# **PHOSPHATE REMOVAL FROM WASTEWATER USING SLAG AND ION EXCHANGE RESINS**

**By** 

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#### **CERTIFICATE OF AUTHORSHIP/ ORIGINALITY**

I certify that the work in this thesis has not previously been submitted for a degree nor has it been submitted as part of requirements for a degree except as fully acknowledged within the text.

I also certify that the thesis has been written by me. Any help that I have received in my research work and the preparation of the thesis itself has been acknowledged. In addition, I certify that all information sources and literature used are indicated in the thesis.

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## **TABLE OF CONTENTS**



### **CHAPTER 1 - INTRODUCTION 1**











## **LIST OF FIGURES**



**Figure 4.6** Langmuir isotherm model calculation for phosphate removal by (a) Purolite A860S, (b) Dowex 21K XLT, and (c), Dowex 21K XLT\_Cu **77 Figure 4.7** Freundlich isotherm model calculation for phosphate removal by (a) Purolite A860S, (b) Dowex 21K XLT, and (c) Dowex 21K XLT\_Cu **78**  Figure 4.8 Effect of contact time on the removal of phosphate (phosphate initial concentration: 20 mg /L, resin dose:  $1.5 g/L$ **82 Figure 4.9** Pseudo-first order kinetic model calculation for phosphate removal by (a) Purolite A860S, (b) Dowex 21K XLT, and (c) Dowex 21K XLT Cu (resin dose: 1.5  $g/L$ ) **86 Figure 4.10** Pseudo-second order kinetic model calculation for phosphate removal by (a) Purolite A860S, (b) Dowex 21K XLT, and (c) Dowex 21K XLT Cu (resin dose: 1.5  $g/L$ ) **87 Figure 4.11** Elovich model calculation for phosphate removal by (a) Purolite A860S, (b) Dowex 21K XLT, and (c) Dowex 21K XLT Cu. (resin dose:  $1.5$  g/L) **88 Figure 4.12** Intraparticle diffusion kinetic model by (a) Purolite A860S, (b) Dowex 21K XLT, and (c) Dowex 21K XLT\_Cu (resin dose: 1.5 g/L) **89 Figure 4.13** Breakthrough curve for different resins, filtration velocity =20 ml/min, initial phosphate concentration (a)  $20$ mg/L and (b) 30 mg/L **91 Figure 4.14** Phosphate removal efficiency with different inlet concentration (a) 20 mg/L and (b) 30 mg/L **93** 

- **Figure 4.15** Thomas model calculation for (a) Purolite A860S, (b) Dowex 21K XLT and (c) Dowex 21K XLT-Cu (initial phosphate concentration: 20 mg /L) **95**
- **Figure 4.16** Thomas model calculation for (a) Purolite A860S, (b) Dowex 21K XLT and (c) Dowex 21K XLT-Cu (initial phosphate concentration: 30 mg/L) **96**
- **Figure 4.17** Yoon-Nelson model calculation for (a) Purolite A860S, (b) Dowex 21K XLT and (c) Dowex 21K XLT-Cu (initial phosphate concentration: 20 mg/L **98**
- **Figure 4.18** Yoon-Nelson model calculation for (a) Purolite A860S, (b) Dowex 21K XLT and (c) Dowex 21K XLT-Cu (initial phosphate concentration 30 mg/L) **99**
- **Figure 4.19** The efficiency of phosphate desorption from Dowex 21K  $XLT$  Cu using Na<sub>2</sub>SO<sub>4</sub>, NaCl and NaOH solutions **102**
- **Figure 4.20** SEM images of (a) SFS and (b) M-SFS **105**
- **Figure 4.21** Schematic diagram of modification of SFS in the NaOH solution (You et al., 2011) **106**
- **Figure 4.22** Effect of adsorbent dosage on (a) phosphate removal rate and (b) adsorption capacity of SFS and M-SFS (initial phosphate concentration  $= 20$  mg/L, mixing time: 24h) **109**
- **Figure 4.23** Freundlich isotherm model calculation for phosphate removal by (a) SFS, (b) M-SFS **111**
- **Figure 4.24** Langmuir isotherm model calculation for phosphate removal by (a) SFS, (b) M-SFS **112**
- **Figure 4.25** Comparison of phosphate removal from synthetic wastewater by SFS and M-SFS (Initial P concentration = 20 mg/L, amount of adsorbents  $= 4g/L$ ) **115**
- **Figure 4.26** Pseudo-first order kinetic model calculation for phosphate removal by (a) SFS and (b) M-SFS **117**
- **Figure 4.27** Intraparticle diffusion kinetic model for phosphate removal by (a) SFS and (b) M-SFS **118**
- **Figure 4.28** Intraparticle diffusion kinetic model for phosphate removal by (a) SFS and (b) M-SFS **120**
- **Figure 4.29** Elovich kinetic model for phosphate removal by (a) SFS and (b) M-SFS **121**
- **Figure 4.30** Breakthrough curve for (a) SFS and (b) M-SFS, filtration velocity =  $20$  mL/min, adsorbent amount =  $630g$ , initial P concentration =20mg /L **123**
- **Figure 4.31** Adsorption efficiency by SFS and M-SFS column, filtration velocity =  $20$  mL/min, adsorbent amount =  $630g$ , initial P concentration =20mg /L **124**
- **Figure 4.32** Thomas column model calculation for (a) SFS, (b) M-SFS (initial P concentration 20 mg/L) **126**
- **Figure 4.33** Yoon-Nelson column model for (a) SFS, (b) M-SFS (initial P concentration: 20 mg/L) **127**

