

## Research Article

# Analysis of Spectroscopic, Morphological Characterization and Interaction of Dye Molecules for the Surface Modification of TiB<sub>2</sub> Nanoparticles

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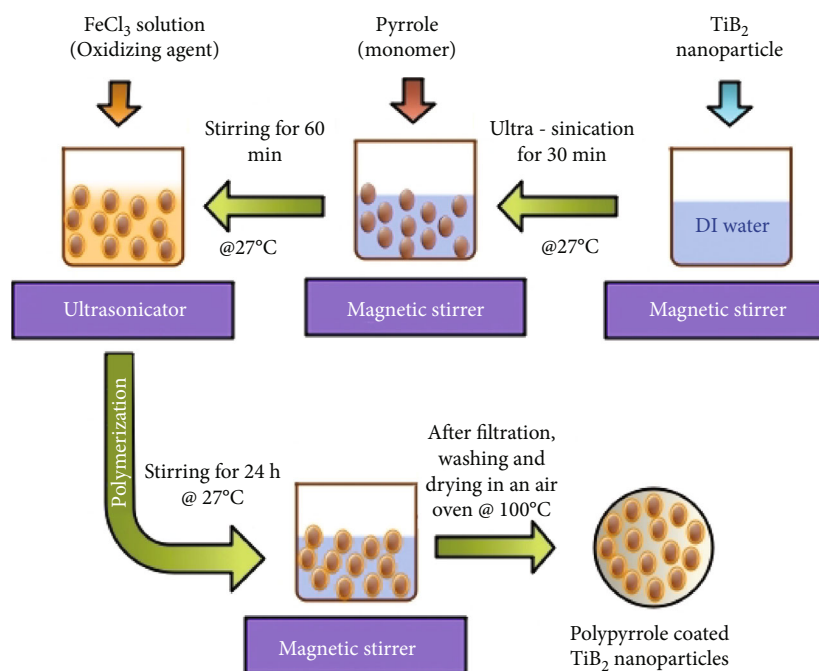
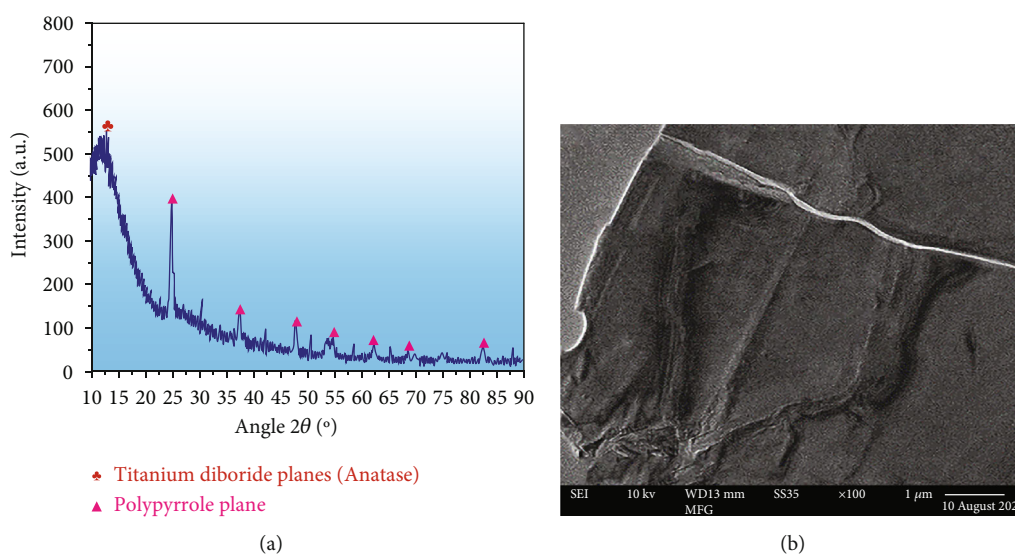
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Nanoparticles of titanium diboride (TiB<sub>2</sub>) coated with a conductive polymer and subjected to an oxidizing agent was carried out in this research. TiB<sub>2</sub> nanoparticles coated with polypyrrole (PPy) were studied using XRD and TEM techniques. Nitrogen adsorption and desorption properties of nanoparticles are covered with modified polypyrrole to understand better the surface zone, structural features, and pore geometries of the nanoparticles. The water-based hazardous anionic Congo red (CR) dye was removed using polypyrrole-coated titanium diboride (PPy@TiB<sub>2</sub>) nanoparticles. Numerous cutting-edge experimental techniques, including FTIR, FE-SEM, EDXS, and element mapping analysis, were used to confirm the CR color's adhesion to the PPy@TiB<sub>2</sub> nanoadsorbent study. While conducting batch experiments, coated TiB<sub>2</sub> nanoadsorbents with polypyrrole enhanced the adsorption behaviour. One of the factors evaluated in the adsorption tests was pH; another was contact time, and a third was dose. At pH = 4, 98.75% of Congo red dye was detached using 60 mg of PPy-coated TiB<sub>2</sub> nanoadsorbent. Several sorption-desorption cycles were performed on this nanoadsorbent to determine its reusability. An excellent adsorption capacity for water treatment is reported in PPy-coated TiB<sub>2</sub> nanoadsorbent.

