

## Supporting information

### Application of $\text{Fe}_3\text{O}_4@\text{THAM-CH}_2\text{CH}_2\text{Cl}$ magnetic nanoparticle as a new adsorbent in ultrasonically assisted removal of Congo red from aqueous solutions

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#### 1. Methods

Design-Expert 10 software was applied in this research to design the experiments and analyze data. RSM was applied to optimize the variables to remove Congo red from the water environment by the proposed adsorbent. Standard stock solutions of Congo red ( $500 \text{ mg L}^{-1}$ ) were prepared, and working solutions were prepared daily by diluting the standard stock solutions with double distilled water. The chemical structure of CR is shown in Fig. S1.

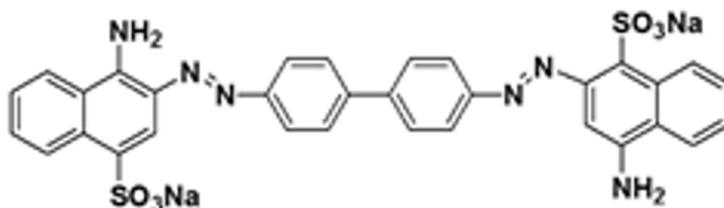


Fig. S1. The chemical structure of CR.

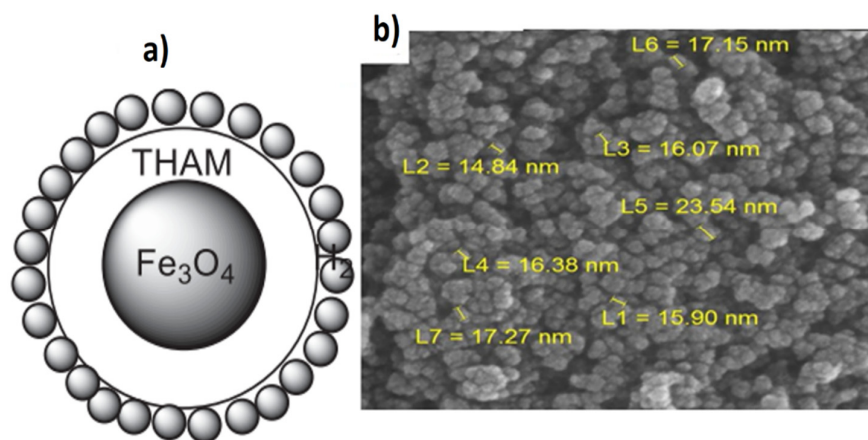
#### 2. Synthesis and properties of nanoparticle adsorbent

$\text{Fe}_3\text{O}_4@\text{THAM}$  MNP was synthesized as a chemical intermediate for the preparation of the MNPs catalyst [25]. The synthesis and characterization of  $\text{Fe}_3\text{O}_4$  MNPs have been discussed and confirmed using techniques such as FT-IR, EDS, MAP, SEM, XRD, BET, and VSM [25].

Briefly, the preparation of  $\text{Fe}_3\text{O}_4@\text{THAM-CH}_2\text{CH}_2\text{Cl}$  MNPs includes three steps: (1) First,  $\text{Fe}_3\text{O}_4$  nanoparticles were prepared by the co-precipitation method. (2) Then, the magnetite  $\text{Fe}_3\text{O}_4$  NPs were coated by tris(hydroxymethyl) aminomethane (THAM). (3) Finally,  $\text{Fe}_3\text{O}_4@\text{THAM-CH}_2\text{CH}_2\text{Cl}$  was prepared.

Fig. S2 shows the Scheme and FE-SEM image of the prepared  $\text{Fe}_3\text{O}_4@\text{THAM-CH}_2\text{CH}_2\text{Cl}$  NPs.