## **LIST OF TABLES**

**CHAPTER 2** 





#### **Nomenclature**

- a = Langmuir constant
- AC = activated carbon
- $Al^{3+} =$  Aluminum
- $b =$ Langmuir constant
- $BW = boron waste$
- $C$  = the bulk phase concentration (mg/L)
- $Ca(OH)<sub>2</sub> = Caclcium hydroxide$
- $C_e$  = equilibrium concentration of adsorbate (mg/L)
- $C_0$  = initial concentration of adsorbate (mg/L)
- $C_0$  = inlet adsorbate concentration (mg/L)
- $C_s$  = the concentration on the external surface (mg/L)
- $C_t$  = concentration of adsorbate at time t (mg/L)
- $C_t$  = outlet adsorbate concentration at time t (mg /L)
- $C_{Ek}$  = the concentration of potassium ions in solution at equilibrium

 $Ca^{2+} = Ca$ lcium (II)

- $Cl =$ chloride
- $CO<sub>3</sub><sup>2-</sup> =$  carbonate
- $Cu^{2+} = Copper (II)$
- $D_m$  = molecular diffusion coefficient

 $D_s$  = the surface diffusion coefficient (m<sup>2</sup>/s)

 $EXAFS = X-ray$  absorption fine structure

 $Fe^{3+} = i$ ron (III)

 $F =$  the linear velocity calculated by dividing the filtration velocity by column section area

 $FeO =$  zero-valent iron

 $g/L$  = gram per litre

 $HAIX = hybrid$  anion ion exchange

 $H_3PO_4$  = Phosphoric axit

H2PO4 - =dihydrogen phosphate ion

- $HC = hydrochloric acid$
- $HCO<sub>3</sub>$  = bicarbonate
- $HFO = iron (iii) oxide$

 $hr = hours$ 

 $K = a$  constant called the "half value".

 $K^+$  = Potassium

 $k_f$  = the external mass transfer coefficient (m/s)

 $K_F$  = Freundlich constants (mg/g)

 $K_L$  = Langmuir constant related to the energy of adsorption (L/mg)

- $k_{Th}$  = Thomas rate constant (mL/min.mg)
- $k_{YN}$  = rate velocity constant (1/min)
- $k_1$  = equilibrium rate constant of pseudo-first-order sorption (1/min)
- $k_2$  = equilibrium rate constant of pseudo-second-order (1/min)
- $k_{AB}$  = kinetic constant, (L/mg.min)

 $KNO<sub>3</sub> = Potassium nitrate$ 

 $KH<sub>2</sub>PO<sub>4</sub> = Monopotassium phosphate$ 

KCl = Potassium chloride

LDHs = layered double hydroxides

 $M =$  mass of dry adsorbent (g)

