

**Facile cascade-anchored synthesis of ultrahigh metal loading single-atom for
significantly improved Fenton-like catalysis**

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Supplementary Note 1: Chemicals and Materials

Iron nitrate nonahydrate ($\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$, 99%) and melamine ($\text{C}_3\text{H}_6\text{N}_6$, 99%) were bought from Macklin Ltd. (Shanghai, China). Sodium sulfate (Na_2SO_4 , 99%), potassium peroxymonosulfate (PMS, $\text{KHSO}_5 \cdot 0.5\text{KHSO}_4 \cdot 0.5\text{K}_2\text{SO}_4$), sodium bicarbonate (NaHCO_3 , 99%), sodium dihydrogen phosphate (NaH_2PO_4 , 99%), sulfuric acid (H_2SO_4), sodium hydroxide (NaOH , 99%), potassium thiocyanate (KSCN , 99%), 5,5-dimethyl-1-pyrroline N-oxide (DMPO, 97%), methyl phenyl sulfoxide (PMSO) and Furfuryl alcohol (FFA) were purchased from Aladdin Ltd. (Shanghai, China). Methanol (MeOH , 99.9%) was purchased from Tianjin Kemiou Chemical Reagent Co., Ltd (Tianjin, China). Tertbutyl alcohol (TBA, > 99.5%), Nafion solution (5 wt.%) and oxalic acid (OA, 99%) were purchased from Sinopharm Group Chemical Reagent Co., Ltd. (Shanghai, China). 2,2,6,6-Tetramethylpiperidine (TEMP) was bought from Tianjin Sinos Opto Technology Co., Ltd (Tianjin, China). The studied model pollutants, bisphenol A (BPA), phenol (BP), Rhodamine B (RhB), tetracycline (TC), methyl orange (MO) and sulfamethoxazole (SMX) were purchased from Aladdin Ltd. (Shanghai, China). All chemicals were of analytical grade and were used without further purification. All solutions used in this work were prepared using ultrapure water.

Supplementary Note 2: Characterization methods

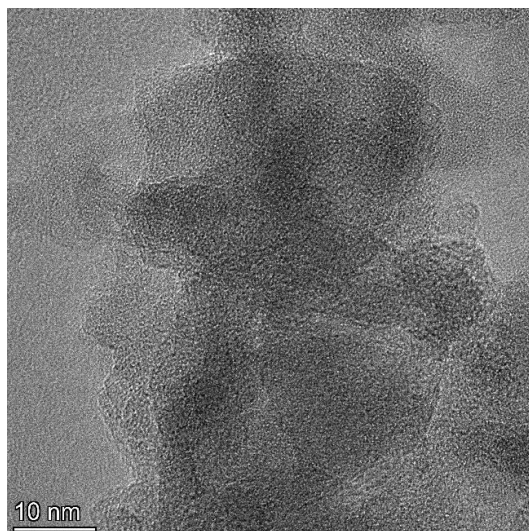
XAS measurements were carried out in transmission mode using a commercial laboratory device (easyXAFS300, easy XAFS LLC, Renton, WA). This instrument is based on Rowland circle geometries and is equipped with spherically bent crystal analyzers (SBCAs) and a silicon drift detector (AXAS-M1, KETEK GmbH, Munich, Germany). The morphologies of catalysts were observed by the transmission electron microscopy (TEM, JEM-2800). Aberration-corrected high-angle annular dark-field scanning transmission electron microscopy (AC-HAADF-STEM) images were recorded on a high resolution transmission electron microscope (JEM-ARM200F) with a field-emission gun at 200 kV. X-ray photoelectron spectroscopy (XPS, Thermo Scientific ESCALAB 250Xi) was utilized to analyze the bonding state of elements on the catalyst surface. Fourier transformed infrared spectroscopy (FTIR) was conducted by Thermo Scientific Nicolet IS50 FTIR (Thermo Fisher, USA). In-situ FTIR was carried out using an external reflection configuration and the Ge crystal was used as prism. Raman analysis was carried out using a confocal microscopic Raman spectrometer (LabRam HR, Horiba, France). X-ray diffraction (XRD) patterns of catalysts were characterized by a Philips-12045 B/3 diffractometer. BET specific surface areas and pore distribution of the catalysts were measured through a Micromeritics ASAP 2460 instrument. The atomic ratios of Fe in the samples were analyzed using an inductively coupled plasma atomic emission spectrometer (ICP- AES) (IRIS Intrepid II XSP) to reveal the catalyst composition. Electron paramagnetic resonance (EPR) was used to analyze the reactive species by a Bruker EMX Nano (Germany).