## 1. Introduction

Every continent has a textile industry, and this sector contributes significantly to global economic growth [1]. Due to

the coloring, printing, and washing processes carried without the appropriate pretreatment measures, textile wastewater is increasingly dyed. There are various concentrations of colored components in textile wastewater, which contribute

FIGURE 1: Diagram for the production of PPy@TiB<sub>2</sub>.FIGURE 2: (a) XRD and (b) TEM analysis of PPy@TiB<sub>2</sub>.

to its pollution [2]. Therefore, textile effluent is clean since it has a high concentration of pigment molecules. Due to CR's ability to convert to benzidine, the anionic synthetic diazo dye is carcinogenic. Because of its aromatic structure, CR dye is both physiochemically and thermally stable [3]. CR's color changes depending on the pH, turning red at a pH of less than 5. Toxic to humans and animals, industrial effluent contains significant levels of COD and BOD, suspended particles and colloids, salts, and some other hard materials [4]. To deal with textile wastewater treatment issues, energy, water, and chemicals are required. There are numerous advantages to adsorption over other water treatment methods that are investigated as a viable option [5].

Various adsorbent forms are tried, including polar and nonpolar adsorbents. High surface-area adsorbent materials have effectively extracted different dye compounds from textile effluent. Various essential uses for nanomaterials have also been demonstrated [6–8]. Nanoparticles of titanium diboride (TiB<sub>2</sub>) were employed in multiple material science applications: high-performance solar cells and photocatalytic devices using TiB<sub>2</sub>-based nanotubes or nanoflowers, TiB<sub>2</sub> nanoparticles, and a PVDF membrane covered with a TiB<sub>2</sub> layer amplified for photocatalytic wastewater purification [9–11]. A polymer matrix alters the material's properties, such as combining polypropylene, polypyrrole, polyamide, or polylactic acid. Environmental scientists have

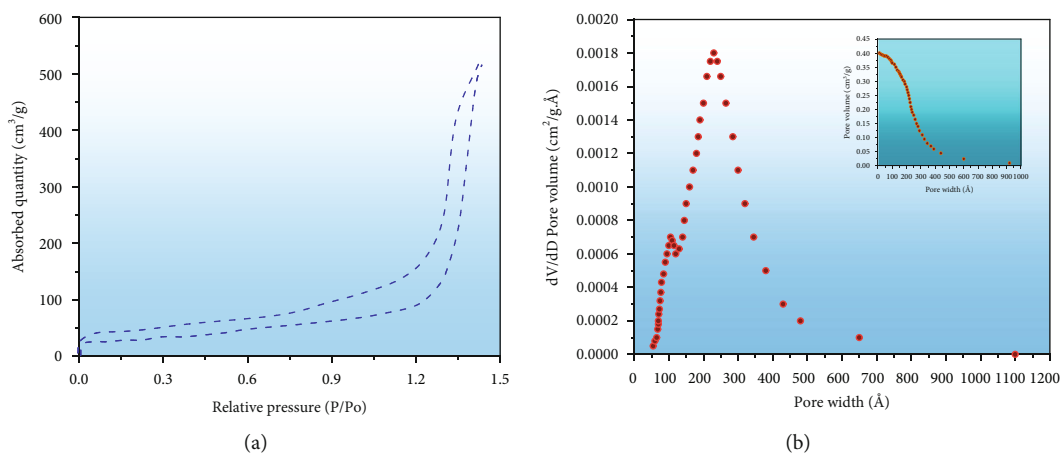


FIGURE 3: (a) Isotherm of N-adsorbed and desorbed of PPy@TiB<sub>2</sub> and (b) grain size dispersal plot and comparison of pore width and volume plot.

