New Synthesis of BaO nanoparticles Using the UV-Irradiation Technique for Removal Cibacron Blue Dye from their Aqueous Solution

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Abstract

This study describes a straightforward and affordable photolysis approach for the large-scale production of barium oxide nanoparticles (BaO-NPs). BaO-NPs were synthesized Using the UV-Irradiation Technique. The structural, morphological, compositional, and optical characteristics were studied by XRD, SEM, EDX and PL. the The BaO-NP's crystallite size was determined to be 14.6 nm. the morphology of BaO nanoparticles which demonstrated the existence of nanoparticles in the produced materials. and predicted bandgap of the nanoparticles was determined to be 3.26 eV by photoluminescence spectroscopy. BaO NPS that have been produced exhibit desirable adsorbent behavior for cibacron blue dye in their aqueous solutions, the effects of contact time, adsorbent mass, temperature, and starting cibacron blue dye concentration were investigated. The dye isotherm adsorption demonstrated good Freundlich agreement. The ∆H, ∆S, and AG parameters were calculated using a thermodynamic Van't Hoff equation, and the results were 9.69701 kJ/mol, 0.813 kJ/mol K, and 9.403 kJ/mol, respectively. Additionally, the kinetic investigation revealed that the adsorption exhibited pseudo-second-order behavior.

Keywords: UV-Irradiation, Photolysis, BaO NPs, photoluminescence

1. Introduction

The vast applications of nanoscience and nanotechnology, which depend on their structure and scale, give them a prominent role in scientific research¹. Nanoparticles are useful in many different fields because of their unique physical, chemical, electrical, mechanical, medical, biological, catalytic, optical, and magnetic capabilities². Precipitation, sonochemical³, solid state, sol-gel⁴, microwave⁵, hydrothermal⁶, supersaturated reverse micelles, and microemulsion method are only a few of the many techniques used to create nanoparticles⁷. Metal oxides have a special role in the evolution of nanospecies because of the diverse range of characteristics that allow them to be used in so many different contexts⁸. Semiconductor nanoparticles, in particular, have promising uses in catalysis, sensors, photoelectron, highly functional, and efficient devices⁹. Among different metal Due to its broad bandgap¹⁰, barium oxide in particular occupies a special position¹¹. The standard of water throughout Rapid expansion of the population. A nation's development depends on its industrial expansion¹². As various sectors reject life-threatening sewage entering aquatic systems, which is not good influences the water's quality¹³. The fabric, printing, papermaking, cosmetics, food processing, and the pharmaceutical industry releases its effluents into the water bodies with high levels of colors and dyes¹⁴. It has been discovered to be neurotoxic, chronically toxic, and carcinogenic to both people and animals. Dye treatment from industrial effluents is necessary before they pose a threat to the environment¹⁵. In addition to harming the beauty of the river, dyes also hinder sunlight's ability to pass through water, which has an impact on how aquatic plants synthesize food. In the present study, BaO NPS adsorption of cibacron blue dye. BaO NPS prepared by photolysis methods and were characterized through SEM, EDX, XRD, PL. and study its kinetic with thermodynamic properties.
2. Experimental
2.1. Materials
Barium acetate (CH₃COO)₂Ba, urea (NH₂)₂CO, sodium hydroxide (NaOH) and Cibacron blue dye with deionized water were used throughout the preparation period.

2.2. Methods
All chemicals used in this study were purchased from sigma Aldrich without any purification. BaO nanoparticles were synthesized by the photolysis method. As shown in fig. (1), a photocell was used to irradiate the source of BaO NPS. Immersed U.V source (125 w) is used with λ=365nm. The cell contains a quartz tube like jacket for immersion U.V source in the solution barium. Pyrex counting was used as a reactor. An ice bath cools the reactor to avoid rising temperature due to U.V lamp. Accordingly, 100 ml of 0.4 mole urea was added slowly (drop per second) to 100 ml of 0.2 mole Barium acetate under a magnetic stirrer for 15 min at 25 °C then, a photocell irradiated the Barium solution for 30 min. The white precipitate is formed it is separated and washed several time with D.W the powder was dried at 120 °C for 3 h and calcinated at 600 °C for 3 h. A white powder of BaO NPs is obtained.

2.3. Adsorption cibacron blue on BaO NPS
Using 100 mg/L Cibacron Blue dye stock solutions made with distilled water, batch adsorption tests were carried out. The Cibacron Blue dye was added to the solution and extra diluted to the proper quantities (5–25 mg/L). In 200 ml glass tubes, 100 ml of dye solution and 0.01 g of BaO NPS were combined to complete the tests. The glass tubes were set on a shaker (HZQ-C) at different temperatures (15, 25, 35, 45, and 55 °C) and shook for 90 minutes. A UV-visible spectrophotometer was used to calculate the dye concentration. Equation 1 was used to quantify the amount of dye that had been adsorbed in the solution. Qₑ indicates the equilibrium capacity of adsorption (mg/g), and C₀ is the initial concentration of the dye Cibacron Blue (mg/L).

\[
Q_e = (C_0 - C_e) \frac{V_{sol}}{m}
\]

where Qₑ represents the equilibrium capacity of adsorption (mg / g), C₀ is the initial Cibacron Blue dye concentration (mg / L), Cₑ is the equilibrium concentration of Cibacron Blue dye (mg / L), V is the volume of the solution (L), and m is the amount of the BaO NPS.

