the progress of your paper by logging on to Editorial Manager as an author. The URL is https://www.editorialmanager.com/espri/.

Thank you for submitting your work to our journal.

Kind regards,

Editorial Office Environmental Science and Pollution Research ####Revision Required

#### ESPR: Your manuscript ESPR-D-19-11554 - [EMID:59fd1ee3c2f73982] (External) Inbox ×

Ta Yeong Wu <em@editorialmanager.com> to me ▼

CC: wu.ta.yeong@monash.edu, tayeong@hotmail.com

Ref.: Ms. No. ESPR-D-19-11554 Aluminum leaching from water treatment sludge and kinetic study Environmental Science and Pollution Research

Dear Dr. Putra,

Editor and reviewers have now commented on your paper. You will see that there are a number of issues that need to be addressed significantly.

We ask that you give the comments raised by the referees your careful consideration and that you submit a revised version of your manuscript as well as an itemized reply to each of the Editor's and reviewers' comments. Please make sure to submit your editable source files (i. e. Word, TeX)

Your revision is due by 18 Mar 2020.

If you decide to revise the work, please submit a list of changes or a rebuttal against each point which is being raised when you submit the revised manuscript.

If you need extra time to revise your manuscript, please contact Mrs. Berbon + joanne.berbon@springernature.com

To submit a revision, go to https://www.editorialmanager.com/espr/ and log in as an Author. You will see a menu item called 'Submissions Needing Revision'. You will find your submission record there.

I am looking forward to receiving you revised manuscript!

Yours sincerely,

Assoc. Prof. Ta Yeong Wu Editor Environmental Science and Pollution Research × 8 0

Mon, Feb 17, 2020, 11:39 PM 🔥 🕤 🚦

#### Editor:

Apart from both reviewers' comments, other concerns which must be considered by the authors are:

•Page 1, Lines 58-61: Make sure the Keywords used in the revised manuscript are not repeating the words used in the Title. Thus, please replace "aluminum", "kinetic" and "leaching" with other suitable Keywords which are not found in the title. Make sure 6 to 8 Keywords are provided and abbreviations such as HCI and WTS should be dismissed.

•Page 3, Lines 34-44: Please highlight the novelty of this work. Although sulfuric acid is commonly used, why hydrochloric acid was chosen in this study?

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## ###RESPONSE AND REBUTTAL TO REVIEWERS AND EDITOR ###REVISED MANUSCRIPT

## **##COVER LETTER FOR SUBMISSION OF REVISION**

Professor Ta Yeong Wu Editor Environmental Science and Pollution Research

Ref: ESPR-D-19-11554 Title: Aluminum leaching from water treatment sludge and kinetic study

Dear Prof. Wu,

We are very grateful for your email of February 17, 2020 regarding our above mentioned paper that was submitted to Environmental Science and Pollution Research. We are also grateful to the editor and the reviewers for their valuable comments that drew our attention to points that would improve the quality of our paper. We have taken into consideration all comments raised by the reviewers and revised the manuscript accordingly. An itemized response to their comments is listed in the response to reviewers' comments. We do hope that the revised version with changes marked by yellow highlight will come to your satisfaction.

Thanks for your attention. We do hope to hear positively from you soon.

Best Regards.

M.D. Putra, Corresponding author, on behalf of the coauthor

Meilana Dharma Putra, Ph.D. Chemical Engineering Department Lambung Mangkurat University Banjarmasin 70123 Indonesia Mobile: +6282281226215 Email: mdputra@ulm.ac.id;

## **Environmental Science and Pollution Research**

# Aluminum leaching from water treatment sludge using hydrochloric acid and kinetic study --Manuscript Draft--

Manuscript Number:	ESPR-D-19-11554R1					
Full Title:	Aluminum leaching from water treatment sludge using hydrochloric acid and kinetic study					
Article Type:	Research Article					
Corresponding Author:	Meilana Dharma Putra, Ph.D. Universitas Lambung Mangkurat Banjarbaru, Sauth Kalimantan, Indonesia INDONESIA					
Corresponding Author Secondary Information:						
Corresponding Author's Institution:	Universitas Lambung Mangkurat					
Corresponding Author's Secondary Institution:						
First Author:	Agus Mirwan					
First Author Secondary Information:						
Order of Authors:	Agus Mirwan					
	Meilana Dharma Putra, Ph.D.					
	Jhy-Chern Liu, Ph.D.					
	Susianto Susianto, Ph.D.					
	Ali Altway					
	Renanto Handogo, Ph.D.					
Order of Authors Secondary Information:						
Funding Information:	Kementerian Riset, Teknologi dan Pendidikan Tinggi	Dr. Agus Mirwan				
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Additional Information:	
Question	Response
§Are you submitting to a Special Issue?	No

