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Continuous Removal of Cr(VI) by Lab-Scale Fixed-Bed Column Packed with Chitosan-Nanomagnetite Particles

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Hexavalent Chromium species are classified as hazardous compounds due to their high toxic potential, considering also their remarkable solubility and redox potential. Various processes have been developed to remove/recover Cr(VI) species from polluted groundwater, such as membrane processes, ion-exchange and adsorption and chemical or biochemical reduction. Indeed, the reduction/removal process of Cr(VI) through iron-based materials usually leads to a pH increase of the reaction medium, allowing to facilitate the subsequent precipitation of the Cr(III) species. In this context, the use of iron based nano-particles (IBNs) supported on bio-polymer matrix allowed to maximize the Cr(VI) removal capacities of iron-based materials, leading to the production of high active and eco-compatible nano-materials. The use of chitosan as surface-modified agent, allows the reduction of aggregation forces among the produced IBNs, leading to higher surface active areas and chemical reactivity. At the same time, the use of a bio-polymer increases the eco-compatibility of the IBNs, reducing the possible interaction with bacteria and microorganisms during the treatment process. In this work chitosan-nanomagnetite particles were synthetized and employed as packing material inside fixed-bed lab-scale column (height 25 cm and diameter of 1.5 cm) to remove, in continuous, Cr(VI) species from synthetic wastewaters. The tests were performed at different inlet flow-rate values (2, 5 and 7 mL/min) at fixed Cr(VI) initial concentration (20 mg/L) and varying the solution pH (pH=4 and 7). The obtained breakthrough curves were then modeled according to the classical dynamic Thomas model.

* 1. Introduction

The use of nanoparticles in various sectors, such as in civil (Di Palma et al., 2015), in the cosmetic (Stoller et al., 2017a), in the catalyst-industrial (Chinh et al., 2018) and in the environmental one (Vilardi et al., 2017a) has been extensively studied, as demonstrated by the increasing number of publications on this field (Bavasso et al., 2016). Various researchers successfully tested the use of metallic nanoparticles for the removal of various pollutants, such as heavy metals (Gueye et al., 2016), organic compounds (Vilardi et al., 2018a) and inorganic anions (Muradova et al., 2016). Heavy metals pollution is considered an environmental problem of great concern (Di Palma et al., 2007) and in particular hexavalent chromium still represents a severe environmental problem in the Mediterranean Area (Vilardi et al., 2018b) since it is characterized by a high carcinogenetic and remarkable solubility and mobility in the environment. The most suitable processes for its removal are represented by adsorption (Bavasso et al., 2018), membrane treatments (Di Palma et al., 2018a), also combined with other treatments (Stoller et al., 2018a) and reduction/precipitation processes (Vilardi et al., 2018c). The adsorption process has already demonstrated its suitability for the removal and recovery of various heavy metals (Lu et al., 2017), through the use of classical or biological waste materials (Vilardi et al., 2018d). Regarding the latter process, it consists of reducing Cr(VI) to the low toxic and low soluble Cr(III) species, that usually tend to co-precipitate in mixed iron or aluminum hydroxides (Vilardi et al., 2017b). Considering the increasing necessity to develop environmentally friendly and economic technologies for a sustainable remediation procedure of polluted environments (Stoller et al., 2016), the production of nanocomposites (Stoller et al., 2018b), constituted by both organic and inorganic compounds, may play a fundamental role to reach such outstanding goals (Chinh et al., 2019). The use of iron-based nanoparticles has already proved to be very effective towards various contaminants in very complex wastewater (Vilardi et al., 2019a), such as tannery wastewaters (Vilardi et al., 2018e), heavy metals-polluted groundwater (Vilardi et al., 2018f) and in the presence of both nitrates and Cr(VI) pollutants (Maharramov et al., 2017). Therefore, the production of iron-based bio-nanocomposite represents an important field of research to develop environmentally friendly and effective materials for the removal of different pollutants in aqueous media (Vilardi and Di Palma, 2017). The aim of this work was to investigate on the removal efficiency of Cr(VI) using a bio-nanocomposite material, produced using chitosan as bio-polymer and nanomagnetite as iron based nanoparticles and employing them as packing material in lab-made column. Kinetic experiments were conducted at different inlet flow-rate values (2, 5 and 7 mL/min) at fixed Cr(VI) initial concentration (20 mg/L) and varying the solution pH (pH=4 and 7). The obtained breakthrough curves were then modeled according to the classical dynamic Thomas model, considering the mass transfer at liquid/solid interface the process limiting step (Vilardi, 2019).