Supplementary Note 3: Electrochemical measurements

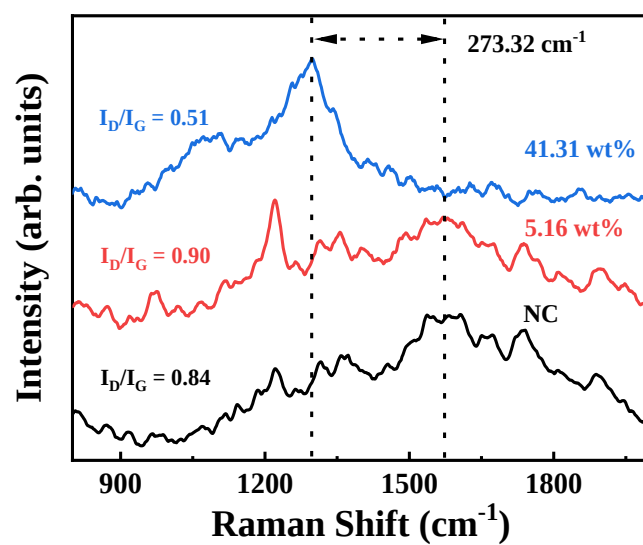
The electrochemical performance was tested by a CHI 760E potentiostat (CH Instruments, Chenhua, Shanghai, China). The catalyst was coated on a glassy carbon electrode as a working electrode (5 mg of catalyst, 50 μ L of Nafion and 1 mL of ethanol were mixed by ultrasonication, and 10 μ L of the mixture was dropped onto the glassy carbon electrode to dry naturally, the catalyst loading was 0.25 mg cm⁻²), a platinum sheet was used as the counter electrode, and Ag/AgCl electrode was used as the reference electrode. The electrode potential was converted to a reversible hydrogen electrode (RHE) reference scale using $E_{RHE}=E_{Ag/AgCl}+0.197+0.0592(\text{pH})$. The linear sweep voltammetry (LSV) curve was obtained at the scanning rate of 10 mV s⁻¹ in the electrolyte of Na₂SO₄ (50 mM, pH 7). Chronopotentiometry and chronoamperometry analyses were performed to obtain the surface oxidation potential and current response in the reaction system. PMS and SMX samples were added into the electrolyte at stated intervals, with final concentrations of 2 mM and 10 mg L⁻¹, respectively.

