Fabrication of perovskite-type oxide nanopowders as a novel adsorbent in removal of Bromo Thymol Blue dye from aqueous solutions

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Article published on April 30, 2015

Key words: Nanopowder, Perovskite-type oxide, Bromo thymol blue, Isotherm.

Abstract

Nanoparticles of perovskite-type Gd$_{0.5}$Ca$_{0.5}$CrO$_3$ was investigated for its ability to perform as a suitable sorbent for anionic dye from aqueous solution. The nanoperovskite were characterized using X-ray powder diffraction (XRD), Fourier transform infrared spectroscopy (FTIR) and scanning electron microscopy (SEM). The effects of various experimental factors; adsorbent dose, contact time, pH and dye concentration were studied by using the batch technique. Experimental results indicate that the prepared Gd$_{0.5}$Ca$_{0.5}$CrO$_3$ nanopowder can remove 91.38% of BTB dye under optimum operational conditions of a dosage of 0.02 g, pH 2, contact time of 25 min and initial dye concentration of 50mg/L. Adsorption isotherms have been modeled by Langmuir, Freundlich, Temkin and Dubinin-Radushkevich (D-R). The Langmuir model displayed the best fit for the isothermal data. The maximum predicted adsorption capacities were 300 mg/g for Bromo Thymol Blue (BTB) dye. The results showed that the sorption kinetics of BTB removal from aqueous solutions onto nanopowder fitted well with the pseudo-second-order model.

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Introduction
The family of perovskite-type oxides generally formulated as ABO$_3$ (A is a rare earth metal with large ionic radius or alkali earth metals; B is a transition metal with a small ionic radius) Significant catalytic applications of these perovskites include reactors for partial and total oxidation of hydrocarbons and in devices for the removal of pollutants from combustion gases (Tejuca et al., 1993; Schaak et al., 2002) and could be considered as an adsorbent/catalyst material for the removal of dyes (Jeong et al., 2006; Carbajo et al., 2006).

Dye effluents from textile industries and photographic industries are becoming a serious environmental problem because of their toxicity, unacceptable color, high chemical oxygen demand content, and biological degradation (Siddique et al., 2009). Dyes can be classified as anionic (direct, acid, and reactive dyes), cationic (basic dyes) and non-ionic (disperse dyes) (Mishra and Tripathy, 1993). Waste waters offer considerable resistance for their biodegradation due to presence of these heat and light stable dyes, thus upsetting aquatic life (Wong and Yu, 1993). Several treatment methods including coagulation, chemical oxidation, membrane separation and adsorption techniques have been proposed for the treatment of dye waste water. Considerable amounts of the literature reports the adsorption of dyes on various adsorbent surfaces especially nanomaterials (Yazdaniakhsh et al., 2010; Wu et al., 2004).

The first aim of the present study is to fabricate and characterize of perovskite-type oxide nanoparticles Gd$_{0.5}$Ca$_{0.5}$CrO$_3$. Investigation of the efficiency of the perovskite-type oxide nanoparticles as adsorbent for the removal of Bromo Thymol Blue dye (BTB), from aqueous solutions is the second goal and to explain the adsorption of BTB onto nanoparticles Gd$_{0.5}$Ca$_{0.5}$CrO$_3$ using isotherm models.

Material and methods
Reagents
The chemicals and reagents used in this work include Gd(NO$_3$)$_3$.6H$_2$O (99.9%), Ca(NO$_3$)$_2$.4H$_2$O (99.9%) and Cr(NO$_3$)$_3$.9H$_2$O (99.9%) obtained from Merck, Germany; citric acid (CA) (99.5% purity), was purchased from Aldrich, USA. 2.1. The chemical structures of Bromo Thymol Blue (BTB) dye is given in Fig. 1.

![Fig. 1. Molecular structures of Bromo Thymol Blue (BTB) dye.](image)

Preparation of Gd$_{0.5}$Ca$_{0.5}$CrO$_3$ powders
For preparing the nanoperovskite Gd$_{0.5}$Ca$_{0.5}$CrO$_3$, proportional amounts of Gd(NO$_3$)$_3$.6H$_2$O, Ca(NO$_3$)$_2$.4H$_2$O, and Cr(NO$_3$)$_3$.9H$_2$O were dissolved in 40 ml of deionized water. Then, citric acid (CA) was added to the metal solution by stirring at room temperature. The solution was refluxed with stirring for 90 min to convert it to a stable complex. The gel was heated at 80°C for 3 h and then dried at 112°C for 19 h. It was grounded in an agate mortar to change into a powder and calcinated at 700°C in air for 8 h.