* 1. Materials and Methods
		1. Materials

All the reagents were purchased from Sigma Aldrich (Milan) and were of analytical grade or higher. The solutions were prepared in deionized water. The following reagents were used in the experiments: NaOH, NaCl, chitosan, K2Cr2O7, FeCl3·6H2O, FeSO4·7H2O, NH4OH (33% v/v) Diphenylcarbazide, and H2SO4. The chromium salt was dissolved in deionized water to prepare a solution with an initial Cr(VI) concentration equal to 20 mg/L, whereas the NaOH and H2SO4 solutions were diluted up to 0.1 M and subsequently used to modify the initial Cr(VI) solution pH, measured by a Crison pH-meter (Di Palma et al., 2018b).

* + 1. Bio-Nanocomposite Synthesis

The followed procedure is a modification from Tran et al. (2010), since the spinning disk reactor was used instead of classical stirred tank vessels. In brief, the iron precursors were dissolved in the aqueous chitosan solution (prepared dissolving 2 g of chitosan in 180 mL of deionized water and 20 mL of H2SO4 0.1 M) using a molar ratio of 2:1 among Fe(III) and Fe(II) species (0.1 M of FeSO4·7H2O and 0.2 M of FeCl3·6H2O). The solution was stirred at 300 rpm for 30 min and subsequently the Chitosan-NanoMagnetite particles (CNM) were synthetized in continuous using a Spinning Disk Reactor (SDR) according to the optimal operating parameters values reported in a previous work (Vilardi et al., 2017c). The use of rotating device allows to increase mass transfer rate and to reduce the residence time inside the reactor (Di Palma and Verdone, 2009). In detail, the inlet flowrate of chitosan-iron solution was 150 mL/min, that one of NH4OH solution was 37.5 mL/min (added according to the stoichiometric molar ratio of OH-/Fe(II)=8 mol/mol), the injection point distance over the disk was set to 2 cm and the rotational velocity of the disk was fixed to 146.5 rad/s. The CNM particles were then characterized by Dynamic Light Scattering (Brookhaven), showing a mean dimension of 40 ± 2.5 nm. Point of Zero Charge (PZC), was determined by suspending different material amounts (0.01, 0.1, 1, 5, 10, 20 % wt) in 0.1 M NaCl solution and measuring the solution pH after 24 h of contact time, according to (Chung et al., 2012). The pH of zero charge measured was equal to 4.5, that was really close to the value of 4.8 reported by Chung et al. (2012).

* + 1. Experimental set up

A borosilicate glass column of 25 cm of height and 1.5 cm of internal diameter was used in the continuous experiments (Vilardi et al., 2019b). The column was packed with 11.4 g of the CNM material to reach a bed height h (cm) of 10 cm, corresponding to a bed volume, Vb (mL), of 17.7 mL. The bed porosity, =0.55, was calculated as the ratio between void volume Vv (measured by the volume, mL, of the added deionized water in the packed column at the selected h, according to a previous work (Vilardi et al., 2018g)) and the related Vb. A supporting layer of 1.2 cm of glass wool was placed on the packed bed top, to prevent the sorbent floating. A peristaltic pumps (LAMBDA) was used in up-flow mode at a desired flow-rate (2, 5 and 7 mL/min), to send the Cr(VI) solution through the column (Figure 1).

Six experiments adopting the above mentioned three different inlet flow-rate values (Q (mL/min)) and two different pH value (3 and 7) were conducted to obtain the breakthrough curves for Cr(VI) compound, by fixing the sampling time to 5 min and stopping the column operation until the final Cr(VI) concentration exceeded a value of 95% of the initial Cr(VI) concentration in the synthetic solution. Once the column resulted saturated, the packing material was recovered and substituted with freshly prepared material. At selected time intervals a liquid sample was withdrawn and the CNM particles were separated through an ultra-centrifuge (14000 rpm for 5 min) and the liquid phase was withdrawn to proceed with the Cr(VI) measure through dyphenilcarbazide method (Vilardi et al., 2017b).



*Figure 1: experimental set up scheme.*

* + 1. Data analysis and mathematical modeling

The breakthrough curve shape and the breakthrough time, tb (min), are the key-parameters for the description of the column process dynamic behaviour. The former occurs when the Cr(VI) concentration in the effluent reaches a pre-determined value, usually related to the specific concentration limit for effluent disposal according to the specific environmental regulation (in this study the value of 0.2 m/L was taken into account). Another important parameter is the bed saturation time, te (min), that is the required process time to reach a Cr(VI) concentration in the effluent equal to 0.95Cr(VI)0, where the subscript 0 indicates the initial conditions. The maximum column capacity, qtot (mg), is expressed by the equation reported below:

where Cr(VI)ads (mg/L) is the adsorbed Cr(VI) concentration. The adsorption capacity of the column, q (mg/g), can be calculated as the ratio between qtot and the CNM used mass, m(g).