Supplementary Note 4: Quantum chemical calculations

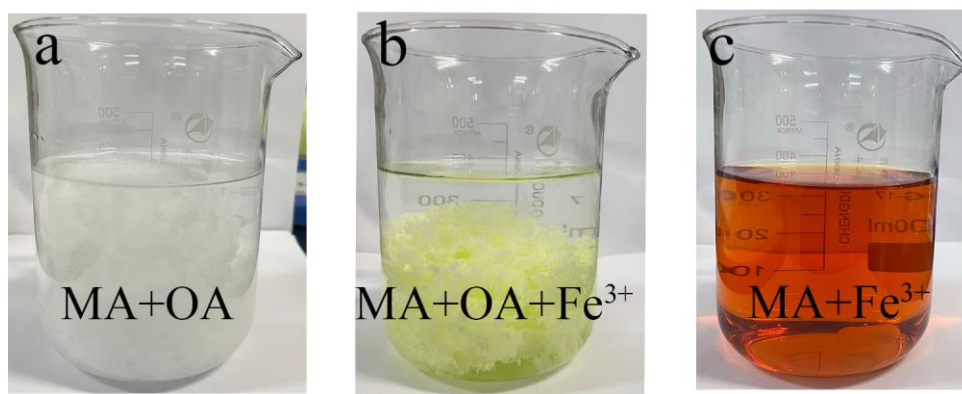
All spin-polarized density-functional theory (DFT) was calculated through projective enhanced wave method in the Vienna ab initio simulation package (VASP 5.4.1) based on the plane wave base set^{1,2}. Generalized gradient approximation (GGA) and PBE parameterization were used to deal with the exchange correlation potential³. Grimme's DFT-D3 model was applied for Van der Waals correction (6). To avoid interaction between adjacent images, a vacuum area of about 18 Å was used. The energy cutoff was set at 450 eV. The Brillouin-zone integrals were sampled using a Γ -centered Monkhorst-Pack lattice ($2 \times 2 \times 1$). Considering the strong d-electron correlation of 3d TMs (4th row in the periodic table), PBE + U is necessary. The corresponding U value of Fe was set to 3.29 according to Ref.^{4,5}. The structure was completely relaxed until the maximum force on each atom was less than 0.02 eV/Å, with an energy convergence criterion of 10^{-5} eV. The free energies (G) of different intermediates are defined as $G = E_i - E_{\text{reactant}}$ (E_i is the energy of intermediates and E_{reactant} is the total energy of reactants) and finally obtained by $G = E_{\text{total}} + E_{\text{ZPE}} - TS$, where E_{total} , E_{ZPE} , and TS are the ground-state energy, zero-point energies, and entropy terms, respectively, with the latter two taking vibration frequencies from DFT.



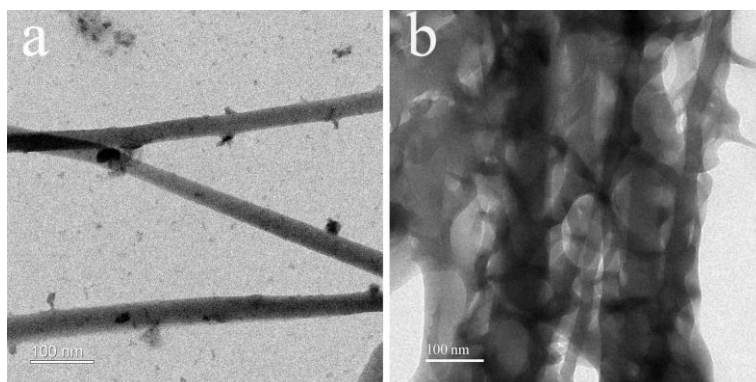
Supplementary Fig. 1 | HRTEM images of Fe-SAC-41.31.



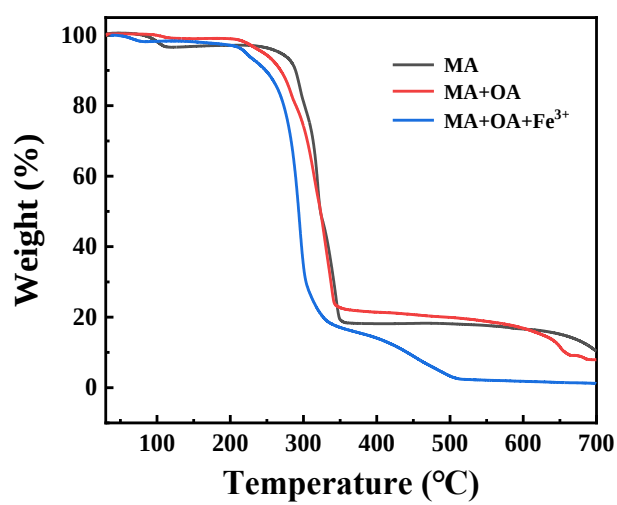
Supplementary Fig. 2 | Raman spectra of as prepared catalysts with different Fe loading.