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## Aluminum leaching from water treatment sludge using hydrochloric acid and kinetic study

Agus Mirwan<sup>a</sup>, Meilana Dharma Putra<sup>a\*</sup>, Jhy-Chern Liu<sup>b</sup>, Susianto<sup>c</sup>, Ali Altway<sup>c</sup>, Renanto

Handogo<sup>c</sup>

<sup>a</sup> Department of Chemical Engineering, Faculty of Engineering, Lambung Mangkurat University, Banjarbaru 70714, Indonesia

<sup>b</sup> Department of Chemical Engineering, National Taiwan University of Science and Technology, Taipei 106, Taiwan

<sup>c</sup> Department of Chemical Engineering, Faculty of Industrial Technology, Institut Teknologi Sepuluh Nopember, Surabaya 60111, Indonesia \*Corresponding author: Email: mdputra@ulm.ac.id

#### Abstract

Water treatment sludge (WTS) is abundantly produced in the world; the waste contributes to the environmental problems. Therefore for WTS utilization, aluminum leaching was employed using hydrochloric acid in this study. Al leaching efficiency increased from 72% to 80% as hydrochloric acid concentration increased from 1M to 4M. Decreasing the particle size and increasing the temperature increased Al leaching efficiency. The proposed kinetic model revealed that the rate controlling step followed a series of two leaching mechanisms: initially controlled by product-layer diffusion and then by a chemically controlled reaction. For instance, at 70 °C, the initial stage is well fitted by product-layer diffusion ( $R^2$ =0.87) compared to  $R^2$ =0.60 for chemical reaction; while for the second stage  $R^2$ =0.95 was observed via chemical reaction compared to  $R^2$ =0.74 for product-layer diffusion. The activation energies in these two stages were 9.58 kJ/mol and 10.73 kJ/mol, respectively. The proposed model was well validated by using data from literature and thus will be useful for other applications of leaching and extraction processes.

Keywords: Waste, environmental problem, mechanism, model, chemical surface, diffusion, removal.

#### Introduction

Water treatment plants (WTP) involves a coagulation process by using aluminum salts, e.g., polyaluminum chloride or aluminum sulphate, that cause hydrolyzation in raw water to form aluminum hydroxide precipitates. This process results in the production of voluminous sludge known as WTS. Globally, the total production of WTS can reach 10,000 tons/day (Okuda et al. 2014). Being a non-toxic material, most of WTS is disposed for land application and landfilling (Babatunde & Zhao 2007; Nair & Ahammed 2015). On the other hand, recycling WTS should be avoided because of organic solubilization (Meng et al. 2018). In recent years, many studies have been focused on Al recovery from WTS by leaching processes for possible reuse (Haynes & Zhou 2015). Various methods such as acidification, basification, ion exchange, and membranes have been employed for the recovery (Ahmad et al. 2016b; Nair & Ahammed 2014; Ooi et al. 2018). Ahmad et al. (2016a) indicated that Al recovery might not be simple, even though laboratory and plant scale tests have showed that it is feasible and economical. Intensive research in acid leaching has been under development due to the efficient and low-cost process compared to others methods (Jiménez et al. 2007). Thus, further exploration of means to recover Al from WTS is deemed important.

Acids have been examined for Al leaching from WTS. The commonly used sulfuric acid (H<sub>2</sub>SO<sub>4</sub>) is one of the effective solvents to extract aluminum. Jiménez et al (Xu et al. 2009) reported that sulfuric acid could extract 70% of Al within 30 min at pH=2. The leaching efficiency of Al reached 83.6% using 1M H<sub>2</sub>SO<sub>4</sub> within 30 min (Cui et al. 2015). Okuda et al. (Okuda et al. 2014) reported that more than 80% of Al was extracted within 2 h using H<sub>2</sub>SO<sub>4</sub> at pH=1 under ambient temperature. However, due to its toxicity, sulfuric acid can bring some negative effects on environment and could lead to self-inhibition effect due to the new interaction during leaching process (Seidel & Zimmels 1998). The use of hydrochloric acid as a solvent in leaching is gaining grounds in research due to its lower acute toxicity compared to sulfuric acid (Hagen & Järnberg 2009). The leaching process

using HCl continues to improve to attain high efficiency. It was reported that the sulfuric acid and hydrochloric acid lead to corrosion effect in vessel; however, corrosion rate for both acid can be significantly reduced by using hard chrome plating (Ajeel et al. 2012). The leaching rate of Al from secondary aluminum dross at 75 °C for 600 min was about 7-18% (Yang et al. 2019a). On the other hand, the extraction efficiency of 94% was obtained from raw kaolin using HCl at 70 °C for 6 h; however, sodium chloride was also formed (Bhattacharyya & Behera 2017). The leaching efficiency using HCl varied depending on the raw materials and process condition such as temperature, time, etc.