The data obtained by the performed experiments were then interpreted through the dynamic Thomas model, that assumes: (i) constant separation factor and (ii) that the sorption process follows the Langmuir kinetic mechanism, occurring with negligible axial dispersion, because the process is limited by the mass transfer at the liquid/solid interface and not by the chemical reaction. The model can be described by the following equation:

where KT (mL/min mg) is the Thomas rate constant and q0 (mg/g) is the maximum sorption capacity of the CNM material. The data fitting was accomplished in Excel environment, using the non-linear solver function in order to avoid the errors due to linearization (White and Verdone, 2000).

* 1. Results and Discussion
		1. Influence of pH and Q on process performances

Figure 2 displays the obtained breakthrough curves at different operating parameter values.



*Figure 2: breakthrough curves obtained at different initial Q for pH=3 (a) and pH=7 (b) (h=10 cm, temperature=25°C, Cr(VI)0=20 mg/L, the Cr(VI) concentration reported on y-axis is the Cr(VI) in the outlet of the column).*

The shape of the curves changed with both pH and Q, since at higher Q the residence time among liquid phase and solid phase decreased as also reported in a previous study (Vilardi et al., 2018g) and in Table 1 (see te and tb values). As a consequence, to an increase in the Q value also the tb and te decreased. As regard the pH influence, it can be clearly observed from the graphs reported in figure 2, at pH=7 the curves shape slightly changed but the te values were substantially lower in comparison to those obtained at pH=3 (see also Table 1). The effect of pH on the Cr(VI) adsorption behavior can be explained considering the pH of zero charge: Cr(VI) compounds are present as oxyanions, thus, a positively charged surface of the adsorbent material can foster their adsorption. Since the pH of zero charge of the material is 4.5, for pH<4.5, i.e. pH=3, the surface’s adsorbent resulted positively charged and the adsorption is enhanced. The chemical reduction of Cr(VI) to Cr(III) is possible onto the surface of the sorbent or in the liquid bulk, since the reducing agent is represented by the ferrous ions released by the nanoparticles or present onto the sorbent surface bonded with ferric species in the iron oxide. The reaction causes the oxidation of 3Fe(II) to 3Fe(III) and the reduction of Cr(VI) to Cr(III) species.

Table 1: parameters obtained from breakthrough data analysis (Veff=te\*Q is the total effluent volume treated).

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| --- | --- | --- | --- | --- | --- | --- | --- |
| h (cm) | pH | Q (mL/min) | Veff (mL) | te (min) | tb (min) | qtot (mg) | qe (mg/g) |
| 10 | 3 | 2 | 490 | 245 | 30 | 5.43 | 0.48 |
| 4 | 860 | 215 | 20 | 9.53 | 0.84 |
| 6 | 1110 | 185 | 15 | 13.99 | 1.23 |
| 7 | 2 | 420 | 210 | 20 | 4.61 | 0.40 |
| 4 | 720 | 180 | 15 | 9.53 | 0.84 |
| 6 | 960 | 160 | 5 | 12.82 | 1.12 |

* + 1. Breakthrough data fitting to Thomas model

Figure 3 shows the fitted Thomas model to the experimental data obtained at the optimal pH value.



*Figure 3: breakthrough curves modeling through Thomas model (pH=3, h=10 cm, temperature=25°C, Cr(VI)0=20 mg/L).*

Thomas model was able to well describe the breakthrough data and the asymptotic experimental behavior. The regressed model parameters are reported in Table 2, as well as the correlation coefficient values.

*Table 2: regressed Thomas parameter values and correlation coefficients.*

|  |  |  |  |
| --- | --- | --- | --- |
| Q (mL/min) | KT (mL/min mg) | q0 (mg/g) | R2 |
| 2 | 2.05 | 0.42 | 0.982 |
| 4 | 0.984 |
| 6 | 0.994 |

The obtained KT and q0 values are in line with those obtained in a previous study (Vilardi et al., 2018g), as expected also considering the similar breakthrough curve shapes and te values. The R2 values were close to 0.99, with a higher value for the data from the experiments conducted at higher Q. However, Thomas model demonstrated to be suitable to describe the experimental behavior observed at different inlet flow-rate values.

* 1. Conclusions

The obtained CNM nanoparticles were tested as packing material at fixed bed height (10 cm) and varying the inlet solution pH=3 and 7, whereas the inlet flow-rate was varied as 2, 4 and 6 mL/min. The experimental results showed that the pH influenced the overall process efficiency, since for pH<pH of zero charge the adsorbent’s surface resulted positively charged and the sorption of oxyanions, such as Cr(VI) species, resulted enhanced too. The increase of inlet flow-rate caused a decreased in the contact time between solid and liquid phase, causing a decrease of effluent time and, as a consequence, of the effluent treated volume. Finally, Thomas dynamic model was able to well describe the experimental data and the shape of the obtained breakthrough curves; the regressed model parameter values were in line with the expectations considering also a previous work regarding the Cr(VI) removal in more complex wastewaters.

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