3. Results and discussion
3.1. Characterization
BaO-NPs as-synthesised were examined for phase and crystallinity using the XRD method and Cu Ka radiations (λ = 1.5406). Figure 2: As-produced BaO nanoparticles' XRD patterns in the 10–80° range. Fig. (2) displays the XRD spectra of BaO-NPs that were created. The observed XRD pattern contains numerous well-defined peaks that are connected to and well-matched with the tetragonal phase of BaO-NPs and are in good agreement with the "JCPDS" card No. 26-0178. These peaks include (200), (223), (211), (102), (375) and (212), among others. The produced nanoparticles are highly crystalline in character, as indicated by sharp and strong peaks. The Debye–Scherrer formula, which can be written as

\[
D = \frac{K \lambda}{\beta \cos \theta}
\]

is used to calculate the sample's average crystallite size (D), where β is the full-width at half maximum, K is the form factor (0.90), and is the wavelength of Cu Ka radiation (λ=1.5406). The diffraction angle is . BaO-NPs are discovered to have an average grain size of about (14.6nm).

Fig. (3) show the morphology of the synthesized BaO-NPs as determined by FE-SEM. A low-resolution image of prepared BaO-NPs with a flower-shaped cluster morphology is shown in a high-resolution SEM picture, demonstrates the extremely crystalline growth of the produced nanomaterials. The SEM pictures also show that some NPs are arranged in structures resembling nanorods and nanosheets and are connected by accumulating on top of one another. By integrating EDS and SEM, the elemental composition analysis of produced NPs was evaluated. The chemical purity of the produced BaO-NPs was validated by EDS spectra. Where an analysis (EDX) as shown in fig.(4) shows that the nano-barium oxide is of high purity. When an optically excited substance emits light spontaneously, it is called photoluminescence (PL). In order to investigate various excitation types, determine the bandgap energy, find specific defects for radiative transitions, and determine the impurity levels in the nanomaterial sample, the excitation energy and intensity can be selected. The analysis method of photoluminescence spectroscopy is non-destructive fig. (5) depicts the photoluminescence spectra of BaO-NPs as it was measured at an excitation wavelength of 230 nm. The spectrum displays two significant emission bands (Blue-violet and red), distinct peak are identified: a strong emission peak at 380nm.

The energy gap can be calculated according to the following equation:

For example \( E_v = \frac{1240}{\lambda} \ldots \) \[3\]

Where \( E_v \) is the energy gap, and \( \lambda \) is the maximum wavelength.

And by the wavelength and according to the above equation, the energy gap of barium oxide is \( 3.26 \text{ ev} \).

In previous studies, Barium nitrate \( \text{Ba(NO}_3\text{)}_2 \) and sodium bicarbonate \( \text{NaHCO}_3 \) were used by Manauwar Ali Ansari et al. to prepare barium oxide nanoparticles using the co-precipitation method. The resultant of barium oxide nanoparticles from this method had a crystal size of 16–15 nm and energy gap 4.65 \( \text{ev} \).

In another study, The author Mohamed E. and his group prepared nano barium oxide (BaO NPS) using \( \text{BaCl}_2.2\text{H}_2\text{O} \), \( \text{NaOH} \) and \( \text{(NH}_2\text{)}_2\text{CO} \) with continuous stirring at a temperature of 50 and burning the precipitate at a temperature of 500°C, the NPs Size was to be 50 \( \text{nm} \).

The author E.Sundharam and his group prepared nano barium oxide using ultrasonication method using \( \text{BaCl}_2.2\text{H}_2\text{O} \) and \( \text{NaOH} \) temperature 400°C (4hr) Crystal size nanoparticles were 19.6 \( \text{nm} \).

As the author Arvind k.Bhakta and his group prepared the nano-barium oxide using infrared irradiation and diazonium chemistry, and the crystal size was (3-18\( \text{nm} \)).

In our method, The photolysis approach differs from other methods in that it is straightforward, quick, and inexpensive. In addition, it produces a lot of nanoparticles that are highly pure, have small crystal sizes, and are produced in vast amounts, as demonstrated by the current study.

3.2. Adsorption isotherms

To determine how the BaO NPS and cibacronblu interact, fit isotherm adsorption with the adsorption results. This project by Freundlich The correlation value \( R^2 \) demonstrates that Freundlich isotherm equations suit the adsorption findings as displayed in fig. (6) quite closely. The following equation represents the Freundlich adsorption's linearized form26

\[
\log(Q_e) = \log(K_f) + \frac{1}{n} \cdot \log(C_e)
\]

Where \( K_f \) and \( n \), which are called the Freundlich constants, are the indicators of adsorption capacity and adsorption intensity, respectively27. The \( K_f \) is gotten from the intercept and the \( n \) is obtained from the slope computed 1/n,
of cibacron blue (25 ppm) in a sequence of tests. The mixture was shaken at 200 rpm under 298 K temperature. Fig. (7) shows the adsorption was extremely rapid at the beginning of 15–90 min. The fast adsorption came from a strong link between active BaO NPS and cibacron blue dye. Because of the occupancy on the BaO NPS surface, the adsorption rate of cibacron blue dye became constant after 90 min.