Kinetic models are important tools to describe the mechanism of the leaching process. Some studies have used the shrinking core model to define the dissolution kinetics of aluminum from fly ash and kaolin (Cheng et al. 2012; Tang et al. 2010). A kinetic process of aluminum leaching is described via diffusion or chemical reaction. However, the exact rate limiting step for the kinetics is still undetermined for a general aluminum leaching process. Hence, the kinetics should be studied in detail to explore the role of the rate limiting step in the system.

Many studies have been devoted to assess the optimal conditions for acid leaching of Al from the WTS. However, most of them were focused on the use of  $H_2SO_4$  (Cheng et al. 2016; Cheng et al. 2012; Keeley et al. 2016). Some leaching processes have been also carried out from raw materials using HCl; thus, the Al leaching process from WTS using HCl would be investigated thoroughly in this study. On the other hand, the information on the mechanism of leaching process seems insufficient. In the present study, the effects of some main parameters, such as acid concentration, particle size, and temperature on Al leaching have been investigated using HCl. The kinetics of Al leaching from WTS has been further studied using the shrinking core (SC) model. A modified kinetic model has also been developed in this study.

#### Material and methods

#### Materials and characterization

WTS was collected from the water treatment plant in Banjarmasin, South Kalimantan, Indonesia. It was washed and dried under direct sunlight for 48 h. The material was further crushed, screened, and separated into the size fractions of 70-120 mesh (0.210-0.125 mm), 120-200 mesh (0.125-0.074 mm), and 200-325 mesh (0.074-0.044 mm). Samples were then stored in plastic bottles for experiments without any further treatment.

WTS was characterized by X-ray diffractometer (XRD, Philips X-pert powder model, Netherlands) using powder diffraction database file-2 (PDF-2). The results showed the major mineral phases of kaolinite (Al<sub>2</sub>Si<sub>2</sub>O<sub>5</sub>(OH)<sub>4</sub>), quartz (SiO<sub>2</sub>), hematite (Fe<sub>2</sub>O<sub>3</sub>), and corundum (Al<sub>2</sub>O<sub>3</sub>) (Fig. 1). The surface functional groups of each constituent solids were characterized using Fourier Transformation Infrared Spectroscopy (FTIR, Shimadzu, Japan), and the results of which are shown in Table 1. Flake structure of WTS was found when being analyzed by scanning electron microscopy equipped with X-ray microanalysis (SEM-EDX, SEM EVO MA 10, Germany). The results showed the presence of dominant elements, namely Al, Si, Fe (color difference) with the composition of 32.78%, 49.15%, and 18.07%, respectively as shown in Fig. 2. Similar observation of WTS sample was also shown in other literature containing Fe<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> (Ahmad et al. 2016b; Cheng et al. 2012).

Aluminum content in the samples was determined by using the inductively coupled plasmacluster optical emission spectrometer (ICP-OES, 9060-D Teledyne Leeman Labs, USA). Each analysis was triplicated and the average value was reported. The Al leaching ratio (x) can be calculated as:

$$x = X/X_0$$

(1)

where  $X_0$  denotes total Al content and X refers to the amount of Al leached (Yang et al. 2019b).

#### Leaching process

The leaching process was conducted in a 500-mL Pyrex reactor with a thermostat water bath (Fig. 3). Mixing was achieved by using a magnetic stirrer at 300 rpm. Al leaching from WTS was carried out by placing 5 g of WTS into 250 ml of HCl (37%, Sigma-Aldrich) solution at various concentrations (1, 2, 4, and 6 M). WTS samples with different particle size (70-120, 120-200, and 200-325 mesh) were used in experiments conducted at different reaction temperatures (30, 50, 70, and 90 °C). Total reaction time was controlled at 60 min. At selected time interval, samples were collected and then filtered by a syringe for the analysis of Al content.

#### Shrinking core (SC) model

The shrinking core (SC) model describes the kinetic characteristics of non-catalytic, heterogeneous, solid-liquid reactions consisting of three processes: film diffusion, product-layer diffusion, and surface chemical reaction (Keeley et al. 2016). In this study, only product-layer diffusion and surface chemical reaction were considered due to the fast process of film diffusion. The kinetic equation for a leaching process controlled by a surface chemical reaction is presented as follows:

$$\frac{t}{\tau_c} = 1 - (1 - x)^{1/3} \tag{2}$$

If the leaching process is controlled by the diffusion through product-layer, the kinetic equation is described as follows:

$$\frac{t}{\tau_i} = 1 - 3(1 - x)^{2/3} + 2(1 - x)$$
(3)

where  $\tau_c$  is the complete leaching time for process controlled by surface chemical reaction,  $\tau_c = \rho_B R / b M_w k_c C_A$ ,  $\tau_i$  is the complete leaching time for a process controlled by productlayer diffusion,  $\tau_i = \rho_B R^2 / 6b M_w D_e C_A$ , x is the fraction of Al leached out, t is the reaction time,  $\rho_B$  is the solid density, R is the radius of initial particle, b is the stoichiometric coefficient,  $M_w$  is the molecular weight (g/mol),  $k_c$  is the factor of mass transfer,  $C_A$  is the hydrochloric acid concentration, and  $D_e$  is the coefficient of product-layer diffusion.

