Efficient Adsorption of Trivalent Rare Earth Elements Using Nano Copper Ferrite: Kinetics, Isotherms and Thermodynamics

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ABSTRACT

Rare earth elements (REEs) have versatile applications in various industries and are thrown as industrial waste waters, which leads to serious environmental issues. This paper describes the adsorption efficiency of nano copper ferrite on REEs namely lanthanum and cerium in their trivalent states at room temperature and at the optimum conditions of pH, contact time, adsorbent, adsorbate dosage, temperature etc. The batch adsorption studies were conducted for the adsorption studies using ICPAES (Inductive coupled plasma atomic emission spectrophotometer). In addition the data is studied for Langmuir and Freundlich models and observed that data well fitted to Langmuir monolayer adsorption. From the graph and intercept values of theoretical and experimental values, the pseudo second order kinetics are followed. Thermodynamic parameters reveal that reaction is spontaneous and endothermic. From the over all observations like fast kinetics, chemical stability, high adsorption capacity it is concluded that nano CuFe₂O₄ has maximum efficiency for the adsorption of REEs.

Keywords: Nano spinel copper ferrite, Trivalent rare earth elements, Adsorption, Langmuir adsorption isotherms, Thermodynamic parameters.

INTRODUCTION

REEs consists of lanthanide series and scandium plus yttrium. They have 4f orbital as inner electronic configuration and outer shell has 5d¹⁻¹⁰ and 6s² electrons. This electronic configuration makes lanthanides more special for fast reactions. Their physical characteristic properties are almost close due to similar electronic configuration¹⁻³. They have difference in chemical properties because of lanthanide contraction of ionic radius and electrostatic effect associated with increase of shielded nuclear charge through electrons. The trivalent form {M(III)} is the most thermodynamic stable form⁴⁻⁶. Lanthanides have many applications in various fields such as super conductors, fertilizers, Lasers, catalysts and optical fibers. As they are unable to pass through cell membrane, they are...
non toxic. In the present work La(III) and Ce(III) are selected which are from light REEs family.

Lanthanum is the 57th element in the periodic table, it causes nervous system disorder in humans. Also the petroleum industry waste, which has lanthanum, dumped into waste waters. This causes severe damage to plants and animals. The atomic number of cerium is 58 and the pure metal itself can ignite if scratched with knife. It is strong reducing agent and causes itching, sensitivity to heat and skin lesions are experienced. The main interest in selecting La(III) and Ce(III) is that they are 80% global consumption and also present in industrial waste water. The main aim is to obtain a potential adsorbent for REE extraction from aqueous media. Till date several methods are applied for the extraction of REE mainly using solvent extraction, electro chemical precipitation, Ion exchange and adsorption. Among them adsorption is preferred due to its reversibility, eco friendly nature, cheap, simplicity and hence applicable for the REE ions in aqueous samples. However the investigation of REE from aqueous solution is limited. Adsorption is treated as efficient, quick and low cost procedure in the recovery of metals.

Nano spinel copper ferrites are used for the adsorption of REE from aqueous solution. As these spinel ferrites are famous for their thermal stability, chemical and magnetic properties. They are widely used as catalysts, magnetic fluids and microwave radiation isolators. Nano spinel copper ferrite is prepared using co-precipitation method which is most widely used in the synthesis of magnetic nanoferrites. Co-precipitation is simple with high crystalline and textural properties.

Synthesis of CuFe$_2$O$_4$

Aqueous solution of iron nitrate Fe(NO$_3$)$_3.9$H$_2$O and copper nitrate Cu(NO$_3$)$_2.3$H$_2$O with stoichiometric ratio Cu:Fe = 1:2 were stirred for one hour (for each 1 g of nitrate, 3.3 mL of milli Q water are used. 1 molar NaOH solution is added, for each 1 g of nitrate, followed by further stirring for 15 minutes. The material is annealed for 2 h at 700°C, at ordinary atmosphere in the muffle furnace. This is further characterized using analytical techniques. Concentration of REE metal ion in both initial and with drawn samples are calculated through ICP-AES technique.