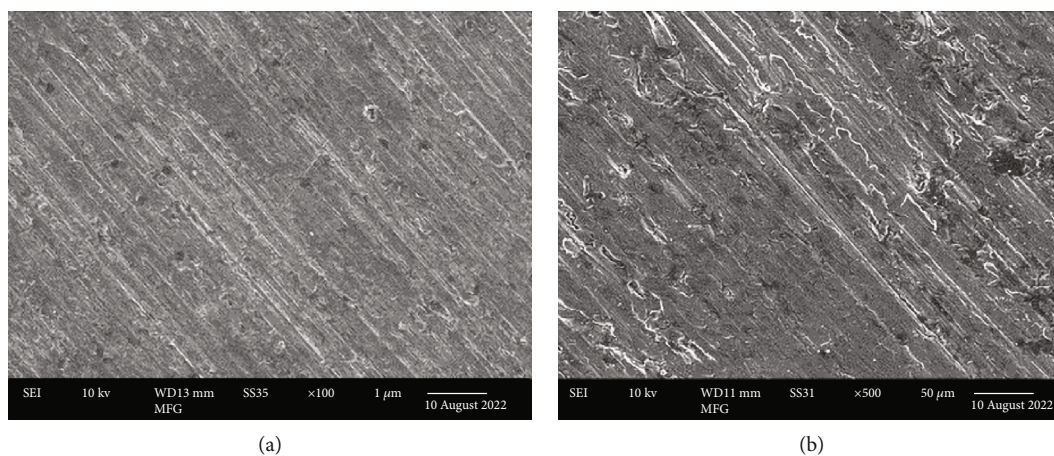


FIGURE 4: FESEM images of various magnifications of (a) PPy@TiB<sub>2</sub> and (b) CR-adsorbed PPy@TiB<sub>2</sub>.

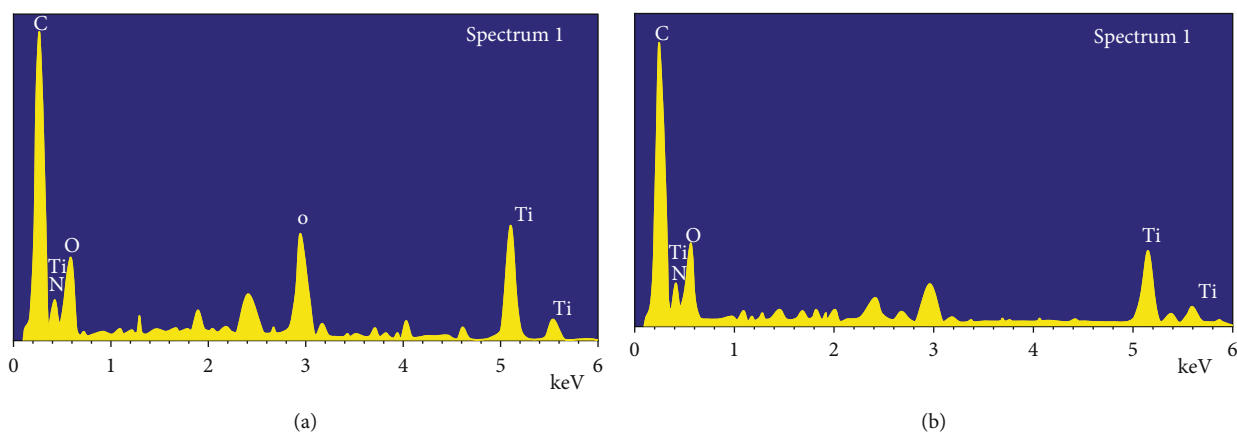


FIGURE 5: Energy dispersive X-ray analysis of (a) PPy@TiB<sub>2</sub> and (b) Congo red-adsorbed PPy@TiB<sub>2</sub>.

been considering polymeric nanoadsorbents based on polypyrrole for several years.

The polymer has many properties that make it an ideal solution for water pollution problems [12]. Scientists have found that polypyrrole-based nanocomposites effectively

clean the environment of heavy metal ions and radioactive pollutants, pigments, chemical compounds, and pesticides. Recent studies highlight photovoltaic efficiency, optical features, degradation capacity of photocatalytic materials, physical characteristics, and electric characteristics [13–15]. The