#### **Results and discussion**

#### Effect of acid concentration

Figure 4 shows the amount of extracted aluminum with time at the temperature of 90 °C, particle size of 200-325 mesh and various concentrations. Aluminum (Al) is the major element of the WTS in the form of Al(OH)<sub>3</sub> and the solubility is affected by pH. The main reaction in the leaching process is shown as follows:

 $Al(OH)_{3(s)} + 3HCl_{(aq)} \rightarrow AlCl_{3(aq)} + 3H_2O$ 

As shown in Figure 4, the efficiency of Al leaching increased from 72% to 80% as HCl concentration changed from 1M to 4M. Although at 6M of HCl, it slightly decreased to 78% indicating the key role the pH plays in Al leaching from WTS. This acidification process (with low pH) is more convenient and efficient compared to the basification process (Baba et al. 2009). As reported by Cheng et al. (Cheng et al. 2012), metal such as Al is easily dissolved at pH<2.5. On the other hand, the solubility of aluminum chloride (AlCl<sub>3</sub>) decreases with increasing HCl concentration at concentration higher than 5M (Seidel & Zimmels 1998); this notion is similarly in accordance with our finding here (>4M). It was plausible due to the formation of other metal-chloride (Raza et al. 2015). When AlCl<sub>3</sub> is saturated in the reaction system, dissolution and precipitation of Al(OH)<sub>3</sub> might reach a dynamic equilibrium. The HCl concentration of 4 M was therefore chosen for subsequent experiments.

#### Effect of particle size and temperature

Fig. 5 shows the amount of extracted aluminum with time at the concentration of 4M, at the temperature of 90 °C and with various particle sizes. The efficiency of Al leaching increased as particle size decreased. It indicated that smaller particle sizes lead to faster

leaching process. It could be reasonably related to the acceleration of the mass transfer process (Brantley et al. 2008) thus minimizing the effect of internal diffusion. Also, smaller particle size increased the surface area; thus, it consequently resulted in larger contact between particle and solvent (Adekola et al. 2017). Fig. 6 shows the amount of extracted aluminum with time at acid concentration of 4M, particle size of 200-325 mesh and various temperatures. The Al leaching efficiency increased with temperature and an optimum process was obtained at 4M and 90°C for 60 min with the efficiency of 82%. Increasing the temperature from 70 °C to 90 °C significantly increased the Al leaching efficiency (from 67% to 82%) compared to the increasing temperature from 50 °C to 70 °C (62% to 67%) and from 30 °C to 50 °C (58% to 62%). This revealed that the temperature range 70-90 °C is crucial for the leaching process; this situation will plausibly continue at a higher temperature. Significantly higher leaching efficiency (>70%) was observed by increasing temperature at above 70 °C (Raza et al. 2015; Shalchian et al. 2018). On the other hand, a leaching efficiency of about 99% was obtained at 70 °C using succinic acid, an expensive solvent (Raza et al. 2015). The different results could be attributed to differences in used solvents and extracted components.

#### Statistical Analysis

The resulted data were also affirmed by conducting statistical test using Anova analysis. Significant effect of HCl concentration on the Al leaching ratio (p=0.0007) was obtained. Significant effect were also obtained in each range of concentration between 1M and 2M (p=0.0474), 2M and 4M (p=0.0343); while insignificant effect was observed in the concentration range of 4M and 6M (p=0.2924) due to the drop of Al leaching ratio at higher concentration.

For size particle effect, significant effect was also achieved at total range  $(p=1.57 \times 10^{-7})$  as well as in each range between +70/-120 and +120/-200 (p=0.0027) and +120/-200 and +200/-325 (p=6.84x10<sup>-5</sup>). The statistical results also revealed the significant effect of

# temperature on Al leaching ratio at total temperature range (p=1.29x10<sup>-8</sup>) and the range of 30 °C and 50 °C (p=0.0124), 50 °C and 70 °C (p=0.0284) and 70 °C and 90 °C (p=0.0005).