Characterization methods
The dried powders were analyzed by Fourier transform infrared spectroscopy using thermo Nicollet Nexus 870 FTIR spectrometer. The crystallization and microstructure of the oxide powders were characterized with XRD in a 2θ range from 20 to 80°, using CuK a radiation (λ = 1.5418 Å) on a Rigaku D/MAX RB XRD diffractometer equipped with a curved graphite monochromator. A scanning electron microscope (SEM, LEO1450 VP, V = 26 kV) was used to study the morphology of the calcined powders.
Dye removal experiments
Dye adsorption by the GCCO nanopowder was determined in batch experiments at different contact times (2–30 min), dye concentrations (50-300 mg/L), different doses of adsorbent (0.005-0.02 g) and pH of solution (1–12). The initial dye solution concentration for all batch experiments was adjusted to 50 mg/L except for the experiments in which the effect of the initial dye concentration was tested. All experiments were conducted at 25°C.

Results and discussion
FT-IR studies
Figure 2a and b shows the FT-IR spectra of the dry gel before calcination and calcined powder at 700°C for 8 h, respectively. As shown in Fig. 2a, the broad absorption band around 3415 cm⁻¹ corresponds to the \( \nu_{\text{O-H}} \) stretching vibrations of water molecules (Kuznetsov et al., 2002) and characteristic bands at about 1226 and 1384 cm⁻¹ corresponds to the stretching mode of M–O–C group (Kuznetsov et al., 2002, Kim and Honma, 2004) and the anti-symmetric \( \nu_{\text{NO}} \) stretching vibration (Liu et al., 2007), respectively. The absorption band around 1081 cm⁻¹ is assigned to C–O bond (Nakamoto, 1978). In the FT-IR spectrum of the GCCO nanopowder (Fig. 2b) a strong peak appears at 590 cm⁻¹, which is assigned to the M–O bond.

X-ray diffraction studies
Fig. 3 shows the XRD patterns of the GCCO nanopowder prepared by sol–gel method. The diffractograms reveal that the crystalline perovskite structure is the main phase for synthesized nanopowder. The most intense diffraction peak (2θ = 32.58) is a single peak for all of the studied perovskites (Merino et al., 2006). The crystalline sizes of the GCCO nanopowder were determined by means of an X-ray line-broadening method using the Scherrer equation:

\[
D_{\text{hkl}} = \frac{0.9\lambda}{\beta_{\text{hkl}} \cos \theta_{\text{hkl}}}
\]

Where \( D_{\text{hkl}} \) is the particle size in nanometers, \( \lambda \) is the wavelength of the radiation (1.54056 Å for CuK\( \alpha \) radiation), \( k \) is a constant equal to 0.9, \( \beta_{\text{hkl}} \) is the peak width at half-maximum intensity and \( \theta \) is the peak position. The crystalline size of the GCCO nanopowder calcined at temperature of 700°C was found to be 30 nm.

Fig. 2. (a) FT-IR spectra of the GCCO dry gel before calcination. (b) FT-IR spectra of the calcined powders at 700°C.

Fig. 3. The XRD patterns of the GCCO nanoparticles at 700°C.

Microscopic analyses
Scanning electron microscopy (SEM) of the GCCO nanoparticles prepared by the sol–gel method is
shown in Fig. 4. This image exhibits typical morphologies for prepared powder. Based on the SEM images, porosity of the surface is evident and it seems that the particles have not grown with uniform size. The particles size of GCCO that were propagated on the surface seems to be in the range of 29–97 nm.