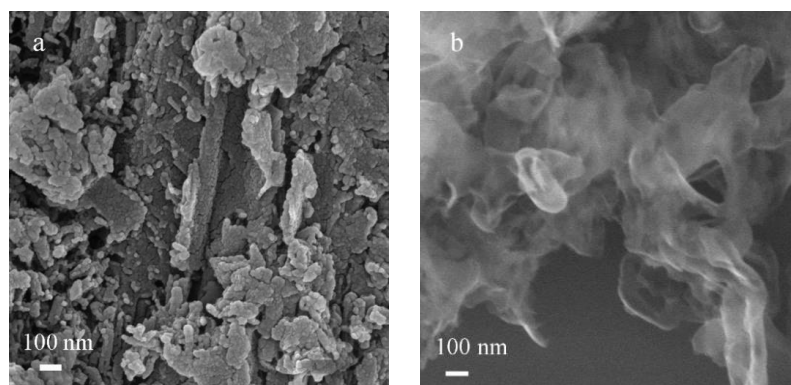
Supplementary Fig. 3 | **a**, Supramolecular organic skeleton generated by the complexation reaction between MA and OA; **b**, SAC precursor after anchoring Fe³⁺; **c**, SAC precursor prepared by conventional steam-drying solvent method.



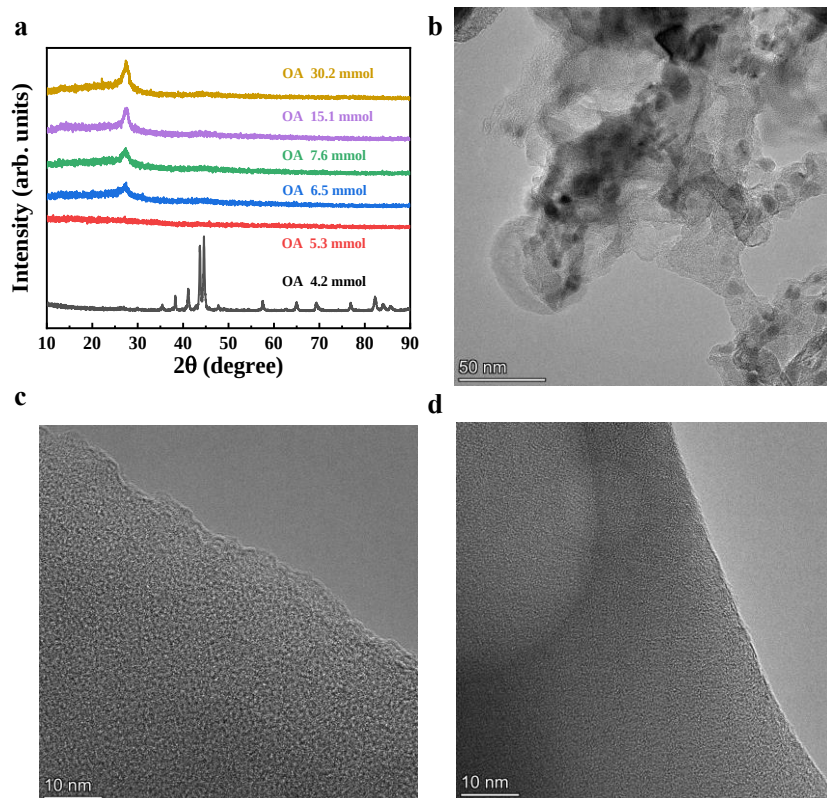
Supplementary Fig. 4 | TEM images of Fe-SAC precursor obtained by **a**, conventional solvent evaporation method; **b**, cascade anchoring strategy.



