

Temporal control of RAFT polymerization *via* Magnetic Catalysis

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- **Characterization data of the magnetic $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$ catalyst**

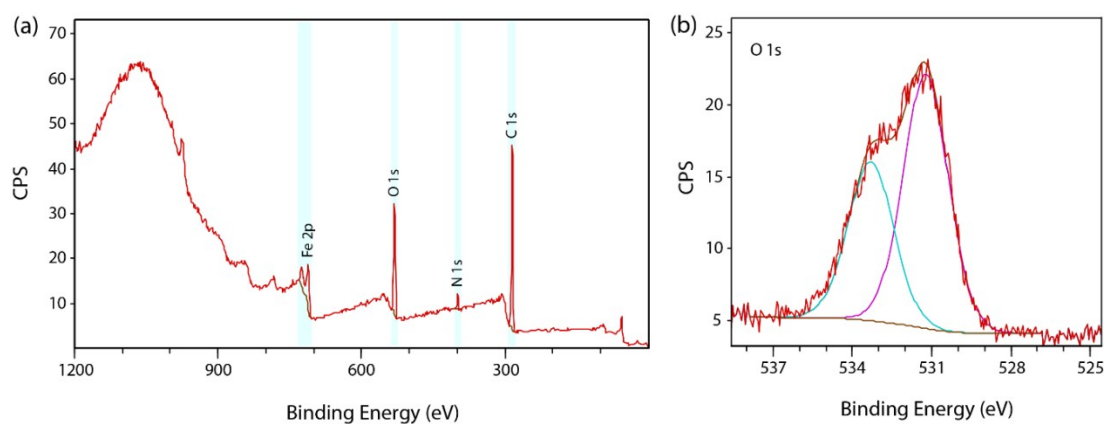


Fig. S1 Survey XPS spectrum of $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$ (a), O1s orbital (b).

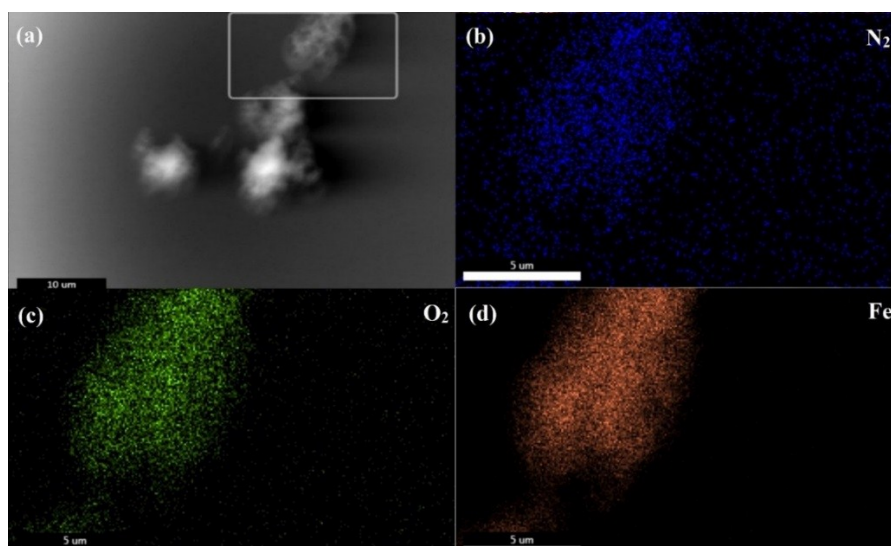


Fig. S2 (a) STEM-HAADF images of $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$, and (b-d) the corresponding EDS mappings.

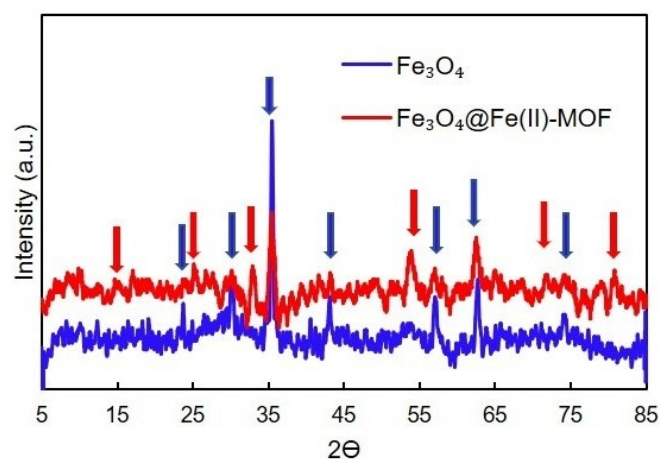


Fig. S3 XRD of Fe_3O_4 and $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$ particles.