**Fig. 4.** SEM images of GCCO nanoparticles.

**Effects of various conditions on adsorption**

**The Effect of pH on dye adsorption**

One of the most important factors in adsorption studies is the effect of acidity on the medium (Calvete et al., 2010). Different species may present divergent ranges of suitable pH depending on which adsorbent is used. The effect of initial pH on the adsorption capacity of BTB dye using GCCO adsorbent was evaluated within pH range between 1 and 12 with a stirring time of 25 min. The initial concentrations of dye and adsorbent dosage were set at 50 mg/L and 0.02 g, respectively. The percentage of dye removal is defined as:

\[
\text{Removal rate } \% = \frac{C_0 - C_e}{C_0} \times 100
\]

where \(C_0\) and \(C_e\) represent the initial and equilibrium concentrations of dye at given time, respectively. Fig. 5 shows that the effect of pH on the adsorptive removal of BTB by GCCO nanopowder. The percentage removal of BTB (anionic dye) was maximum at acidic pH (pH 2) and decreased with further increase in pH. BTB dye is an acidic dye that has negative charge in solution, which favors electrostatic interactions between the GCCO surface and dye molecules leading to the strong adsorption. Therefore, to have the optimized condition to remove BTB, acidic pH should be applied and pH 2 seems to lead to the best result; so this pH was selected to run further experiments.

**Fig. 5.** Effect of pH on the removal of dye by GCCO nanopowder (contact time = 25 min, initial dye concentration = 50 mg/L, GCCO dosage: 0.02 g).

**Effect of adsorbent amounts and contact time**

By increasing amounts of GCCO nanopowder, the removal efficiency increased due to increase in the accessible sites for the adsorption of the analyte. Fig. 6 shows that the removal efficiency of BTB dye with the initial dosage of 0.005 g, increased from 63.40 % at the second minute of contact to 87.33% at time equals to 30 min by keeping constant stirring, however, with increasing GCCO dosage to 0.02 g, obtained removal efficiency in the second minute of stirring was 86.46%, and in the 25 minute of stirring was 91.38%. In addition, increasing the contact time from 2 to 30 min with different dosages of adsorbents led to decreasing in the concentration of BTB dye.

**Fig. 6.** Effect of stirring time on removal of dyein different doses (●) 0.005 g, (●) 0.01 g, (▲) 0.02 g of GCCO (initial dye concentration=50 mg/L, initial pH = 2).
**Effect of dye concentration**

The initial dye concentration is another important variable that can affect the adsorption process. The effect of initial BTB concentration on dye removal efficiency by GCCO nanoparticles was studied by varying the initial dye concentration from 50 to 300 mg/L at pH 2, a catalyst dosage of 0.02 g and contact time of 25 min, as shown in Fig. 7. Results show that removal of textile dye BTB decreases with increasing initial concentration. As it is obvious, the percentage removal of BTB decreased from around 91.38% at a concentration of 50 mg/L to 48.28% when the concentration was increased to 300 mg/L. This behavior reveals the dependency of adsorption to initial concentration of BTB.

![Fig. 7. Effect of initial adsorbate concentration of on removal of dye by GCCONanopowder (contact time = 25 min, GCCO dosage = 0.02 g, pH = 2).](image)

**Adsorption isotherms**

The equilibrium isotherm of a specific adsorbent represents its adsorptive characteristics and analysis of isotherm data is so important to predict the adsorption capacity of the adsorbent, which is one of the main parameters required for designing the adsorption processes (Afkhami and Moosavi, 2010). The amount of dye adsorbed onto GCCO nanoparticles has been calculated based on the following mass balance equation:

\[ q_e = \frac{V(C_0 - C_e)}{m} \]

Where \( q_e \) was the equilibrium adsorption capacity of dye adsorbed on unit mass of nanoparticles (mg/g), \( V \) is the volume of the dye solution (L), \( C_0 \) and \( C_e \) were the initial dye concentration (mg/L) and dye concentration (mg/L) at equilibrium and \( m \) (g) is the mass of dry GCCO nanoparticles added.

Analysis of the equilibrium data is important to develop an equation which accurately represents the results and which could be used for design purposes. The equilibrium adsorption data of BTB dye onto GCCO adsorbent was analyzed by the use of well known models given by the Langmuir, Freundlich, Temkinand Dubinin–Radushkevich, isotherm models.

**Langmuir isotherm model**

In its formulation, this empirical model assumes monolayer adsorption (the adsorbed layer is one molecule in thickness), with adsorption can only occur at a finite (fixed) number of definite localized sites, that are identical and equivalent, with no lateral interaction and steric hindrance between the adsorbed molecules, even on adjacent sites (Vijayaraghavan et al., 2006). In its derivation, Langmuir isotherm refers to homogeneous adsorption, which each molecule possess constant enthalpies and sorption activation energy (all sites possess equal affinity for the adsorbate) (Kundu and Gupta, 2006), can be represented as follows:

\[ \frac{C_e}{q_e} = \frac{1}{b q_{max}} + \frac{C_e}{q_{max}} \]

where the \( q_{max} \) (mg/g) is the surface concentration at monolayer coverage which illustrates the maximum value of \( q_e \) and it can be attained as \( C_e \) is increased. The values of \( q_{max} \) and \( b \) can be determined from the linear regression plot of \( (C_e/q_e) \) versus \( C_e \).