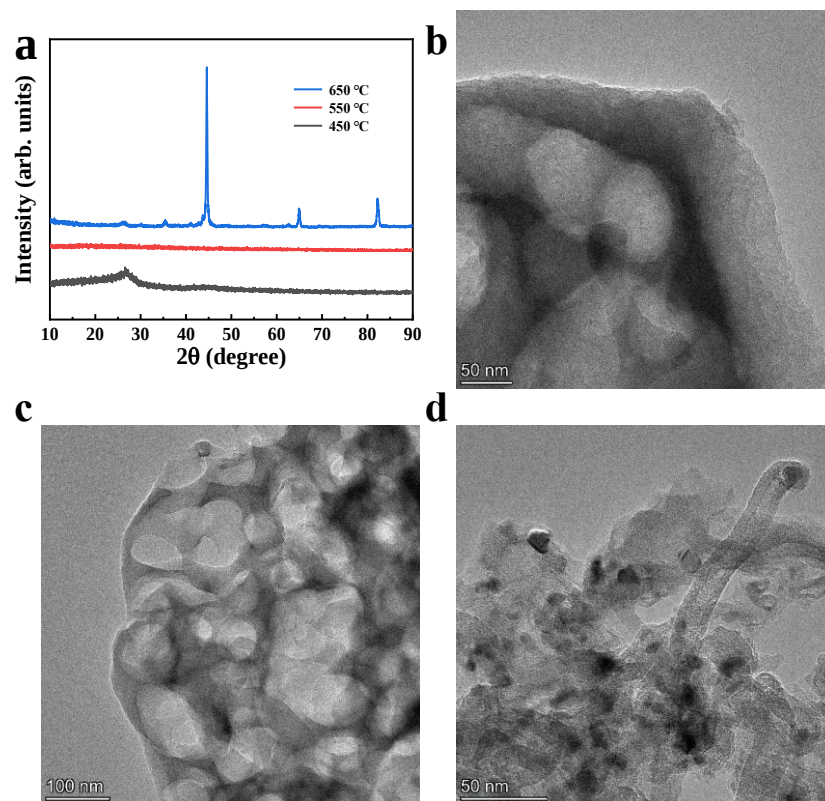
Supplementary Fig. 5 | Thermogravimetric curves of MA, MA+OA supramolecular organic skeleton, and Fe-SAC precursor.



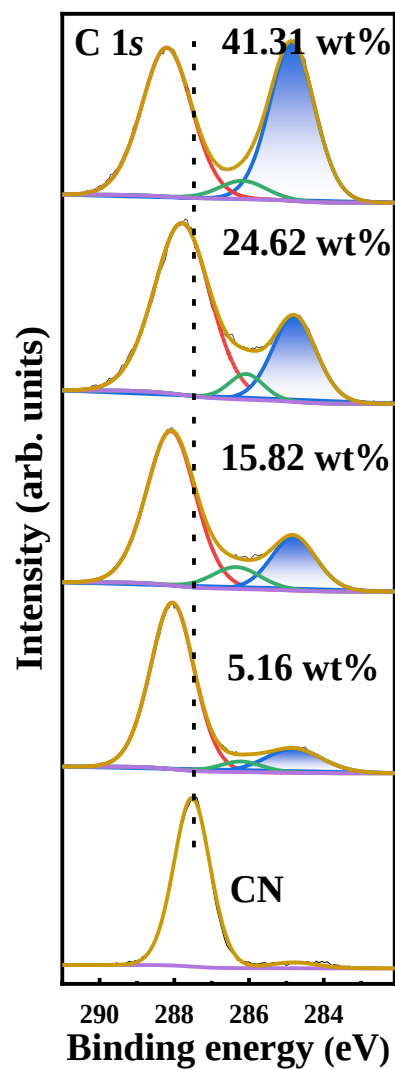
Supplementary Fig. 6 | SEM images of C₃N₄ prepared by pyrolysis of MA **(a)** or MA+OA supramolecular polymers **(b)**.