Adsorption Experiments

The equilibrium adsorption experiments can be performed individually at room temperature. The batch experiments are carried out to evaluate the adsorption process and REE mobilization. Every time a particular parameter is varied by fixing all other parameters at their optimum values and the effect of temperature, kinetics, isotherms and thermodynamic parameters are calculated. Also the parameters like effect of pH, nano spinel ferrite (adsorbent) dose, REE’s dose, contact time are optimized. However % of REE adsorbed can be calculated using the formula given as

$$\% \text{REE adsorption} = \frac{(\text{Initial O.D} - \text{Final O.D})}{\text{Final O.D}} \times 100$$

here O. D = optical density of REE

RESULTS AND DISCUSSION

Effect of pH

The pH is key factors during the adsorption process. The amount of REE’s adsorption onto nano copper ferrite was studied at pH=2 to 11. Results showed that when pH was higher than 7, the adsorption capacity was maximum. At lower pH, the adsorption capacity was not maximum, but the results are in agreement with the observations.
of other researcher’s group. At lower pH values the protons are in competition with adsorbates (REEs) and nano copper ferrites retain very small amount of lanthanides, as a result the adsorption of La(III) and Ce(III) is very less. Also at when pH is increased adsorbate-adsorbent electrostatic repulsion decreases and adsorbed Lanthanides seems stable. A pH value of 7.8 is fixed as optimum for the entire study.

**Adsorbent dosage (nano spinel copper ferrite)**

The nano spinel copper ferrite dosage is varied from 0.1 to 1 g under the optimum conditions of other parameters (pH = 7.8, contact time = 30 min for both REEs and at room temperature). Relationship between nano spinel copper ferrite dosage and removal efficiency of the REE from their aqueous solutions is presented in Fig. 3. Nano spinel copper ferrite shows maximum removal efficiency of more than 90% ions at 500 mg. The rate of adsorption increases with dose of adsorbent is due to availability of more adsorption sites with further growth of surface area. This reduced with increase in adsorbent dose. Overlapping of adsorption sites as a result of overcoming of copper ferrite particles. By increasing the initial adsorbent concentration in solution, the quantity of adsorbed lanthanides gradually increases. But increase starts to ease after 0.2 g/L. Trend is probably because of the adsorption sites saturation at higher concentrations.

**Effect of contact time**

Adsorption capacity increases with time can be explained in 2 steps. That is in first 20 min a steep elevation in the sorption. Around 95% of the overall sorption takes place with in first 30 min of contact time. Proper reaction rate, diffusion mechanism are very significant in explaining the role of contact time.

**Dose of REEs**

Effect of REEs dosage on adsorption can be done by preparing aqueous solution of lanthanum nitrate and cerium nitrate respectively. The range is from 5ppm to 30ppm. The dose of nano copper ferrite is 0.5 g throughout. The total volume is 30.0 mL. The temperature of the system is 300K with a pH of 7.8. The solution was centrifuged and the supernatent liquid concentration is observed using ICPAES, with time until the system approaches equilibrium. The same procedure is repeated until the error is minimized and attains reproducibility. The maximum adsorption was observed at 10 and 8ppm respectively for the REE ions. This is because, establishment of saturated more ionic layer at lower concentration of metal ions, makes extra interaction near the adsorbent hindered in high concentration solution due to the interactions among REE cations.
existence on the copper ferrite medium. Also, tendency for an ion to remain in Stern layer depends on the electro porosity of the metal ion.

Spinel copper ferrite at various conditions and concentrations, to design and optimise an operating procedure, the analysis of equilibrium data is very important, because the adsorption is a mass transfer operation and is elaborated by a rate process. However, to examine the characteristics of adsorption process, kinetic data, potential rate controlling data are analyzed using pseudo first order and pseudo second order. They can be represented as

\[ q_t = q_e \cdot e^{-k_1t} \]  
\[ q_t = \frac{tq_2^2k_2}{1 + tq_2k_2} \]

\( k_1 \) is the rate constant for pseudo first order kinetics.

\( k_2 \) is the rate constant for pseudo second order kinetics.

From reports, the kinetics of the elements do follow pseudo second order kinetics for both REEs. However, R² for pseudo second order at different concentrations are above 0.99. Adsorption is chemisorption through sharing of electrons among copper ferrite and adsorbate. From this data, it is suggested that adsorption rates are quick at lower concentrations for lanthanides. In other words, time required for the equilibrium adsorption increases as RE ions concentration increases.