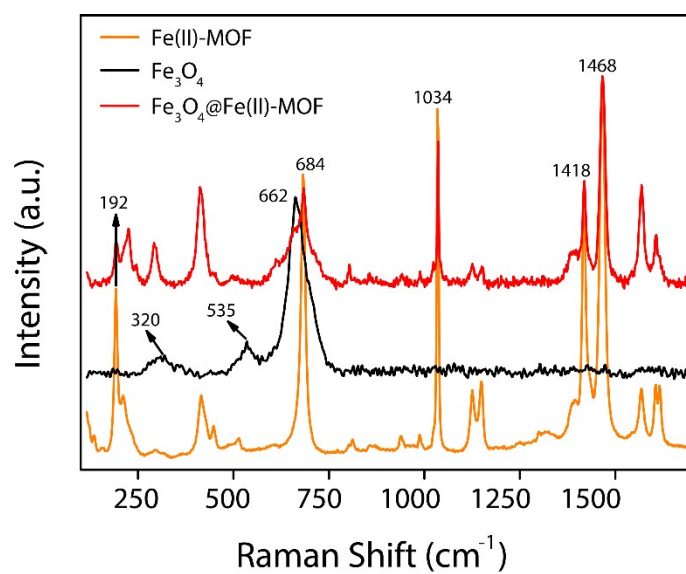


Fig. S4 Raman spectra of Fe_3O_4 , Fe(II)-MOF , and $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$ particles.

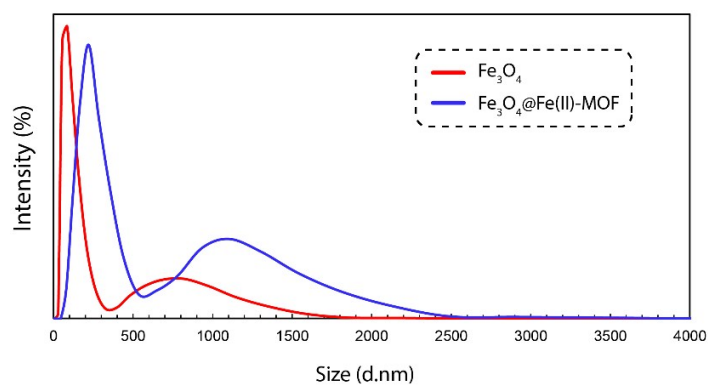


Fig. S5 DLS spectra of Fe_3O_4 and $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$ particles.

- ***Zeta potential of the magnetic catalysts***

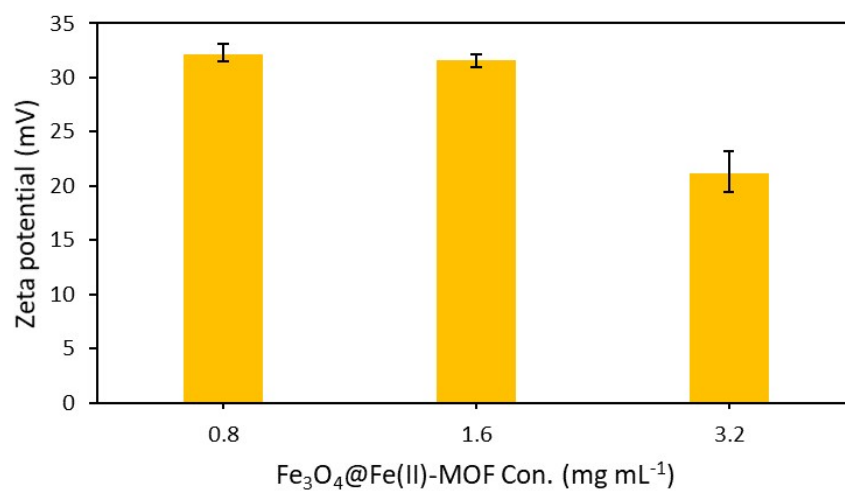


Fig. S6 Zeta potential of the Fe₃O₄@Fe(II)-MOF NPs at different concentrations as 0.8, 1.6, and 3.2 mg mL⁻¹ obtained from three runs.