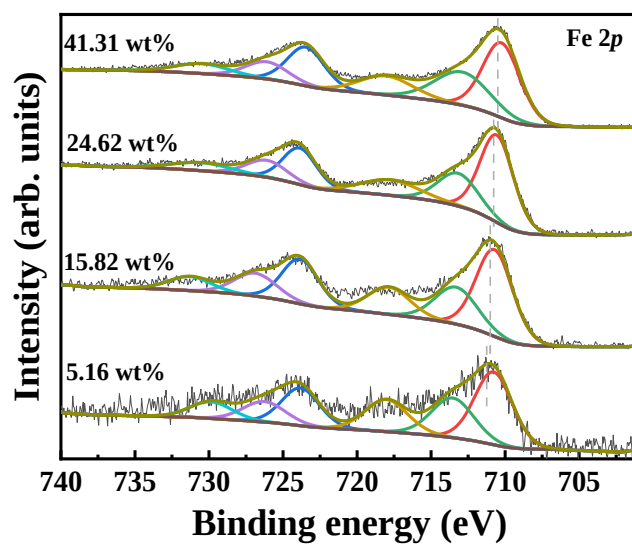
Supplementary Fig. 7 | **a**, XRD patterns of catalysts prepared with different OA amount. **b-d**, HRTEM images of Fe-SAC at OA addition of 4.2, 5.3, and 30.2 mmol, respectively.



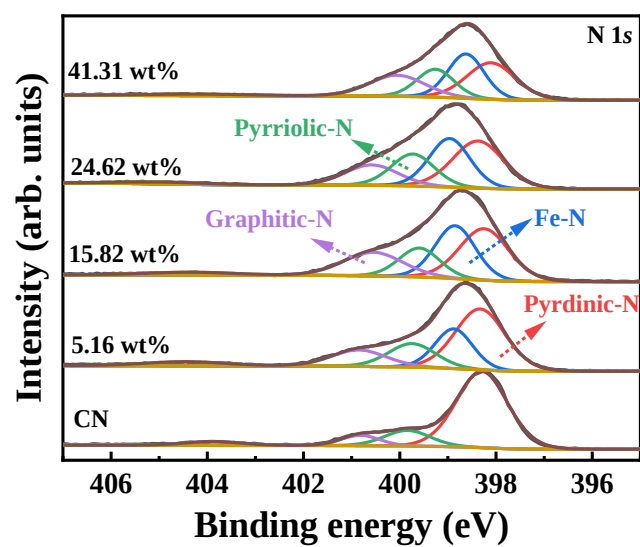
Supplementary Fig. 8 | a, XRD patterns of catalysts calcined at different temperatures.
b-d, HRTEM images of catalysts calcined at 450, 550 and 650°C, respectively.



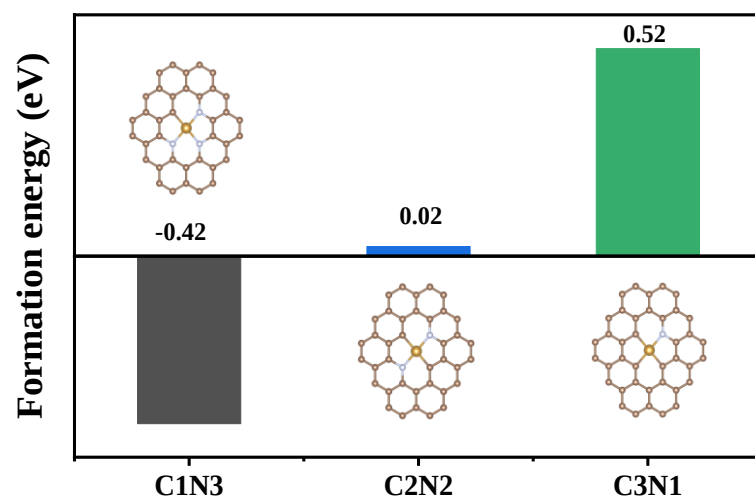
Supplementary Fig. 9 | High-resolution XPS spectra of C 1s for the Fe-SAC-*x* samples.