#### Kinetic study and mechanism

Kinetic study is important to elucidate the various processes in the system. The shrinking core model used in the study includes the product-layer diffusion and chemical reaction. The kinetic model using shrinking model has been applied to fit the experimental data. The correlation coefficient values  $(R^2)$  for product-layer diffusion and chemical reaction were 0.75-0.84 and 0.57-0.67, respectively for all experimental data of concentration, particle size and temperature. Although the fitting results of the kinetic model and experimental were unsatisfactory, the product-layer diffusion as a rate limiting step was found more appropriate for the system. It was reported that for the process controlled by surface chemical reaction a plot of  $1 - (1 - x)^{1/3}$  should be a straight line with slope of  $1/\tau_c$  and for the process controlled by the product-layer diffusion, a plot of  $1 - 3(1 - x)^{2/3} + 2(1 - x)$  should be a straight line with slope of  $1/\tau_i$  (Tian et al. 2018; Yang et al. 2020; Yang et al. 2019b). However, the low values of  $R^2$  obtained here were, probably, because both the exponential rate and linear rate were observed in the leaching process thereby it couldn't be tackled by the shrinking model. On other hand, the chemical reaction controlled the kinetic process (Brantley et al. 2008); contrarily, the kinetic model of product-layer diffusion was proven to fit the experimental data (Cheng et al. 2012). Those different results of the rate of limiting step were due to the used solvent, the extracted component, the raw materials and the temperature affecting the kinetic process (Brantley et al. 2008). In fact, the kinetic model in those literatures was only evaluated during the exponential rate of leaching; while, the subsequent leaching rate cannot be ignored as the increase in the leaching efficiency was observed at about 14-20% (Cui et al. 2015). On the other hand, researchers revealed that the experimental data did not conform to both kinetic models of chemical reaction or productlayer diffusion (Yang et al. 2019b); they concluded that the mathematical approach failed to explain the dynamic situation due to finer particle size, grain shape, and reactant activity (Shalchian et al. 2018). In fact, they had both stages of exponential and linear leaching process. Hence, their difficulty of interpretation can be clearly described in this study later. In the next discussion, we will clearly explore the effect of product-layer diffusion and chemical reaction as rate limiting step on both proposed regions of leaching rate as shown in Fig. 7. The exponential rate (t< 20 min) is used in the first-stage and the linear rate (t>20 min) in the second-stage. The mechanisms of leaching process were further assessed separately for each stage and the integration limits of the corresponding differential mass balance equation consequently changed. For the first-stage process, the equations (2) and (3) were applied for surface chemical reaction and product-layer diffusion, respectively. For the second-stage, the equation based on surface chemical reaction as rate of limiting step led to:

$$\frac{t - t_1}{\tau_c} = 1 - \left(\frac{1 - x}{1 - x_1}\right)^{1/3} \tag{4}$$

$$\tau_{c} = \rho_{B} R \left( 1 - x_{1} \right)^{1/3} / b M_{w} k_{c} C_{A}$$
(5)

where *t* is the total reaction time,  $t_1$  is the first-stage reaction time, *x* is the total Al leached out, and  $x_1$  is the Al leached out in the first stage. Both  $t_1$  and  $x_1$  values were obtained from experimental data as the initial conditions. The kinetic equation for the rate limiting step based on product-layer diffusion can be further expressed as follows:

$$\frac{t-t_1}{\tau_i} = 1 - 3 \left( \frac{1-x}{1-x_1} \right)^{2/3} + 2 \frac{(1-x)}{(1-x_1)}$$
(6)

$$\tau_{i} = \rho_{B} R^{2} \left( 1 - x_{1} \right)^{2/3} / 6b M_{w} D_{e} C_{A}$$
(7)

Table 2 shows the constants of leaching rate in the first and second stages of various HCl concentrations. It was clear that in the first stage, the leaching process of Al was controlled by the product-layer diffusion since the correlation coefficient value ( $R^2$ ) for the rate constant of product-layer diffusion ( $K_i$ ) was higher than that of surface chemical reaction

( $K_c$ ). For the second-stage process, it was found that the  $R^2$  value of  $K_c$  was slightly higher than that of  $K_i$  especially at a higher acid concentration (> 1M). This finding implies that Al leaching could be appropriately controlled by chemical reaction. A similar finding was observed at various particle sizes for both first and second stages as presented in Table 3; furthermore, the  $R^2$  value of  $K_i$  for second stage was very low hence the diffusion could not be justified to control the process. This revealed that the diffusion process was dominant at the beginning of process for short time. Hence, at the second stage, after the particle experienced the shrinkage by time, the diffusion rate would be very fast, and the chemical reaction subsequently became the rate limiting step. Furthermore, the leaching rate process depends more on the particle size compared to acid concentration as the increase in rate constants became twofold when the particle size was reduced from 120-200 to 200-325 mesh.