### Kinetic models

To compare the efficiency of nano adsorption isotherm models

Table 1: Parameters for PFO and PSO kinetics for the adsorption of REE on nano spinel copper ferrite

<table>
<thead>
<tr>
<th>Parameter</th>
<th>La(III)</th>
<th>Ce(III)</th>
<th>La(III)</th>
<th>Ce(III)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ci (mg/L)</td>
<td>10</td>
<td>8</td>
<td>10</td>
<td>8</td>
</tr>
<tr>
<td>K₁ (min⁻¹)</td>
<td>0.026</td>
<td>0.024</td>
<td>2.45x10⁻⁴</td>
<td>2.37x10⁻⁴</td>
</tr>
<tr>
<td>qₑ (mg/g)</td>
<td>119.6</td>
<td>116</td>
<td>116</td>
<td>112.4</td>
</tr>
<tr>
<td>R²</td>
<td>0.980</td>
<td>0.986</td>
<td>0.996</td>
<td>0.996</td>
</tr>
</tbody>
</table>

Fig. 4. Dosage of La(III) on adsorption by nano spinel copper ferrite

Fig. 5. Dosage of Ce(III) on adsorption by nano spinel copper ferrite

Fig. 6. Pseudo first order kinetics for the adsorption of REE on nano spinel copper ferrite

Fig. 7. Pseudo second order kinetics for the adsorption of REE on nano spinel copper ferrite

### Adsorption isotherm models

To establish equilibrium data of adsorption, the experimental data is studied with respect to Langmuir and Freundlich models. From the Table 2,
the experimental data is fitted for Langmuir model and hence adsorption is mono layer\textsuperscript{27}. Due to the ionic radius closeness of La(III) = 1.051\text{Å} and Ce(III) = 1.034\text{Å}. The qm values for REEs is almost equal. As the Langmuir model assumes mono layer adsorption the data fits more into this. The linearised form is

\[ q_e = \frac{Q_0k_Fc_0^{1/n}}{1+k_Fc_e} \]

\( q_{\text{max}} = \text{adsorption capacity (µg g}^{-1}\text{)} \) and b is energy of adsorption.

\( q_e = k_Fc_e^{1/n} \)

\( k_F \) is adsorption capacity and “n” is adsorption intensity.

\( K_L \) and \( q_m \) are from linear fit of \( c/c_e vs c_e \). Langmuir model is the most widely used and covers monolayer adsorption. This model is appropriate with specific parameters for adsorption of REEs on copper ferrites\textsuperscript{28}. Maximum mono layer adsorption capacities are found to be 333 and 345 mg/g for lanthanum and cerium respectively. However E values from 8 to 16 KJ/mol explain chemical adsorption.

\[ \Delta G^0 = -RT\ln K_L \]

\[ \ln K_L = \frac{-\Delta H^0}{RT} + \frac{\Delta S^0}{R} \]

\( \Delta G^0 \) is the change in Gibb’s free energy, \( R \) ideal gas constant (8.314J/mol/K), T absolute temperature (K), \( K_L \) Langmuir constant. \( H^0, S^0 \) are changes in enthalpy and entropy respectively. Effect is examined on the adsorption of REE by nano copper ferrite at the range of 300K 310K and 320K at pH 7.8 and constant time of 60 minutes. The concentration of La (III) and Ce(III) are 10 and 8 ppm respectively at the concentration of adsorbent is at 0.5 gram.

Both the REE are temperature dependent and at higher temperature the increase in the dose of adsorbed lanthanides is because of the higher affinity of the available active sites. Positive values of enthalpy change (\( H^0 \)) decrease in the Gibb’s free energy change with rise of temperature encourages sorption mechanism is endothermic\textsuperscript{29}.

More over positive values of entropy change (\( \Delta S^0 \)) means disorder of the system increases after metal sorption. Increase of entropy is due to release of water during the metal sorption of RE ions. This fact that \( \Delta H^0 \) is less than \( T\Delta S^0 \). This explains that sorption process is through entropy rather than enthalpic changes\textsuperscript{30}.

The thermodynamic studies are more important to understand the mechanism of adsorption for the practical applications. To study these effects Gibb’s free energy and Van’t Hoff equation are followed.
The capacity of 983 and 973 mg/g for La(III) and Ce(III) respectively. The optimum conditions for the parameters are obtained. The pH of 7.8 is opted for both the elements to reach the equilibrium in 30 minutes. Langmuir isotherm model is fitted well followed by pseudo second order kinetics. The system is thermodynamically spontaneous, endothermic and randomness increases with temperature. Hence to conclude the adsorption is economical, environmental benign procedure.

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Conflict of interest

Authors have no conflict of interest regarding work and publication.

REFERENCES


