Contents lists available at SciVerse ScienceDirect



Chemical Engineering Journal

Chemical Engineering Journal

journal homepage: www.elsevier.com/locate/cej

Adsorption performance and mechanism of Cr(VI) using magnetic PS-EDTA resin from micro-polluted waters

Ning Mao, Liuqing Yang, Guanghui Zhao, Xiaoli Li, Yanfeng Li*

State Key Laboratory of Applied Organic Chemistry, College of Chemistry and Chemical Engineering, Institute of Biochemical Engineering & Environmental Technology, Lanzhou University, Lanzhou 730000, PR China

HIGHLIGHTS

- ► A novel magnetic chelating resin with EDTA functionality was prepared.
- ▶ It shows easy separation and excellent adsorption for Cr(VI) in aqueous solution.
- ▶ 100% of Cr(VI) could be removed at the low metal concentration (<40 mg/L).
- ► XPS shows that the mechanism contains Cr(VI) reduction and electrostatic interaction.
- ► A conceptual model for Cr(VI) adsorption by this resin is proposed to illustrate the mechanism.

ARTICLE INFO

Article history: Received 6 April 2012 Received in revised form 20 June 2012 Accepted 20 June 2012 Available online 28 June 2012

Keywords: Magnetic resin Hexavalent chromium Adsorption

ABSTRACT

Adsorption of Cr(VI) from aqueous solution onto a magnetic chelating resin with EDTA functionality (magnetic PS-EDTA) was investigated in a batch system. Various factors affecting the uptake behavior such as pH, contact time, initial concentration of the metal ions and dosage of resins on Cr(VI) removal were studied. The magnetic modified resin showed higher adsorption capacity and shorter adsorption equilibrium time for Cr(VI) compared with the raw PS-EDTA resin. The equilibrium data were analyzed using the Langmuir, Freundlich and Tempkin isotherm models among which Langmuir isotherm model was found to be suitable for the monolayer adsorption process. The monolayer adsorption capacity values of 123.05 mg/g for raw PS-EDTA and 250.00 mg/g for magnetic resin were very close to the maximum capacity values obtained at pH 4.0. Kinetic studies showed that the adsorption followed a pseudo second-order reaction. The mechanism was further identified by fitting intraparticle diffusion was also involved in adsorption. XPS spectra confirmed that reduction of Cr(VI) by Fe₃O₄ nanoparticle on the resin occurred, while the electrostatic interaction between protonated amine groups and Cr(VI) anion played an important role in the adsorption. Furthermore, the resin could be regenerated through the desorption of the Cr(VI) anions using 0.5 M NaOH solution and could be reused to adsorb again.

© 2012 Elsevier B.V. All rights reserved.

1. Introduction

Many industries such as electrodeposition, leather manufacturing, steel production and wood preservative industries generate a huge amount of toxic metals. The discharge of such effluents cause increased adverse effects on human beings and environment. Chromium is one of the most important toxic metals which could damage upper respiratory tract and has chronic toxicity [1]. Chromium exists in trivalent (Cr(III)) and hexavalent (Cr(VI)) states. Cr(VI) has been considered more hazardous to public health due to its mutagenic and carcinogenic properties [2]. It is a relatively strong chemical oxidant and could react with the enzymes of the body or biological systems, resulting that its toxicity is about 300 times more than Cr(III) [3].

Because of its high toxicity, Cr(VI) must be substantially removed from the wastewater before being discharged into the aquatic system. Various methods of chromium removal include chemical reduction, filtration, precipitation, adsorption, electrodeposition and membrane systems or even ion exchange process [4]. Among these technologies, adsorption may be considered as preferable in many cases due to its outstanding simplicity, high efficiency, low investment, and potential recovery and reuse of metals [5]. In recent years, investigations have been carried out for low cost, non-

^{*} Corresponding author.

E-mail address: liyf@lzu.edu.cn (Y. Li).

^{1385-8947/\$ -} see front matter @ 2012 Elsevier B.V. All rights reserved. http://dx.doi.org/10.1016/j.cej.2012.06.082