 $MIEX = magnetic$  ion exchange resin

mg  $P/L =$  milligram phosphorus per litre

mg  $PO_4^{3-}$  /  $g = mg$  phosphate per gram

mg  $P/g =$  milligram phosphorus per gram

 $min = minutes$ 

 $mL/min =$  millilitre per minute

 $m/h$  = meter per hour

 $Mg0$  = zero-valent magnesium

 $\mu$  = Solution viscosity

 $N =$ nitrogen

 $Na<sup>+</sup> = sodium$ 

NaCl = sodium chloride

NaOH = sodium hydroxide

 $Na<sub>2</sub>SO<sub>4</sub> = sodium sulphate$ 

 $Na<sub>2</sub>CO<sub>3</sub>=$  sodium carbonate

 $NaHCO<sub>3</sub> = sodium bicarbonate$ 

 $NH<sub>3</sub> = Ammonia$ 

 $N_0$  = saturation adsorbate concentration (mg/L)

n = Freundlich constant

pH = measure of the acidity or basicity of an aqueous solution

 $Q =$  Flow rate (cm3/s)

 $q =$  surface concentration at any radial distance (r) (mg/g)

 $q_e$  = loading of potassium on the resin at equilibrium (mg/g),

 $q_e$  = amount of adsorbate adsorbed per unit mass of adsorbent (mg/g)

- $q_{max}$  = maximum amount of adsorbate adsorbed per unit mass of adsorbent (mg/g)
- $q_0$ = equilibrium adsorbate uptake per g of adsorbent (mg/g)
- rpm = Revolutions per minute
- SEM = Scanning electron microscopy
- SFS = Steel Furnace Slag
- M-SFS = Modified Steel Furnace Slag
- $SO_4^2$  = sulphate
- $t =$  filtration time (min).
- $\tau$  = the time required for 50% adsorbate breakthrough (min)
- $t =$  the time (min)
- $\tau$  = the tortuosity
- $v =$  the linear velocity (cm/min)
- $V =$  volume of the solution (L)
- $Z =$  the bed depth of column

### **ABSTRACT**

Wastewater treatment is an imperative requirement in order to meet effluent quality standards before wastewater can be discharged directly into the rivers, streams and the ocean. Together with the rapid development of industry and society, large volumes of wastewater with hundreds of toxins and impurities, if not treated properly, can severely affect humans and environmental health through different manners such as drinking water contamination and habitat degradation. In particular, the excessive presence of phosphate in water encourages the growth of algae that then leads to eutrophication. Thus, wastewater treatment systems, including phosphate removal, are considered to be a principal tool to reduce the negative effects of wastewater.

The main objectives of this study were to evaluate (i) the capabilities of two commercial ion exchange resins - Purolite A860S, Dowex 21K XLT and a low cost adsorbent - steel furnace slag (SFS) on phosphate removal, and (ii) the effect of modification of these materials on phosphate removal capacities. In this study, two modified materials, Dowex 21K XLT\_Cu and modified steel furnace slag (M-SFS) were developed from original Dowex 21K XLT and steel furnace slag respectively.

The performance of these ion exchange resins on phosphate removal was evaluated in different experimental conditions such as varying adsorbent dose, initial adsorbate concentration, contact time and pH in a series of batch studies. The Langmuir and Freundlich models were found to fit with the experimental data of all resins and slag. The results also show that the modification of ion exchange resin and slag led to a significant improvement of phosphate removal efficiency.

The maximum phosphate adsorption capacities for Dowex 21K XLT\_Cu, Dowex 21 K XLT and Purolite A860S were 97.09; 65.35 and 52.63 mg/g respectively. The adsorption kinetic of these resins was also fitted well with the Pseudo-Second-order kinetic model  $(R<sup>2</sup> = 0.99)$ . The breakthrough data was reasonably described by using two empirical models: Thomas and Yoon-Nelson ( $R^2 \ge 0.93$ ). Phosphate adsorbed by resin was effectively desorbed by using 0.5M NaOH. The adsorbent capacity of resin was found to be recovered well after three adsorption/desorption cycles.

This study also investigated the phosphate adsorption capacity of low cost materials, SFS and M-SFS. The phosphate adsorption capacities of SFS and M-SFS were 22.88 and 30.49 mg/g respectively, lower than those of the resins. For SFS and M-SFS, Langmuir isotherm model gave a better fit than the Freundlich model. The results also revealed that Thomas model can be used to describe the behaviour of the phosphate adsorption on SFS and M-SFS in the filter column test.

In conclusion, the results show that Dowex 21K XLT Cu was the most effective adsorbent for Phosphate removal. It can be regenerated and reused effectively. When the treatment cost is considered, SFS and M-SFS could be the potential adsorbents.