- **Characterization data of the synthesized polymers via the magnetic Fenton-RAFT**

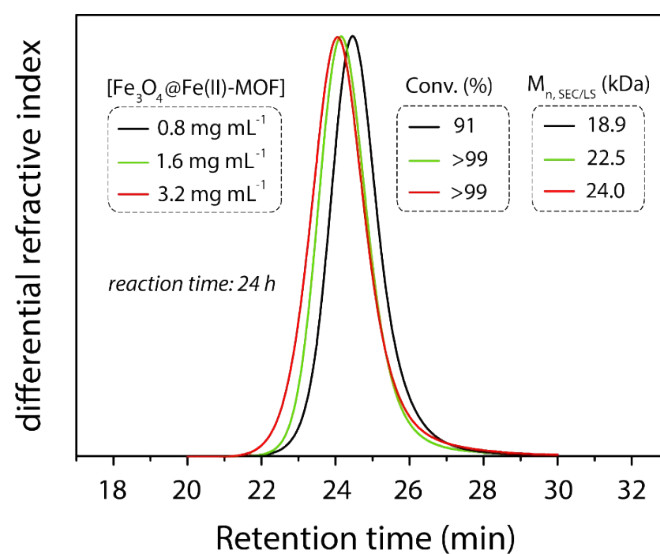


Fig. S7 SEC chromatograms of the synthesized PDMA via the magnetic Fenton-RAFT after 24 h with different concentrations of $\text{Fe}_3\text{O}_4@\text{Fe(II)-MOF}$ catalysts.

- **Kinetic studies**

Table S1. Characterization data of the synthesized polymers within the kinetic studies on the magnetic Fenton-RAFT polymerization of DMA with $[\text{Fe(II)-MOF@Fe}_3\text{O}_4] = 1.6 \text{ mg mL}^{-1}$ and $[\text{H}_2\text{O}_2]_0 = 35 \text{ mM}$.

Time (h)	Conversion (%) ^a	$M_{n,\text{the.}}$ (Da) ^b	$M_{n,\text{SEC/LS}}$ (Da) ^c	$\bar{D}$ (-) ^c
0	0	-	-	-
1	40	8,200	8,500	1.03
2	49	10,000	10,500	1.04
3	55	11,200	11,100	1.04
4	60	12,200	12,000	1.03
5	64	13,000	13,600	1.02
6	65	13,200	13,900	1.02
^a conversion values were calculated from ^1H NMR analysis.; ^b theoretical molecular weights were calculated according to this formula: $M_{n,\text{the.}} = \text{DP} \times \text{conv.} \times \text{MW}_{\text{monomer}} + \text{MW}_{\text{RAFT agent.}}$; ^c experimental molecular weight and dispersity values were obtained from LS analysis coupled with SEC.				

- **Temporal control of Fenton-RAFT polymerization**

Table S2. Characterization data of the synthesized PDMA's within the temporal control of Fenton-RAFT polymerization with $[\text{Fe(II)-MOF@Fe}_3\text{O}_4] = 1.6 \text{ mg mL}^{-1}$ and $[\text{H}_2\text{O}_2]_0 = 35 \text{ mM}$.

Time (h)	Conversion (%) ^a	$M_{n,\text{the.}}$ (Da) ^b	$M_{n,\text{SEC/LS}}$ (Da) ^c	$\bar{D}$ (-) ^c
0	0	-	-	-
1	45	9,200	9,400	1.05
2	45	9,200	9,350	1.05
3	51	10,400	10,350	1.04
4	51	10,400	10,450	1.04
5	59	12,000	11,900	1.03
6	59	12,000	11,950	1.03
7	66	13,350	13,600	1.03
^a conversion values were calculated from ^1H NMR analysis.; ^b theoretical molecular weights were calculated according to this formula: $M_{n,\text{the.}} = \text{DP} \times \text{conv.} \times \text{MW}_{\text{monomer}} + \text{MW}_{\text{RAFT agent.}}$; ^c experimental molecular weight and dispersity values were obtained from LS analysis coupled with SEC.				