Amine functionalized 3D porous organic polymer as an effective adsorbent for removing organic dyes and solvents

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Organic dyes and solvents are emerging pollutants; the development of new materials for their efficient adsorption and removal is thus of great significance. In this work, we report the application of an amine functionalized triptycene-based 3D polymer (TPP-NH₂) as a novel adsorbent for the fast removal of organic dyes in aqueous solution and organic solvents with a view to understanding its adsorption kinetics, adsorption isotherms, desorption and adsorbent regeneration. The adsorption of organic dyes (cationic methylene blue, MEB, and anionic methyl orange, MO) on TPP-NH₂ was fast, and most of the dyes were adsorbed in 60 min. The adsorption of MEB and MO follows pseudo-second-order kinetics and fits the Langmuir model, where the maximum adsorption capacity increased from 204.9 mg g⁻¹ to 560.6 mg g⁻¹ and 213.2 mg g⁻¹ to 803.2 mg g⁻¹ for MEB and MO, respectively, as the temperature increased, suggesting that the adsorption of MEB and MO on TPP-NH₂ is favorable at high temperatures via an endothermic process. The pH has no obvious effect on the adsorption of MEB and MO. The used TPP-NH₂ could be regenerated effectively and recycled at least five times without a significant loss of adsorption capacity. In addition, the TPP-NH₂ can adsorb up to 33 times its own weight in organic solvents while wiping off the water. The high surface area, hierarchical porosity and π–π stacking interactions between the aromatic rings of MEB and MO and the aromatic rings of 3D TPP-NH₂ were responsible for the efficient adsorption. Therefore, the TPP-NH₂ that was synthesized using the facile strategy possesses significant potential in the treatment of wastewater.

1. Introduction

Organic dyes, as important industrial products, have been widely used for the dyeing of silk, cotton, linen and fiber. However, they also have many drawbacks, such as being toxic contaminants, being difficult to degrade and generating lot of pollution.⁴ So far, there are a few methods reported for removing dyes from aqueous solution, such as physical and chemical methods, biological oxidation, incineration and adsorption.⁵–⁷ Adsorption is considered as a promising method among them because it is highly effective, economical, and the simplest.⁵–⁷ However, using traditional porous adsorbents such as activated carbon,⁹ zeolites¹² and oriented clay films¹⁰ for the adsorption of dyes gave low adsorption capacities. Therefore, exploring novel adsorbents for the efficient adsorption and removal of dyes is still of great significance and is challenging.

Recently, porous organic polymers (POPs)¹³–¹⁴ have emerged as a new type of porous material with diverse applications because of their high surface areas, good porosities and structure diversity. In addition, POPs can be designed and constructed as task-specific materials by introducing multiple chemical functionalities into the porous network.¹⁵–¹⁷ A variety of amine functionalized POPs have been synthesized and developed for CO₂ capture and metal catalyst support due to their high surface areas, well defined pores and high physico-chemical stability.¹⁷,¹⁸ In principle, a highly porous polymer with amine functionality should also be an excellent adsorbent for the scavenging and removal of organic dyes and solvents. However, the application of POPs in organic dye and solvent removal has rarely been reported.

Herein, we report the first successful study on the application of an amine functionalized triptycene-based 3D porous polymer TPP-NH₂ as a new adsorbent for the fast adsorption and removal of organic dyes with different charges as well as organic solvents, to our knowledge. The adsorption kinetics, adsorption isotherms and the regeneration of the adsorbent for the removal of MEB and MO from aqueous solution on TPP-NH₂...
were studied in detail. The fast adsorption kinetics, large adsorption capacity, excellent physicochemical stability and good reusability make TPP-NH₂ a highly attractive adsorbent for the removal of organic dyes and solvents from aqueous solution.

2. Experimental

2.1 Reagents and materials

All of the purchased chemicals were of at least reagent grade and were used without further purification. Triptycene (98.0%), anhydrous ferric chloride (FeCl₃, ≥98.0%), 1,2-dichloroethane (DCE, 99.8%) and formaldehyde dimethyl acetal (FDA, ≥98.0%) were purchased from Sigma-Aldrich. All of the other solvents were purchased from TCI America.