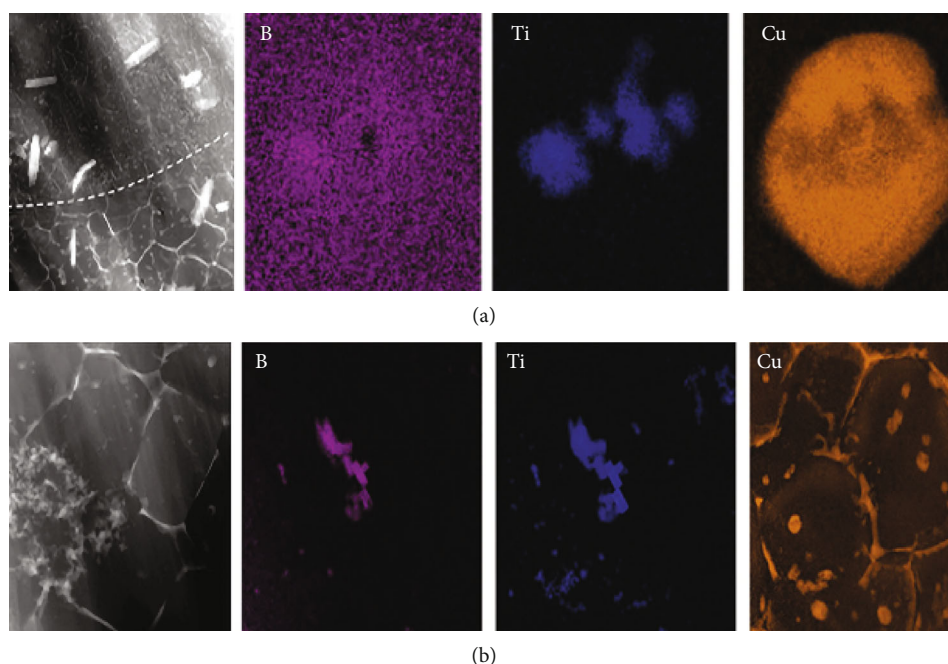


FIGURE 6: (a) PPy@TiB<sub>2</sub> and (b) process after Congo red-adsorbed PPy@TiB<sub>2</sub> elemental mapping analysis.

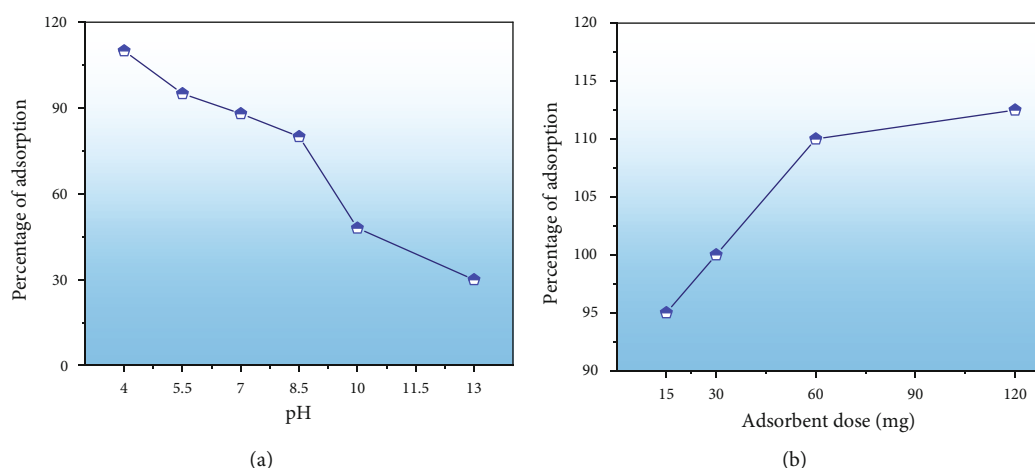


FIGURE 7: (a) Impact of pH and (b) impact of adsorbent dosage.

chemical polymerization production of polypyrrole-coated titanium diboride (PPy@TiB<sub>2</sub>) has positively affected the environment. In the adsorption investigation, several traits and parameters were analyzed. Scientists have used the produced PPy@TiB<sub>2</sub> nanomaterial for wastewater dye removal due to its excellent performance and ease of reusing [16].

## 2. Experimentation

**2.1. Methods and Materials.** In the presence of a PPy monomer, a chemical polymerization approach is used to create a PPy@TiB<sub>2</sub> nanoadsorbent. There were 2 mL of pyrrole monomer and 30 minutes of ultrasonication at 27°C to prepare TiB<sub>2</sub> nanoparticles. The pyrrole monomer was allowed to adsorb on the TiB<sub>2</sub> nanoparticles' surfaces by swirling the dispersion for 60 minutes at 27°C [17, 18]. Iron was added to

the TiB<sub>2</sub> dispersion as an oxidizing agent to polymerize the pyrrole on the nanoparticles' surfaces. The mixture was stirred for an additional 24 hours at 27°C using an electric stirrer. A four-hour oven drying at 100°C removed the contaminants and unreacted oxidants after cleaning the PPy@TiB<sub>2</sub> nanoadsorbent with deionized water and acetone after polymerization. An illustration of TiB<sub>2</sub> nanoparticles coated with polypyrrole is indicated in Figure 1.