The division of two stages was also clearly observed at the temperature effect with significantly different values of  $R^2$  for both product-layer diffusion and chemical reaction as shown in Table 4. It was reported that at a lower temperature the chemical reaction control was more appropriate; conversely at higher temperature, the leaching process was controlled by product-layer diffusion [10]. In this study, the  $R^2$  values were similar for both, the chemical reaction and layer diffusion, at 30 °C in the first stage; the kinetic model of product-layer diffusion fitted well at a higher temperature. In the second stage, the chemical reaction was dominantly observed to be the rate limiting step. It indicated that temperature contributed significantly to the leaching process; in as much as the constants rate can be related to the exponential function of temperature by Arrhenius equation:  $K = A e^{\left(-\frac{E_a}{RT}\right)}$ . A is the pre-exponential factor;  $E_a$  is the activation energy (kJ·mol<sup>-1</sup>); *R* is the gas constant, 8.314 × 10<sup>-3</sup> kJ·mol<sup>-1</sup>·K<sup>-1</sup>, and *T* is the temperature (K).

The activation energies obtained in this work for the first stage and second stage were 9.58 kJ/mol and 10.73 kJ/mol, respectively. Compared with other results (Table 5), the value

of activation energy obtained in the current study was lower than that in those literatures (Cheng et al. 2012; Cui et al. 2015; Tang et al. 2010). It implies that the leaching process of aluminum from WTS using HCl was easier. It was reported (Tang et al. 2010) that dissolution kinetics controlled by surface chemical reaction resulted in a higher activation energy (>42 kJ/mol), whereas a lower activation energy (<20 kJ/mol) suggested that product-layer diffusion is the rate controlling step (Cui et al. 2015). The value of activation energy value in second stage was out of range for chemical reaction as the rate limiting step. It is possible since they only evaluated the leaching process in the first stage; while a further way of the second stage was also observed in this work as the continual process. Hence, due to shrinkage phenomenon that diminishes the diffusion barrier, the chemically controlled process in the second stage would be more plausible to take place thus implying the low activation energy.

To prove the involvement of a series-kinetic control process of chemical reaction and product-layer diffusion, a combined kinetic model was proposed. The combined kinetic model is described as follows:

$$K_{p}t = \alpha \left[ 1 - (1 - x)^{1/3} \right] + (1 - \alpha) \left[ 1 - 3(1 - x)^{2/3} + 2(1 - x) \right]$$
(8)

$$K_{pm}(t-t_{1}) = \alpha \left[ 1 - 3 \left( \frac{1-x}{1-x_{1}} \right)^{2/3} + 2 \frac{(1-x)}{(1-x_{1})} \right] + (1-\alpha) \left[ 1 - \left( \frac{1-x}{1-x_{1}} \right)^{1/3} \right]$$
(9)

$$\alpha = \frac{K_i}{K_c} \tag{10}$$

 $K_p$  is the rate constant of the combined kinetic model. Equation 8 and 9 were applied for the first and second stage process, respectively.  $\alpha$  is the ratio of  $K_i$  (coefficient for diffusion rate) to  $K_c$  (coefficient for chemical reaction rate). The higher value of  $\alpha$  implies chemical reaction rate is controlling. The value of  $\alpha$  is in the range of 0-1.

Figure 8a and 8b show the plot of the combined kinetic model for first and second stages at the temperature of 70 °C, respectively. The figures for other temperatures (not presented here) show similar trends as shown in Figure 8. In Figure 8a, the lower value of  $\alpha$  led to higher value of  $R^2$ ; it means that the product-layer diffusion controlled the process rate. This finding thus confirms the previous conclusion that in the first stage the leaching process was predominantly controlled by product-layer diffusion. In the second stage, the kinetic model with the high value of  $\alpha$  fitted well the data as indicated by high value of  $R^2$ . Hence, the leaching mechanism was mainly affected by a chemical reaction process. The proposed model was validated by using data from literature (Yang et al. 2019b), where the aluminum removal from diamond wire saw powder using hydrochloric acid 2 M at 60 °C was carried out. As shown in Fig. 9, good correlation fittings in the first stage ( $R^2=0.92$ ) and the second stage ( $R^2$ =0.98) that describes the product-layer controlled diffusion and the chemical reaction controlled, respectively were obtained. This finding obviously disproves the previous conclusion (Yang et al. 2019b) regarding the failed mathematical approach at higher temperature ( $R^2$ =0.43-0.70) due to finer particle. This again confirms the previous finding using the separated model. Thus, the new combined kinetic model based on the general shrinkage core model could be well proposed for the system of leaching or extraction process. Moreover, the proposed model can be applied in other applications such as dehydration reaction, hydration kinetic, hydrolysis process, pyrolysis process etc. to predict the experimental data as the rate of shrinkage was shown by exponential and linear curves (Farsi et al. 2019; Huang et al. 2018; Lan et al. 2015; Zhang et al. 2019).