**Freundlich isotherm model**

Freundlich isotherm is the earliest known relationship describing the non-ideal and reversible adsorption, not restricted to the formation of monolayer (Freundlich, 1906). This empirical model can be applied to multilayer adsorption, with non-uniform distribution of adsorption heat and affinities over the heterogeneous surface (Adamson and Gast, 1997). The linearized equation is expressed as:
\[
\log q_e = \log K_F + \frac{1}{n} \log C_e
\]

where \(K_F\) and \(n\) are constants of the Freundlich equation. The constant \(K_F\) represents the capacity of the adsorbent for the adsorbate and \(n\) is related to the adsorption distribution.

**Temkin isotherm model**

The Temkin isotherm equation assumes that the heat of adsorption of all the molecules in layer decreases linearly with coverage due to adsorbent-adsorbate interactions, and that the adsorption is characterized by a uniform distribution of the bonding energies, up to some maximum binding energy (Temkin and Pyzhev, 1940). The Temkin isotherm is represented by the following equation:

\[
q_e = \frac{RT}{b_T} \ln \theta_T + \frac{RT}{b_T} \ln C_e
\]

Where, \(T\) is the absolute temperature (K), \(R\) is the universal gas constant (8.314 J/mol. K), \(A_T\) is the equilibrium binding constant (L/g), \(b_T\) is the variation of adsorption energy (J/mol).

**Dubinin–Radushkevich isotherm model**

Dubinin–Radushkevich isotherm (Dubinin and Radushkevich, 1947), is an empirical model initially conceived for the adsorption of subcritical vapors on to micropore solids following a pore filling mechanism. It is generally applied to express the adsorption mechanism (Gunay et al., 2007) with a Gaussian energy distribution onto a heterogeneous surface (Dabrowski, 2001). The Dubinin–Radushkevich isotherm is represented by the following equation:

\[
\ln q_e = \ln q_{\text{max}} - K \varepsilon^2
\]

\(K\) is Dubinin–Radushkevich isotherm constant (mol²/kJ²), \(\varepsilon\) is the Polanyi potential.

The mean adsorption energy, \(E\) (kJ/mol) is calculated with the help of following equation:

\[
E = \frac{1}{\sqrt{2R}}
\]

where \(R\), \(T\) and \(C_e\) represent the gas constant (8.314 J/mol. K), absolute temperature (K) and adsorbate equilibrium concentration (mg/L), respectively.

The obtained experimental data were fit with Langmuir, Freundlich, Temkin and Dubinin–Radushkevich models; the resulting plots are shown in Fig. 8. Table 1 summarizes the models constants and the determination coefficients. The \(R^2\) values for the Langmuir model are closer to unity than those for the other isotherm models for GCCO nanopowder (\(R^2 = 0.994\)). The Langmuir isotherm equation is therefore expected to best represent the equilibrium adsorption data. This indicates that the adsorption of BTB on GCCO nanoparticles is better described by the Langmuir model. This in turn suggests that adsorption occurs as the monolayer dyes adsorb on to the homogenous adsorbent surface. Due to the steric hindrance associated with the size of the dyes, a higher adsorption capacity can be achieved for small dye molecules (Jindarom et al., 2007) under the same experimental conditions.