**2.2. Characterization.** PPy@TiB<sub>2</sub> nanoadsorbent including TEM, HR-TEM, and fast Fourier transform (FFT) were synthesized. TESCAN FE-SEM, elemental mapping, and EDX analyze the PPy@TiB<sub>2</sub> nanoadsorbent earlier and the subsequent adsorption. X-ray diffraction measurements were determined using Cu K $\alpha$  radiation with a 10° to 90° (2 $\theta$ ) wavelength range [19]. The nitrogen adsorption technique

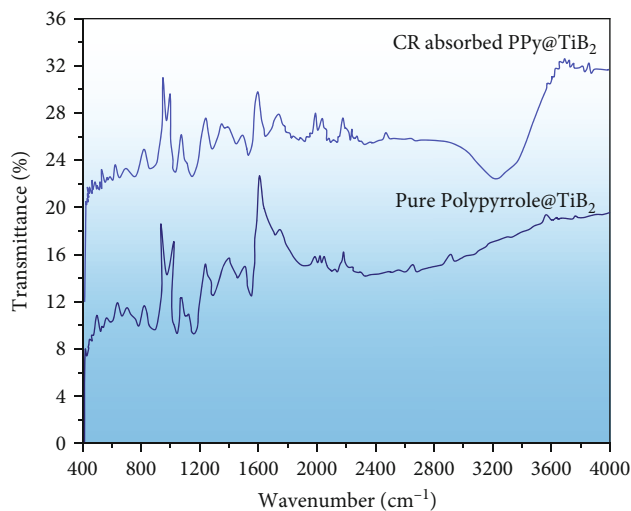


FIGURE 8: Image of attenuated total reflection-Fourier transform infrared spectroscopy.

measures pore size distributions and particular surface area. Using an FTIR spectrometer, it was necessary to conduct pre- and postadsorption studies on the nanoadsorbent, nano-PPy@TiB<sub>2</sub> [20].

**2.3. Adsorption Studies.** When removing Congo red (CR), batch experiments were carried out to test how the as-synthesized PPy@TiB<sub>2</sub> nanoadsorbent performed. The nanoadsorbent PPy@TiB<sub>2</sub> nanoadsorbent was studied to assess the effects of pH (acidity), contact time, and dosage on adsorption performance [21]. A pH experiment was conducted to check the PPy@TiB<sub>2</sub> nanoadsorbent-adsorbed CR at various pH levels (4 to 13). Adsorbent doses of 60 mg and 10 ppm of CR were used to test the adsorption performance. The synthetic nanoadsorbent was evaluated at various doses (15, 30, 60, and 120 mg) in a 10 ppm CR dye concentration to determine the effect of the dose [22]. Color concentrations were measured using a UV-V spectrophotometer during the batch adsorption trials. The formula used to calculate the percentage of CR dye that is eradicated is as follows [23, 24]:

$$\text{Percentage of Removal} = \frac{C_i - C_f}{C_i} \times 100 \quad (1)$$

$C_i$  starting concentration of Congo red color  
 $C_f$  CR dye's ending concentration

### 3. Results and Discussions

**3.1. Properties of PPy@TiB<sub>2</sub> Nanoparticles.** Investigation of PPy@TiB<sub>2</sub> nanocrystalline adsorbent structure was done using XRD. As-synthesized PPy@TiB<sub>2</sub> is indicated in Figure 2(a). Results showed that TiB<sub>2</sub> nanoparticles in composite form with PPy polymer were anatase crystalline. There were no visible phases other than TiB<sub>2</sub> in Figure 2(a), which verify the XRD pattern of its different stages (rutile and brookite). It discovered that PPy@TiB<sub>2</sub> nanocomposite reduced the crystalline phase of TiB<sub>2</sub> and

PPy's amorphous nature by interacting with TiB<sub>2</sub>. XRD data suggest that the surface morphology of TiB<sub>2</sub> nanoparticles with conducting polymer PPy was successful, as demonstrated by these results [25, 26]. Furthermore, TEM and HR-TEM analysis strongly support these findings.

Yet, Figure 2(b) shows the TEM patterns of the synthesized PPy@TiB<sub>2</sub>. The TEM image of the synthesized PPy@TiB<sub>2</sub> showed that only the TiB<sub>2</sub> facet had PPy polymer deposited on it, regardless of the other aspect [27]. A patch of unilluminated PPy polymer showed TiB<sub>2</sub> nanoparticles scattered everywhere, while PPy polymer granules with TiB<sub>2</sub> nanoparticles interacted tightly. TiB<sub>2</sub>'s high contrast made it distinct from PPy, which has lower contrast (Figure 2(b)). This nanoadsorbent formed using TiB<sub>2</sub> in the TEM image inset clearly showed the crystalline structure of the TiB<sub>2</sub> [28]. High resolution-TEM and FFT confirm that TiB<sub>2</sub> nanocomposite PPy@TiB<sub>2</sub> nanocomposite had an anatase crystalline phase as shown in supplemental Figures 3(a) and 3(b).