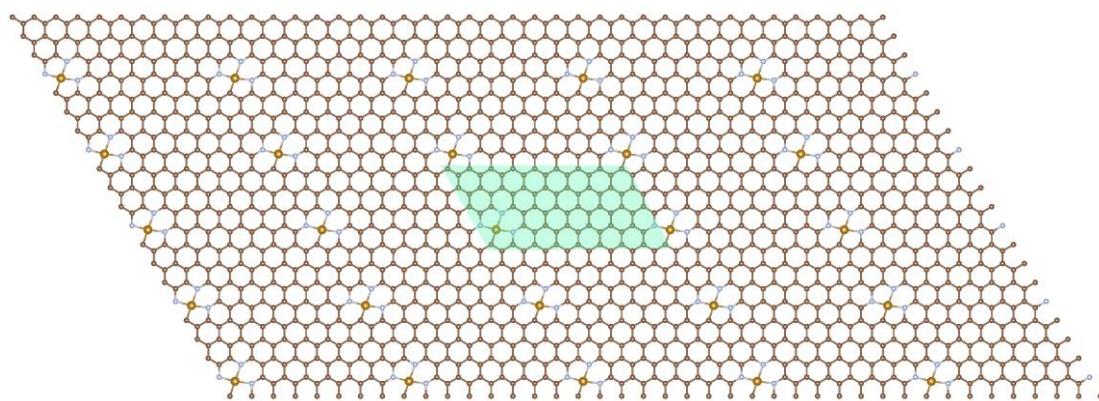
Supplementary Fig. 10 | Fe 2p XPS spectra of Fe-SAC- x ($x = 5.16$ – 41.31 wt%).



Supplementary Fig. 11 | High-resolution XPS spectrum of N 1s.



Supplementary Fig. 12 | The binding energies of the possible CN coordination modes.

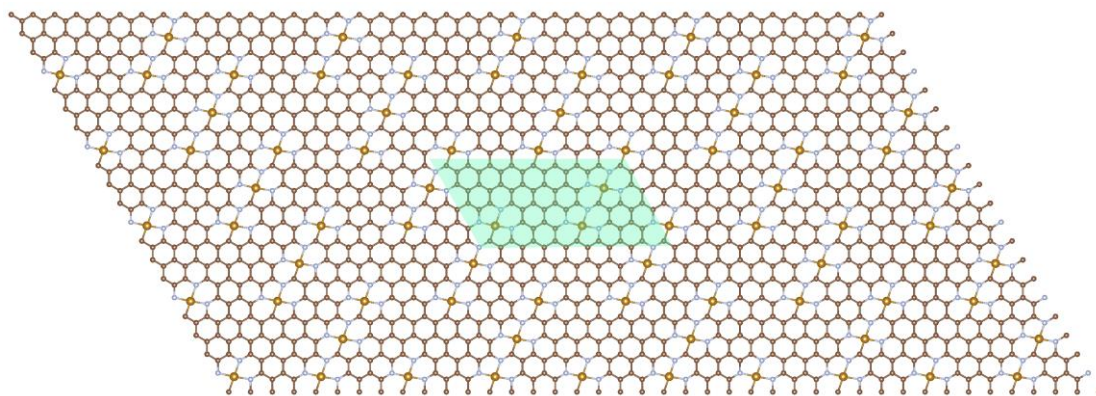


Supplementary Fig. 13 | The scaled-up structure model of Fe-SAC-5.16. The core computing unit cells used in DFT calculations were the green area.