| Table 1. Isotherm parameters for removal of BTB dye onto GCCO nanopowder. |
|-------------|--------|--------|
| Langmuir constants | \(q_{\text{max}}\) (mg.g⁻¹) | \(b\) (L.mg⁻¹) | \(R^2\) |
| 300 | 0.013 | 0.994 |
| Freundlich constants | \(1/n\) | \(K_F\) ((mg/g)(L/mg)^{1/n}) | \(R^2\) |
| 0.665 | 12.50 | 0.989 |
| Temkin constants | \(A_T\) (L/g) | \(b_T\) (j/mol) | \(R^2\) |
| 0.223 | 34.40 | 0.976 |
| (D-R) constants | \(q_{\text{max}}\) (mg.g⁻¹) | \(E\) (KJ.mol⁻¹) | \(R^2\) |
| 161.09 | 2.01 | 0.809 |
The favorability of the BTB dye adsorption process onto GCCO nanopowder was evaluated using a dimensionless parameter (R_L) derived from the Langmuir expression. It is defined as follows:

$$R_L = \frac{1}{1 + \frac{b}{C_0}}$$

where b is the Langmuir constant and C_0 is the highest initial dye concentration (mg/L). The adsorption process can be defined as irreversible (R_L = 0), favorable (0 < R_L < 1), linear (R_L = 1) or unfavorable (R_L > 1) in terms of R_L (Sivaraj et al., 2001). The calculated values of R_L for adsorption of BTB dye fall between 0 and 1, hence the adsorption of BTB onto GCCO adsorbent is favorable.

**Adsorption kinetics**

Several models are available to investigate the adsorption kinetics. In order to investigate the adsorption processes of BTB on GCCO nanopowder, two kinetic models were used, including pseudo-first-order and pseudo-second-order reaction rate equations are the most commonly applied models (Bayramoglu et al., 2006; Hsueh et al., 2007).

The pseudo-first-order equations expressed as follows:

$$\ln (q_e - q_t) = \ln q_e - k_1 t$$

The pseudo-first-order equation is:

$$\frac{t}{q_t} = \frac{1}{k_1q_e} + \frac{t}{q_e}$$

Where k_1 and k_2 are constants of adsorption rate, q_t is adsorption capacity at time t, q_e is adsorption capacity at equilibrium condition.

To describe the adsorption behavior and rate, the data obtained from adsorption kinetic experiments were evaluated using pseudo-first-order and pseudo-second-order reaction rate models. Plots of experimental results of dye fitted to the selected adsorption models are shown in Fig. 9. The kinetic constants along with the correlation coefficients are listed in Table 2. As shown in Table 2, higher values

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**Fig. 8.** Isotherm plots of BTB dye adsorption onto GCCO nanopowders: (a) Langmuir isotherm, (b) Freundlich isotherm, (c) Temkin isotherm and (d) Dubinin–Radushkevich isotherm (Experimental conditions: pH= 2; GCCO dosage = 0.02 g; initial dye concentration = 50, 100, 150, 200, 250, 300 mg/L and contact time = 25 min).
of $R^2$ were obtained for pseudo-second-order, indicating that the adsorption rates of BTB dye onto the GCCO nanopowder can be appropriately described using the pseudo-second-order kinetics.

**Table 2.** Kinetic parameters for the removal of BTB dye onto GCCO nanopowder.

<table>
<thead>
<tr>
<th>Pseudo-first-order constants</th>
<th>$q_e$ (mg/g)</th>
<th>$K_1$ (min$^{-1}$)</th>
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<td>0.974</td>
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<table>
<thead>
<tr>
<th>Pseudo-second-order constants</th>
<th>$q_e$ (mg/g)</th>
<th>$k_2$ (g/(mg.min))</th>
<th>$R^2$</th>
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<tbody>
<tr>
<td>47.61</td>
<td>0.028</td>
<td>0.999</td>
<td></td>
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**Fig. 9.** Plots of first and second-order rates for adsorption of BTB dye onto GCCO nanopowders: (a) pseudo-first-order rate and (b) pseudo-second-order rate.

**Conclusions**

In summary, nanoparticles $\text{Gd}_{0.5}\text{Ca}_{0.5}\text{CrO}_3$ have been fabricated by sol–gel method. The FTIR spectroscopy confirmed the structure of obtained nanoparticles. The XRD reveal that the GCCO nanoparticles prepared by calcinating the gel precursor at 700˚C for 8 h have good crystallinity in perovskite structure. The SEM images show regular morphology and particle size distribution on the surface. In the present study, we demonstrated the perovskite nanoparticles can act as novel adsorbent materials for degradation of BTB dye. The results demonstrated that the BTB dye can be successfully removed from aqueous solutions by the new GCCO nanoparticles. Isotherm modeling showed that Langmuir isotherm best-fits equilibrium data for BTB-GCCO system. The sorption kinetic can be successfully describes by a pseudo-second-order equation for the removal of BTB on GCCO nanoparticles.

**References**


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