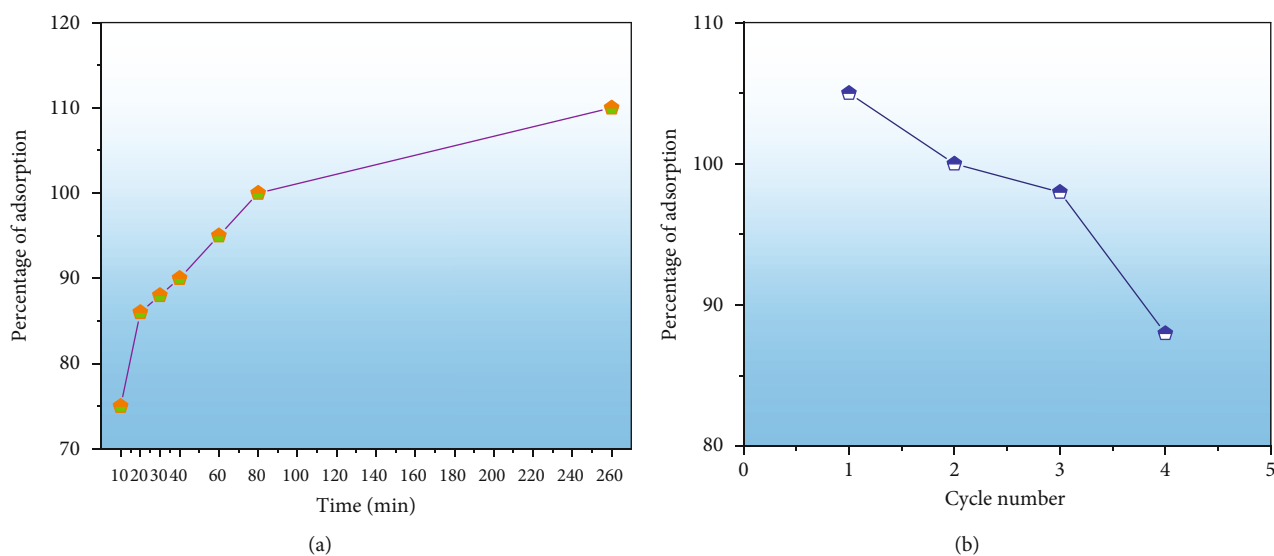
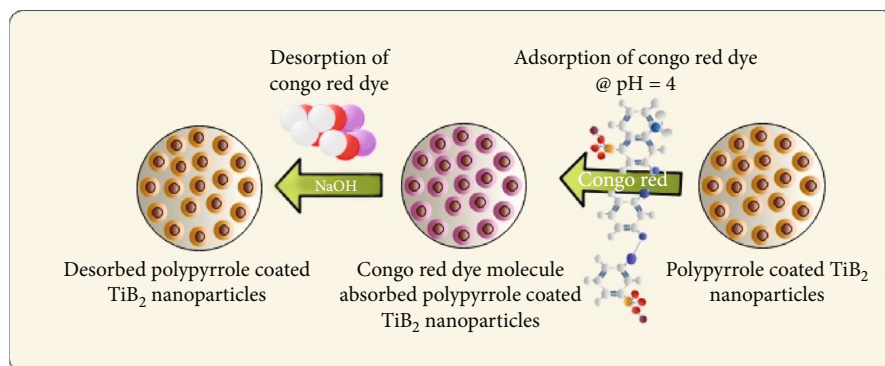
The surface, structural characteristics, and pore geometries of the PPy@TiB<sub>2</sub> nanospecific adsorbent were determined by the isotherm of nitrogen attachment shown in Figure 3(a). It reveals that saturation is observed at higher pressures, with a gradual increase in adsorption up to roughly 0.70 relative pressure (P/Po) [29]. Because of this, mesoporous adsorbents have a unique isotherm. The average pore diameters of the nanoadsorbent with 62.27 m<sup>2</sup>/g surface area are 245.307 Å for adsorption and 216.698 Å for desorption using the nitrogen adsorption-desorption isotherm and the BJH technique [30]. In Figure 3(b), the average BET pore sizes for adsorption and desorption were 232.0041 Å and 247.2938 Å, respectively. According to IUPAC guidelines, mesoporous materials have pore diameters between 20 and 500 nm. Thus, the PPy@TiB<sub>2</sub> nanoadsorbent favours the mesoporous character of the material because of its pore sizes. Desorption is showed to be possible in Figure 3(a), where the desorption sum pore volume (0.385022 cm<sup>3</sup>/gram) is greater than the adsorption sum pore volume (0.361217 cm<sup>3</sup>/gram) at P/Po = 0.982. It shows that reversible desorption is conceivable. The regeneration results confirmed this tendency even more conclusively. A mesopore capillary condensation is present in the isotherm, in accordance with IUPAC's classifications of physisorption isotherm/hysteresis loops [31].

