



BIOSORPTION OF HEAVY METALS BY FLYASH THROUGH MULTICOMPONENT STUDY

Ahmad Ashfaq¹, Mohd Kaifiyan²

¹Civil Engineering Section, ²Faculty of Engineering & Technology
(Aligarh Muslim University, A.M.U., Aligarh, India)

ABSTRACT

The release of large quantities of heavy metals into the natural environment has resulted in a number of environmental problems. The treatment technologies that have been used for treating heavy metal bearing waste streams, such as chemical precipitation, adsorption, ion-exchange, and solvent extraction, have been found to be limited, involve high capital and operational costs and may also be associated with the generation of secondary wastes that also present treatment problems biosorption of heavy metals offers a potential alternative to existing methods for detoxification and recovery of these components from industrial wastewaters and is a subject of extensive studies. The biosorption of Iron(II) and Zinc(II) both ions in combination, by flyash was studied in a batch system as a function of initial pH and dual-metal ion concentrations. The working pH values were determined to the selective uptake of metal ions.

Keywords: *Biosorption, Environment, Flyash, Heavy metals.*

I. INTRODUCTION

The main techniques used to remove heavy metal ions from aqueous streams include ion-exchange chromatography, reverse-osmosis, chemical precipitation, and adsorption. Adsorption as a wastewater treatment process has aroused considerable interest during recent years. Adsorption has evolved as the front line of defense for metal ions, which cannot be removed by other techniques. The adsorption of metals by activated carbon is more complex than for organic compounds because the ionic charges affect their removal rates from solution [2]. Metal ion adsorption by flyash varies with the chemical properties of adsorbate, temperature, pH, ionic strength of the liquid phase, etc. Many activated carbons are available commercially but few are selective for heavy metals and they are expensive. Due to the expense of carbon for water treatment, a search for substitutes is under way. Wastewater treatments require vast quantities of activated carbon, so improved and tailor-made adsorbents are needed for these demanding applications. Such adsorbents should be easily available, economically feasible, and readily and quantitatively regenerated chemically [2]. Since industrial effluents can contain several metals, it is necessary to study the simultaneous sorption of two or more metals and also to quantify the interference of one metal with the sorption of the other. In this research, we investigate the feasibility of coal fly ash to remove heavy metal ions [Zn(II), Fe(II)] from water, modeling the adsorption process of heavy metal ions and investigation of the interactive effects of process parameters such as biosorbent dose, contact time, solution pH, and initial concentration on the sorption capacity towards the metal ions.



II. MATERIAL & METHODS

Efforts have been made to find out the sorption behavior of the flyash exposed to bimetallic solution. Metal solution contained bimetallic in the concentration range of 10-100 mg^l⁻¹ for each metal. Batch equilibrium sorption experiments were carried out using standard batch methodology. Samples were withdrawn at predetermined time intervals and the metal sorbed was determined by subtracting the metal left in the solution after 2.5 h of biosorption from the initial metal ion concentration. In order to determine sorption characteristics of Fe (II) in binary metal mixture, the initial concentration was varied from 10 to 100 mg/l while the Zn concentration in each biosorption medium was held constant at 10, 20, 50, 80 & 100 mg^l⁻¹. The pH of biosorption media was adjusted to the optimum value for the biosorption of Zn with 1 N HCl and 1 N NaOH.

III. RESULTS AND DISCUSSION:

The simultaneous removal of Zn(II) and Fe(II) onto flyash was evaluated. In order to have the selectivity for Zn(II) ions, the pH of biosorption media was adjusted to 6.4, which was considered as the optimum value for the biosorption of metal ions. In the first stage of biosorption studies, the initial Zn(II) concentration was varied from 10 to 100 mg/L, while the initial Fe(II) concentration was held constant at 10, 20, 50 or 60 mg/L. The non-linearized biosorption isotherms of Zn(II) ions in the absence of and in the presence of increasing concentrations of Fe(II) ions are shown in. It can be observed from the figures that the equilibrium Zn(II) uptake increased (3.16-31.92 mg/g) by increasing the initial Zn(II) concentration up to 100 mg/L. In addition, the presence and increasing Fe(II) concentration resulting in shortening of biosorbed Fe(II) (17.38 to 13.04 mg/g) quantities, at equilibrium. The inhibitory effect of Zn(II) ions on the equilibrium Fe(II) uptakes was dominating. Although Zn(II) ions were preferentially biosorbed, Fe(II) ions were also biosorbed by the flyash. This situation shows the partial competition between the Zn(II) and Fe(II) ions. Under study sorption, isotherms of Zn(II) and Fe(II) in the presence of constant concentrations of other metal (10, 20, 50 & 60 mg/g) were computed from the empirical Freundlich isotherm and the comparison between experimental and computed.

IV. CONCLUSIONS

From the experimental data of biosorption of Fe(II) and Zn(II) ions onto flyash, the following points can be concluded. The sorption capacity of metal ions was found dependent on initial pH, contact time, sorbent amount and metal concentration of heavy metal. The maximum sorption capacity of flyash for removal of heavy metals was 15.88 mg/g for Fe(II) and 12.74 mg/g for Zn(II).

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