2.2 Synthesis of TPP-NH₂

TPP-NH₂ was synthesized according to our previous work.²⁷ As shown in Scheme 1, 14-aminotriptycene²⁸ monomer (20 mmol) was dissolved in 20 mL of anhydrous DCE. To this solution, dimethoxymethane (FDA, 80 mmol) and anhydrous FeCl₃ (80 mmol) were added under a nitrogen environment. The resulting reaction mixture was heated to 80 °C for 24 h with constant stirring. After cooling to room temperature, the crude product was collected via filtration and repeatedly washed with methanol until the filtrate was nearly colourless. The product was further purified via Soxhlet extraction in methanol for 24 h. Finally, the product was dried at 120 °C under vacuum to give a brown solid, TPP-NH₂.

2.3 Instruments and characterization

The morphology of the samples was characterized using a Nova NanoS 450 field emission scanning electron microscope (FE-SEM). Transmission electron microscope (TEM) and high-resolution transmission electron microscope (HRTEM) images were recorded on a JEM-2100 transmission electron microscope. The powder samples were treated in ethanol using ultrasound for 20 min and were then dropped and dried on carbon-coated copper grids. The powder X-ray diffraction (XRD) data were collected on a D/Max2550 VB/PC diffractometer (40 kV, 200 mA) using Cu Kα radiation. The N₂ adsorption-desorption isotherms were measured at 77 K using a volumetric adsorption analyzer Micromeritics ASAP 2020. Before taking the adsorption measurements, the samples were degassed at 120 °C for 24 h. The specific surface areas were calculated using the Brunauer–Emmett–Teller (BET) method and the Langmuir method. The total volume was calculated at p/p₀ = 0.99. The micropore volume was derived from the t-plot method.

2.4 Adsorption test

Organic dye adsorption.²⁹ 10 mg of porous polymer TPP-NH₂ was added to 50 mL of MEB and MO solution with an initial concentration from 1 ppm to 1000 ppm under mechanical shaker conditions at a predetermined temperature (T = 30 °C, 40 °C, 50 °C) for 12 h. The residual concentration of the dyes was determined at the maximum wavelength (464 nm for MO and 664 nm for MEB) using UV-vis spectroscopy. The adsorption capacities and removal percentages of the dyes were calculated using the following equations:

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

\[
E\% = \frac{(C_0 - C_e)}{C_0} \times 100\%
\]

where \(q_e\) and \(E\%\) denote the adsorption capacity (mg g⁻¹) and removal percentage of the metal ions, respectively; \(C_0\) and \(C_e\) are the initial and equilibrium concentrations (mg L⁻¹) of the metal ions in aqueous solution, respectively; \(V\) is the volume of adsorption solution (mL); and \(m\) is the weight of the adsorbent (mg).

Organic dye adsorption kinetics. 10 mg of porous polymer TPP-NH₂ was added to 50 mL of MEB and MO solution with an initial concentration of 3 ppm under mechanical shaker conditions at 30 °C. After adsorption for a predetermined time (from 1 min to 540 min), the mixture was filtered with a 0.22 μm Millipore hydrophobic PTFE membrane, and the filtrate was measured using UV-vis spectroscopy.

**pH effect on MEB and MO adsorption.** The pH was adjusted using 0.1 M NaOH and 0.1 M HCl. The value of the pH was ranged from 2 to 9. Then, 10 mg of porous polymer TPP-NH₂ was added to 50 mL of a 1 ppm MEB and 3 ppm MO solution, with different pH values, at 30 °C.

Desorption experiments and reusability of TPP-NH₂. The adsorption/desorption cycles were conducted as follows: each adsorption experiment consisted of 50 mg of TPP-NH₂ with 250 mL of 1 ppm MEB and 3 ppm MO solution for 12 h. After the adsorption experiments, the dye-loaded powder in the flask was collected using filtration, washed with deionized water and stirred in water/ethanol for 24 h to desorb the dyes. The regeneration adsorbent was then filtered and separated from the solution, and washed with deionized water. Finally, the regeneration adsorbent was dried at 120 °C under vacuum. The desorption efficiency was calculated as the percentage of the mass of the analyte in the dyes desorbed to the elution medium of the total mass of the analyte adsorbed on the adsorbent. The above cycle was repeated 5 times.

**Organic solvent adsorption.** Six different kinds of organic solvent, including chloroform, dichloromethane, toluene, methanol, petroleum ether, and ethanol, were used in this

![Scheme 1](image-url) Schematic representation of the synthesis of the amine functionalized polymer TPP-NH₂.
study, and water was also used as a contrast. For example, in a typical adsorption experiment that used chloroform, a certain amount of TPP-NH$_2$ was weighed ($w_1$) and put in a small piece of glass tubing using adsorbent cotton at room temperature, and then the tube was fully immersed in chloroform overnight. A weight measurement ($w_2$) was done immediately after taking it out from the solvent to avoid evaporation. The adsorption capacity value, $W_t$ (w/w)% ($W_t$% = 100($w_2 - w_1$)/$w_1$), was thus calculated. Other solvent adsorption experiments were done in the same way.