These images were taken at three different magnification levels and show the material's granular structure. As the magnification increased, the brightness decreased, indicating that the brightest particles were on the surface while the duldest particles were discovered deep within the layer (Figures 4(a) and 4(b)). It appeared like the character was uniform and smooth. Nanoparticles prepared before CR adsorption became more dispersed and bulkier in their microstructures.

Additionally, the components in the adsorbed PPy@TiB<sub>2</sub> were found out by FESEM-EDX analysis. As depicted in Figure 5, pure and CR-adsorbed PPy@TiB<sub>2</sub> samples have different elemental compositions. In the EDX spectra, carbon, oxygen, nitrogen, and titanium (Ti) are all visible (Figure 5(a)). Because of their relative atomic weights of

TABLE 1: Assignment of FTIR peaks in the pure and Congo red-adsorbed PPy@TiB<sub>2</sub>.

FTIR peaks of PPy@TiB <sub>2</sub> (wavenumber cm <sup>-1</sup> )	Peaks role	FTIR peaks of Congo red-adsorbed PPy@TiB <sub>2</sub> (wavenumber cm <sup>-1</sup> )	Peaks role
1537.64	Carbon-carbon pyrrole bond stretching vibration	1521.56	Carbon-carbon pyrrole bond stretching vibration
1447.79	Conjugated C-N band for stretching	1436.68	Conjugated C-N band for stretching
1291.17	A band of conjugated deformation in the plane of C-H	1272.28	A band of conjugated deformation in the plane of C-H
1142.18	Conjugated carbon-nitrogen stretching vibrations	1136.32	Conjugated carbon-nitrogen stretching vibrations
1034.13	Nitrogen-hydrogen in the planar deformation band	1025.04	Nitrogen-hydrogen in the planar deformation band
992.25, 882.29, 780.08	Deformations of the pyrrole ring G-H in and out of the plane	988.32, 860.58, 754.42	Deformations of the pyrrole ring G-H in and out of the plane
664.89	Characteristic peak of TiB <sub>2</sub>	649.33	Characteristic peak of TiB <sub>2</sub>
478.71	Stretching mode for the titanium-boride bond	477.51	Stretching mode for the titanium-boride bond

FIGURE 9: (a) Impact of contact duration and (b) regeneration capabilities of PPy@TiB<sub>2</sub> for Congo red dye adsorption.FIGURE 10: Diagram for adsorbed and desorbed mechanism of Congo red dye molecule on PPy@TiB<sub>2</sub>.

50.75 and 28.21%, C and O atoms may have an effect on PPy@TiB<sub>2</sub>'s adsorption to CR dye molecules. In Figure 5(b), Na, C, and O atoms with high atomic masses indicate as evidence that CR molecules adsorb onto PPy@TiB<sub>2</sub>. Elements in PPy@TiB<sub>2</sub> and CR-adsorbed PPy@TiB<sub>2</sub> were studied further by an elemental mapping experiment [32, 33]. Before and after CR dye adsorption on PPy@TiB<sub>2</sub>, the elemental mapping is indicated in Figure 6. In the PPy@TiB<sub>2</sub> nanoadsorbent, carbon, oxygen, nitrogen, and titanium were all evenly distributed. C, O, and Na distributions were validated by exposing the nanoadsorbent PPy@TiB<sub>2</sub> nanoadsorbent to CR molecules [32, 33].

### 3.2. Behaviour of Adsorption

**3.2.1. Impact of pH.** When the aqueous phase's pH alters, it alters both the binding sites and ionization processes in this experiment [34, 35]. Figure 7(a) illustrates the percentage of Congo red adsorption by PPy@TiB<sub>2</sub> at different pH values (4.0, 5.5, 7, 8.5, 10.0, and 13.0). The starting pH level of 4.0 (98.75%) had the highest adsorption rate, which decreased as the pH increased. Positively charged PPy@TiB<sub>2</sub> surface was inhibited by electrostatic contact with negatively charged CR (due to the presence of two -SO<sub>3</sub> groups) [36]. Both Congo red and PPy@TiB<sub>2</sub> became negatively charged as the pH increased and the adsorption percentage reduced [37].