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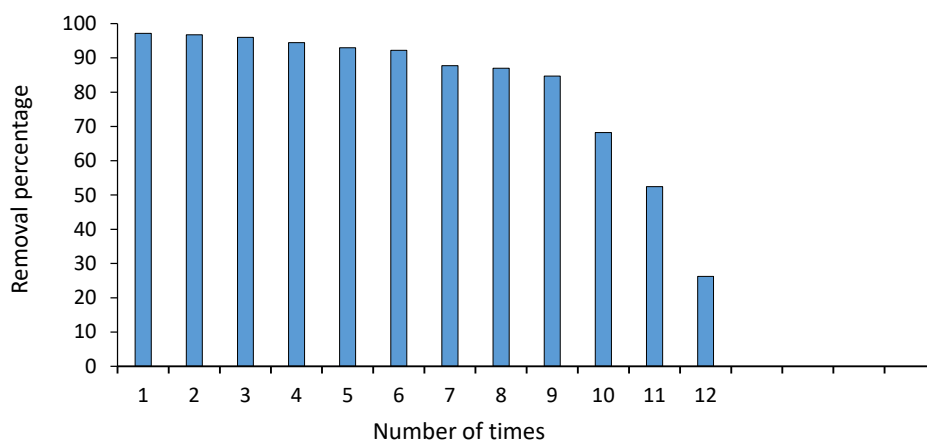
**Fig. S2.** Scheme (a) and FE-SEM images (b) of  $\text{Fe}_3\text{O}_4@$ THAM- $\text{CH}_2\text{CH}_2\text{Cl}$ .

Based on the FE-SEM image,  $\text{Fe}_3\text{O}_4 @$  THAM- $\text{CH}_2\text{CH}_2\text{Cl}$  MNPs show spherical nanoparticles. Based on the VSM, the value of saturation magnetization ( $M_s$ ) of  $\text{Fe}_3\text{O}_4$  MNPs was  $116.6 \text{ emu}\cdot\text{g}^{-1}$ . Also, the BET measurements confirmed the specific surface area of  $121.79 \text{ m}^2\cdot\text{g}^{-1}$ , total pore volume of  $27.9 \text{ cm}^3\cdot\text{g}^{-1}$ , and the mean pore size of  $8.14 \text{ nm}$ . Due to pores on the  $\text{Fe}_3\text{O}_4 @$  THAM- $\text{CH}_2\text{CH}_2\text{Cl}$  adsorbent surface, the proposed adsorbent can remove Congo red from the aqueous environment by trapping the pollutant in its pores.

### 3. Recovery of adsorbent

In this work, the possibility of reusing the proposed adsorbent to remove Congo red was investigated in terms of the number of absorption and desorption times.

According to the results (Fig. S3), about a 5% change in adsorbent efficiency is observed after 6 runs of adsorption and desorption. This confirms that the proposed adsorbent can be regenerated and reused.



**Fig. S3.** Recovery of adsorbent.

Also, the isotherm for the adsorption process of Congo red by the proposed adsorbent was assayed. According to the results (Table S1), the Langmuir adsorption isotherm has a larger correlation coefficient ( $R^2 = 0.9987$ ), which indicates the best fit with the experimental data and that the proposed adsorbent is more suitable for the Congo red adsorption process.

**Table S1.** Isotherm models constants for the adsorption process of Congo dye red on the adsorbent.

<b>Isotherm</b>	<b>Equation</b>	<b>Plot</b>	<b>Parameters</b>	<b>Value</b>
Langmuir	$C_e/q_e = C_e/q_m + 1/(q_m k_L)$	$C_e/q_e$ against $C_e$	$q_m$ (mg. g <sup>-1</sup> )	137
			$k_L$ (L. mg <sup>-1</sup> )	1.4
			$R^2$	0.9987
Freundlich	$\log q_e = \log k_F + (1/n)\log C_e$	$\log q_e$ against $\log C_e$	$n$	1.05
			$k_F$ (L. mg <sup>-1</sup> )	159.8
			$R^2$	0.966