3. Results and discussion

3.1 Characterization of TPP-NH$_2$

The synthesized TPP-NH$_2$ was characterized using SEM, HRTEM, XRD and $N_2$ adsorption–desorption. The SEM image (Fig. 1a) shows the solid sphere morphology of the sub-micrometer particles in TPP-NH$_2$. The HRTEM image (Fig. 1b) shows that TPP-NH$_2$ possesses a large amount of amorphous microporosity. In addition, with a broad band from 10$^\circ$ to 35$^\circ$ in the XRD pattern (Fig. 1c), it confirmed the amorphous nature of TPP-NH$_2$. The porosity of the TPP-NH$_2$ was evaluated using the nitrogen adsorption–desorption isotherm measurements at 77 K. As shown in Fig. 1d, the initial sharp increase in the uptake at low pressures ($p/p_0 < 0.1$) is characteristic of type I isotherms, i.e., TPP-NH$_2$ has significant microporous character. The hysteresis loops up to relative pressures above 0.9 suggest the occurrence of macropores and interparticle voids. The presence of minimal hysteresis in the $N_2$ desorption curves of these polymers suggests partial mesoporous character. In microporous polymers, this behaviour is usually ascribed to the swelling effects of the polymer networks while coming into contact with the adsorbate gas and expanded networks of 3D rigid building blocks.

The BET specific surface area of TPP-NH$_2$ measured from the nitrogen isotherms was 863 m$^2$ g$^{-1}$. The pore size distribution (PSD) curve (Fig. 1d (inset)), calculated based on non-local density functional theory (NLDFT), also confirmed the presence of primary micropores and a few mesopores, which can also be observed in some other polymers derived from using the “knitting” approach.$^{14}$

3.2 Adsorption kinetics of MEB and MO on TPP-NH$_2$

The time-dependent adsorption of MEB and MO on TPP-NH$_2$ was investigated at an initial concentration of 3 ppm. As shown in Fig. 2a and c, the removal percentage increased sharply within 60 min and slowed down thereafter, and, finally, reached equilibrium. It is worth mentioning that more than 95% of both dyes could be removed from the aqueous solution within 60 min at room temperature, indicating that the charge of the dyes plays no significant role in the kinetics of the adsorption. As the adsorption time went on, the adsorption intensities of the MEB and MO aqueous solution became weaker (Fig. 2b and d). Therefore, the optimum contact time was considered to be 60 min.

It was reported that more than 120 min is needed to attain equilibrium for dyes with several adsorbents, such as magnetic cellulose beads,$^{21}$ porous chromium benzenedicarboxylates (Cr-BDC),$^{22}$ and a mesoporous hybrid xerogel.$^{23}$ In contrast, in the present study, the percentage removal of dyes on TPP-NH$_2$ was very high (>95%) in a short amount of time. Therefore, fast adsorption is observed for TPP-NH$_2$, which can be attributed to its large surface area and hierarchical porosity.

Furthermore, the adsorption data are fitted with a pseudo-second-order kinetic model (eqn (3)) (Fig. 2e and f).

$$\frac{t}{q_t} = \frac{1}{k_2q_e^2} + \frac{t}{q_e} \quad (3)$$

where $q_t$ and $q_e$ are the adsorption capacity (mg g$^{-1}$) at a certain time $t$ (min) and equilibrium, respectively, and $k_2$ is the rate
constant for the pseudo-second-order adsorption (g mg⁻¹ min⁻¹). An extremely high correlation coefficient ($R^2 > 0.999$) is obtained for the pseudo-second-order kinetic model (Table 1). This suggests that the rate of adsorption of the organic dyes on the TPP-NH₂ adsorbent depends on the availability of adsorption sites.