**3.2.2. Effect of Adsorbent Dose.** The amount of PPy@TiB<sub>2</sub> used had a beneficial impact on the adsorption percentage (Figure 7(b)). The percentage of adsorption rate was increased to obtain the equilibrium state of maximum adsorption [38]. Adsorption on the PPy@TiB<sub>2</sub> surface increases when the dose increases as the number of adsorption sites increases. The optimum PPy@TiB<sub>2</sub> amount attains to be 0.05 gm due to this study.

The AT-FTIR spectrum of pure PPy@TiB<sub>2</sub> and that of CR-adsorbed PPy@TiB<sub>2</sub> is indicated in Figure 8. Band assignments and wavenumbers are shown in Table 1 for the two samples, including the pure TiB<sub>2</sub> and TiB<sub>2</sub>+CR-adsorbed PPy@TiB<sub>2</sub> FTIR bands. Most peaks experienced a minor shift to the lower wavenumbers after CR dye adsorbs on PPy@TiB<sub>2</sub> [39]. Therefore, these shifting peak values make it clear that the CR dye was successfully adsorbing onto PPy@TiB<sub>2</sub>.

**3.2.3. Effect of Adsorption Time.** In adsorption experiments, adsorption time is critical. CR adsorption is illustrated in various time intervals in Figure 9(a). During the first 30 minutes of the investigation, the percentage of CR slowly adsorbing into the PPy@TiB<sub>2</sub> grew dramatically, reaching a maximum (110%) after 260 minutes. CR adsorption was stable, indicating that it attains the equilibrium phase. For the nanoadsorbent PPy@TiB<sub>2</sub> nanoadsorbent, 260 minutes show the optimal duration for carbon dioxide adsorption. When PPy@TiB<sub>2</sub> is adsorbed, the diffusion process is evidenced by the gradual rise in the adsorbed percentage [40].

**3.2.4. Regeneration Study.** The exhausted adsorbent (Congo red-adsorbed PPy@TiB<sub>2</sub>) was evaluated on how well it can

revitalize in a regeneration study [41]. It is shown in Figure 9(b) that NaOH was used as a desorbing agent for four cycles. This PPy@TiB<sub>2</sub> nevertheless showed outstanding regeneration efficacy with an adsorption decrease from 98.75 to 97% in Figure 9(b). PPy@TiB<sub>2</sub> proves to be a valuable and economical material for removing CR. The reproducibility of PPy@TiB<sub>2</sub> for eliminating CR from textile wastewater was established.

**3.2.5. Adsorption Mechanism.** The dynamic, active sites on the PPy@TiB<sub>2</sub> surface allowed CR to adsorb on it. A schematic diagram of CR dye molecule adsorption and desorption onto PPy@TiB<sub>2</sub> nanoadsorbent is indicated in Figure 10. Chemical polymerization was employed to synthesize nanocomposite using TiB<sub>2</sub> nanoparticles. Dispersed TiB<sub>2</sub> nanoparticles were used to adsorb the Polypyrrole monomer. Then, the pyrrole-adsorbed TiB<sub>2</sub> dispersal is treated with the FeCl<sub>3</sub> solution as an oxidizing agent. Finally, CR adsorption was determined using the polymerized PPy@TiB<sub>2</sub> nanoadsorbent. Carbo-oxygen-nitrogen-titanium elements make up its surface. Adsorption of textile wastewater is made more accessible with the final polymer form's increased surface area and chemical functionality. This positive charge is created by protonating the PPy@TiB<sub>2</sub> surface functional groups at pH = 4. This positive charge is used to interact with CR's negative -SO<sub>3</sub> groups. Due to the negative CR's electrostatic attraction to the TiB<sub>2</sub> surface, the adsorption is reduced as pH rises, whereas deprotonation caused a decrease in surface charge density.

## 4. Conclusion

Chemical polymerization with pyrrole (monomer) and ferric chloride successfully converted tiny titanium diboride nanoparticles (oxidizing agent). Polypyrrole originates on the surface of TiB<sub>2</sub> nanocomposites. PPy@TiB<sub>2</sub> nanoparticles were formed, and CR molecules could adhere to them, according to spectroscopic, structural, and morphological studies. It discovered that pH = 4, 260 minutes of contact hours, 60 mg of adsorbent, and a Congo red dye concentration of 10 ppm were the ideal adsorption conditions. After four cycles, a nanoadsorbent could maintain 90% of the CR color. The outcomes show that polypyrrole-coated titanium oxide nanoparticles are an effective treatment option for colored wastewater containing dye.

### Data Availability

All authors confirmed that all necessary data are available in the manuscript.

### Conflicts of Interest

The authors declare that they have no conflicts of interest.

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