Fig. 7 | Effect of time on adsorption of Siba Crohn F3GA blue dye at 25°C by BaO NPS

4-4 | Effect of adsorbent mass

After adding various amounts of BaO NPS, ranging from 0.01 to 0.2 g to 25 ppm of cibacron blue dye, the influence of the adsorbent mass was calculated. The concoction was 200 rpm shaken at 298 K temperature. The quantity of Fig. (8) displays adsorbed as a function of mass. First of all, the higher amount of the adsorption rate is due to active BaO NPS location. Increased cibacron blue dye absorption was demonstrated by boosting the adsorbent mass.

Fig. 8 | The mass effect of BaO NPS on dye removal percentage

4-5 | Temperature effect and thermodynamic parameter calculation

At specific temperatures of 15°C, 25°C, 35°C, 45°C, and 55°C, the effect of temperature on cibacron blue dye adsorption on the surface of BaO NPS was investigated. With rising temperature, more cibacron blue dye adsorption solution was produced. This indicates that the process was endothermic and that ΔH had a positive mean. Both the absorption and adsorption processes were demonstrated by this. The strong link that binds the adsorbate and the adsorbent was strengthened as a result of diffusion molecules being absorbed into the pores as the temperature climbed. The thermodynamic parameters (ΔH, ΔS, ΔG) give precise information on the associated adsorption-related intrinsic energy modifications, thus they must be evaluated properly. The following modifications were estimated in this study to predict the adsorption process using free energy ΔG:

\[ \Delta G = \Delta H - T \Delta S \]

where \( K_e \) is an equilibrium constant, \( R \) is a gas constant (8.314 J/mol K) and \( T \) is the temperature (K). From the slope, and interception of the plots of van’t Hoff, ΔH and ΔS were identified as shown in Fig (10).

The corresponding ΔH from the slope was 9.697 kJ/mol, which indicated the process was endothermic. The ΔS from the intercept was 0.813 kJ/(mol K) that indicated the adsorbed molecules were still in constant motion on the surface, and they were attributable to absorption as well as adsorption. A higher temperature facilitated the adsorption of cibacron blue dye. The ΔG for adsorbing was determined to be 9.4303 kJ/mol at 298 K, which means spontaneous adsorption.

Fig. 10 | (Van’t Hoff) Scheme of adsorption of Siba Crohn blue dye by nanobarium oxide (BaO NPS) at different temperatures.

6 | Dynamics

To study adsorption, it is important to know the process kinetics to know the rate of pollutant removal from aqueous solutions. In order to determine the most suitable model for the adsorption of Cibacron F3GA blue on the surface of (NPS BaO), the evaluation was carried out using two orders, namely the false and the second order.

The adsorption of Cibacron F3GA blue dye on the surface of (BaO NPS) was applied to the pseudo-first-order by the equation:

\[ \ln(q_e - q_t) = \ln q_e - K_1 t \]

where \( q_t \): the amount of dye adsorbed at each time (t min), \( q_e \): the amount of dye adsorbed at equilibrium (mg/g), \( K_1 \): first-order constant (min\(^{-1}\)). Plotting the relationship between (lnqe-qt) versus time (t), which gives a linear relationship. as in Fig. (11).
The results were fitted well with Freundlich isotherm as in fig. (12).

$$\frac{t}{q_t} = \frac{1}{K_2 q_e^2} + \frac{1}{q_e} t$$  \[\text{(9)}\]

where \(R^2\) is (0.998), K1 (Slope) is (-0.043352) in g/mg units, K2 is 0.196996 in g/mg.min.

Through the results, it is clear that the value of \(R^2\) for the second order of dye adsorption by the BaO NPS is greater than R1 for the first order, and this indicates that the adsorption proceeds according to the pseudo second order.

7 Conclusion

quality of the rutile BaO NPS according to the XRD, SEM,PL characterizations. All the characterization techniques gave identical results to prove that the synthesized nano-BaO their average crystal size was calculated to be 14.6 nm. The nanoparticles proved to have excellent adsorption properties for removing cibacron blue dye from aqueous solutions. Both kinetic and dynamic studies were shown the efficiency of the BaO NPS adsorption. The results were fitted well with Freundlich isotherm model. The calculations of the thermodynamic parameters gave the adsorption values for \(\Delta H\), \(\Delta S\), and \(\Delta G\) equal to 9.69 kJ/mol, 0.813 KJ/mol K, and 9.4303 kJ/mol, respectively. These results prove that the adsorption is an endothermic spontaneous process.

8. References