For future step, the Al produced through leaching process could be precipitated using a base to form AlPO<sub>4</sub>; this could be applied in phosphate industry or fertilizer in agriculture (Muisa et al. 2020; Pradel et al. 2020). On other hand, the Alum sludge can be used as absorbent for heavy metals (Dassanayake et al. 2015). After extraction process using HCl, the non-toxic organic materials remains in the sludge (Smith et al. 2009). This residual sludge essentially provides geopolymeric feedstock (Gomes et al. 2019). Furthermore, the residue becomes secondary raw material for the ceramic and glass industries (Zichella et al. 2020).

The kinetics of Al leaching from WTS was investigated using HCl. Leaching experiments were performed with four different acid concentrations, three different particle sizes, and four different temperatures. Al leaching efficiency increased as HCl concentration changed from 1M to 4M. However, the limited solubility of AlCl<sub>3</sub> at higher concentration (6M) hindered the leaching reaction. In addition, both smaller particle size and higher temperature resulted in higher leaching efficiency. The proposed kinetic model revealed the importance of the dividing the leaching process into two stages. The first stage was more suitable for the kinetic of product-layer diffusion; while the second stage was controlled by the chemical reaction. The proposed model that has been well validated with the literature data showed the prominence to be developed for wide application process using shrinking model.

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#### **Conflict of interest**

The authors declare that there is no conflict of interest.

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Infrared band (cm <sup>-1</sup> )	Transmittance (%)	Functional group
3695.36	29.586	Al–O–H str (kaolinite, illite)
3620.14	26.456	Al–O–H (kaolinite, illite, calcite)
3448.49	24.903	H–O–H str (kaolinite, illite)
1634.56	37.454	H–O–H str ( <i>illite</i> , <i>calcite</i> )
1104.17	16.372	Si–O str (kaolinite, quartz)
1031.85	8.981	Si–O–Si, Si–O str. (kaolinite, illite)
1007.74	10.150	Si–O str (kaolinite, quartz)
913.23	23.281	Al–O–H str (kaolinite, illite, hematite)
779.19	32.612	Si–O–Al str (kaolinite, illite)
694.33	29.459	Si–O str, Si–O–Al str (quartz, kaolinite)
536.17	15.456	Si–O str, Si–O–Al str (kaolinite)
468.67	13.958	Si–O str, Si–O–Fe str. (quartz, kaolinite)
426.24	21.678	Si–O str (quartz)

Table 1 Surface functional groups of FTIR of WTS

	First-	stage of le	eaching p	rocess	Second-stage of leaching process			
Acid								
	Rate co	onstants	Correlation		Rate constants		Correlation	
concentration		1.	2				-2	
	(mi	$n^{-1}$ )	coeffici	ent $(R^2)$	ent $(R^2)$ (min <sup>-1</sup> )		coefficient ( $R^2$ )	
(M)			<b>D</b> <sup>2</sup>				2 2	
	$K_i$	$K_c$	$R^2_i$	$R^2_c$	$K_i$	$K_c$	$R^2_i$	$R^2_c$
	0.0010	0.0000	0.72	0.00	0.0006	0.0005	0.02	0.00
6	0.0212	0.0293	0.72	0.39	0.0006	0.0025	0.93	0.98
4	0.0100	0.0274	0.95	056	0.0006	0.0025	0.05	0.06
4	0.0190	0.0274	0.85	0.50	0.0006	0.0025	0.95	0.90
2	0.0153	0.0242	0.03	0.67	0.0007	0.0027	0.03	0 00
2	0.0155	0.0272	0.75	0.07	0.0007	0.0027	0.75	0.77
1	0.0113	0 0204	0 94	0 74	0.0008	0.003	0.96	0.92
1	0.0110	0.0201	0.71	0.71	5.0000	0.000	0.20	0.72

Parameters of the kinetic model for Al leaching at different acid concentrations

	First-s	tage of leach	ing proce	ess	Second-stage of leaching process				
Particle size	Rate constants (min <sup>-1</sup> )		Correlation coefficient ( $R^2$ )		Rate cons	tants (min <sup>-1</sup> )	Correlation coefficient		
(mesh)							$(R^2)$		
	$K_i$	$K_c$	$R^2_i$	$R^2_c$	$K_i$	$K_c$	$R^2_i$	$R^2_{c}$	
+200/-	0.0179	0.0265	0.89	0.57	0.0010	0.0033	0.80	0.98	
325									
+120/-	0.0164	0.0164	0.97	0.91	0.0003	0.0018	0.95	0.97	
200									
+/0/-	0.0048	0.0125	0.93	0.96	0.0003	0.0018	0.93	0.97	
120									