### 3.3 Effect of pH on the adsorption of MEB and MO on TPP-NH₂

The effect of the pH on the adsorption of MEB and MO was studied in the pH range 2 to 9 (Fig. 3). The percentage removal of MEB on TPP-NH₂ was almost unchanged as the pH increased. However, a lower percentage of MO was observed at pH 2–3. The MO molecule was present in the solution as quinone and azo structures ($pK_a = 3.47$, Scheme 2), depending on the solution pH.²⁴ It is predicted that the azo structure should have a more conjugated π system than its quinone counterpart. Indeed, the adsorption capacity for MO under basic conditions was found to be higher than that under acidic conditions, predicting the role of π–π stacking interactions in the dye adsorption on TPP-NH₂. The above results show that the electrostatic interaction is not the main mechanism involved in the efficient adsorption of MEB and MO. The efficient adsorption performance of 3D TPP-NH₂ was attributed to its good porosity and high surface area, as well as the π–π stacking interactions between the aromatic rings of MEB and MO and the aromatics rings in the chain of TPP-NH₂.

### 3.4 Adsorption isotherms of MEB and MO on TPP-NH₂

The adsorption isotherms were studied at three different temperatures (30–50 °C) in the initial concentration range 1–1000 ppm (Fig. 4). The adsorption capacity of MEB and MO increased with the initial concentration of MEB and MO, indicating favorable adsorption of MEB and MO on TPP-NH₂ at high concentrations. To evaluate the maximum adsorption capacity of MEB and MO, the adsorption isotherms were fitted using the Langmuir model (eqn (4)):

$$\frac{C_e}{q_e} = \frac{C_e}{q_{\text{max}}} \frac{1}{q_{\text{max}} b}$$

where $C_e$ is the equilibrium concentration (mg L⁻¹) of the dyes, $q_e$ is the equilibrium adsorption capacity (mg g⁻¹) of the dyes, $q_{\text{max}}$ is the maximum adsorption capacity (mg g⁻¹), and $b$ is the Langmuir constant. The plots of $C_e/q_e$ against $C_e$ gave good linear plots at all of the test initial concentrations, indicating the adsorption of MEB and MO follows the Langmuir model.

<table>
<thead>
<tr>
<th>Dyes</th>
<th>$q_{\text{exp}}$ (mg g⁻¹)</th>
<th>$q_{\text{est}}$ (mg g⁻¹)</th>
<th>$k_2$ (g mg⁻¹ min⁻¹)</th>
<th>$R^2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>MEB</td>
<td>12.74</td>
<td>12.96</td>
<td>0.0059</td>
<td>0.999</td>
</tr>
<tr>
<td>MO</td>
<td>15.09</td>
<td>15.22</td>
<td>0.0043</td>
<td>0.999</td>
</tr>
</tbody>
</table>

### 3.5 Desorption and regeneration of TPP-NH₂

The pH studies and adsorption isotherms suggest that TPP-NH₂ is an efficient MEB and MO adsorber owing to its good porosity, high surface area and the π–π stacking interactions between dyes and TPP-NH₂, which indicate that the MEB and MO can be released from TPP-NH₂ by rinsing with water/ethanol. The desorption efficiency is up to 94%. To demonstrate the reusability of the TPP-NH₂ adsorbent, an adsorption–desorption cycle was repeated five times. As Fig. 5 shows, the adsorption capacity of the TPP-NH₂ adsorbent does not significantly change during the adsorption–desorption recycling. The results indicate that the TPP-NH₂ adsorbent is suitable for the efficient removal of dyes from aqueous solution.
3.6 Adsorption of the organic solvents on TPP-NH₂

In addition to the dye adsorption capacity, a high organic solvent adsorption capacity is also essential for water treatment. As shown in Fig. 6, the adsorption values of the organic solvents were in the range 1100–3300 wt%, while TPP-NH₂ only uptook less than 100 wt% of water. The organic solvent adsorption capacities on TPP-NH₂ show the sequence chloroform (3300 wt%) > chloromethane (2600 wt%) > toluene (2000 wt%) > methanol (1600 wt%) > petroleum ether (1505 wt%) > ethanol (1380 wt%) > water (85 wt%), which is a little different to the sequence of their polarities. The above results show that the difference in polarity of the organic solvents is not the only decisive factor involved in the efficient adsorption of the organic solvents. The excellent performance of TPP-NH₂ in adsorbing organic solvents might also be due to the hierarchical pores on TPP-NH₂.

4. Conclusions

An amine functionalized 3D porous organic polymer with a high surface area and good porosity was synthesized via a convenient and efficient Friedel–Crafts reaction. The obtained polymer TPP-NH₂ shows high adsorption capacities for organic dyes and solvents and thus can be used as an adsorbent to clean contaminated water. The ease of synthesis and low cost coupled with the efficient and rapid adsorption properties make TPP-NH₂ an attractive adsorbent for a wide range of large-scale applications in water purification and treatment.

Conflict of interest

The authors declare no competing financial interest.

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