Parameters of the kinetic model for Al leaching at different particle sizes

	First-stage of leaching process				Second	d-stage of	leaching J	process
Temperature	Rate constants		Correlation		Rate constants		Correlation	
(°C)	(mi	$n^{-1}$ ) coefficient ( $R^2$ )		$(\min^{-1})$		coefficient ( $R^2$ )		
	$K_i$	Kc	$R^2_i$	$R^2_{c}$	$K_i$	Kc	$R^2_i$	$R^2_{c}$
90	0.0179	0.0265	0.89	0.57	0.0010	0.0033	0.81	0.98
70	0.0128	0.0220	0.87	0.60	0.0002	0.0015	0.74	0.95
50	0.0100	0.0190	0.95	0.79	0.0002	0.0015	0.73	0.94
30	0.0068	0.0150	0.93	0.95	0.0002	0.0120	0.71	0.92

Parameters of the kinetic model for Al leaching at various temperatures

<mark>Time</mark> min	<mark>Temperature</mark> ℃	Model	Rate- controlling step	Activation energy (kJ/mol)	Raw Material	Solvent	Ref.
<mark>0 –</mark> 160	<u>60 – 100</u>	<mark>One</mark> stage	Surface chemical reaction Diffusion process	43 24	Kaolin residue	Hydrochloric acid	(Tang et al. 2010)
<mark>0 –</mark> 60	<u>10 – 70</u>	<mark>One</mark> stage	Inert-layer diffusion	<mark>32.32</mark>	Water purification sludge	Sulfuric acid	<mark>(Cheng</mark> et al. 2012)
<mark>0 –</mark> 120	40 – 80 90 – 106	<mark>One-</mark> stage	Surface reaction Product- layer diffusion	57.65 12.33	Coal mining waste	Hydrochloric acid	<mark>(Cui et</mark> al. 2015)
0 – 15 15 – 60	<mark>30 – 90</mark>	First stage Second stage	Product- layer diffusion Surface chemical reaction	9.58 10.73	Waste treatment sludge	Hydrochloric acid	Present study

Activation energy comparison of leaching process from literature

#### **Caption of Figures**

Fig. 1. XRD pattern of WTS

Fig. 2. SEM pictures of WTS

- Fig. 3. Leaching process and equipment
- Fig. 4. Effect of acid concentration on Al leaching from WTS

Fig. 5. Effect of particle size on Al leaching from WTS

Fig. 6. Effect of temperatures on Al leaching from WTS

Fig. 7. Division of stages in Al leaching process

Fig. 8. Profile of the combined kinetic model against experimental data at 70°C for (a) first-

stage leaching and (b) second-stage leaching

Fig. 9. Validated profile of the combined kinetic model against experimental data from

literature for (a) first-stage leaching and (b) second-stage leaching





















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To: Agus Mirwan <agusmirwan@ulm.ac.id>, liu1958@mail.ntust.edu.tw, alimohad@chem-eng.its.ac.id, susianto@chemeng.its.ac.id, renanto@chem-eng.its.ac.id, susianto.sst@gmail.com, alimohay@yahoo.com

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Dear all,

Congrats to Dr. Meilana Dharma Putra and Dr. Agus Mirwan and other collaborators, for the good work. Please keep the good work to reach highest goals in UNLAM and nationwide. Regards. Renanto

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Dear All,

Alhamdulillah ...

I am very pleased to receive this information. Thank you to all for your help, support and cooperation so far, especially to Dr. Meilana D Putra, Prof. JC. Liu and my supervisors Prof. Renanto, Prof. Ali Altway, and Dr. Susianto. I hope this collaboration will continue in the future.

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Dear Dr. Meilana Dharma Putra,

please be advised that my name is Renanto (first name) Handogo (family name). Thank you.

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To: "Prof. Ir. Renanto, M.Sc.Ph.D(400)" <renanto@chem-eng.its.ac.id> Cc: Agus Mirwan <agusmirwan@ulm.ac.id>, "liu1958@mail.ntust.edu.tw" <liu1958@mail.ntust.edu.tw>, "Prof. Dr.Ir. Ali Altway, M.Sc.(251)" <alimohad@chem-eng.its.ac.id>, "Dr.Ir. Susianto, DEA.(1336)" <susianto@chem-eng.its.ac.id>, "susianto.sst@gmail.com" <susianto.sst@gmail.com>, "alimohay@yahoo.com" <alimohay@yahoo.com>

Dear Prof. Renanto

Thanks for the correction. The mistake is made from the administrator transferring the content to their template, because your name is correctly mentioned in the accepted manuscript as well as in their metadata. Anyway, thank you for the clarification, I will make it in the correction to the publisher.

Best regards,

12/29/22, 9:47 PM

Meilana Dharma Putra Chemical Engineering Department Lambung Mangkurat University

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