Sample: Fe-SAC-5.16

Number of atoms: 25 Fe, 75 N, 1546 C

Theoretical Fe loading: $\text{Fe}(\text{wt}\%) = \frac{25 \times 56}{25 \times 56 + 75 \times 14 + 1546 \times 12} = 6.66\text{wt}\%$

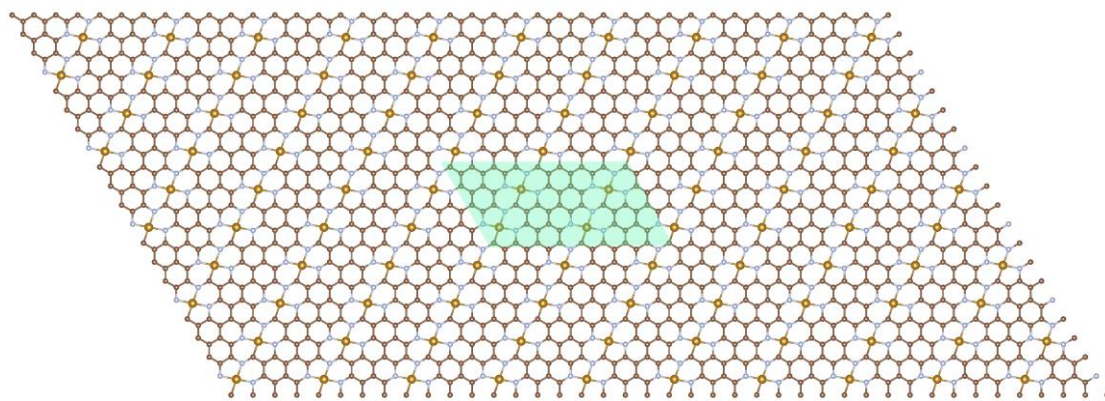


Supplementary Fig. 14 | The scaled-up structure model of Fe-SAC-15.82. The core computing unit cells used in DFT calculations were the green area.

Sample: Fe-SAC-15.82

Number of atoms: 75 Fe, 225 N, 1286 C

Theoretical Fe loading: $\text{Fe}(\text{wt}\%) = \frac{75 \times 56}{75 \times 56 + 225 \times 14 + 1286 \times 12} = 18.43\text{wt}\%$

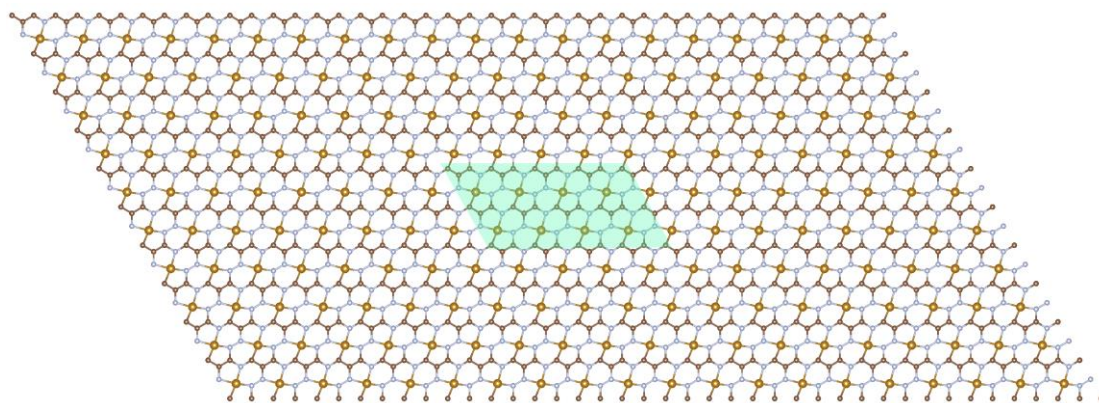


Supplementary Fig. 15 | The scaled-up structure model of Fe-SAC-24.62. The core computing unit cells used in DFT calculations were the green area.

Sample: Fe-SAC-24.62

Number of atoms: 100 Fe, 300 N, 1161 C

Theoretical Fe loading: $\text{Fe}(\text{wt}\%) = \frac{100 \times 56}{100 \times 56 + 300 \times 14 + 1161 \times 12} = 23.59\text{wt}\%$

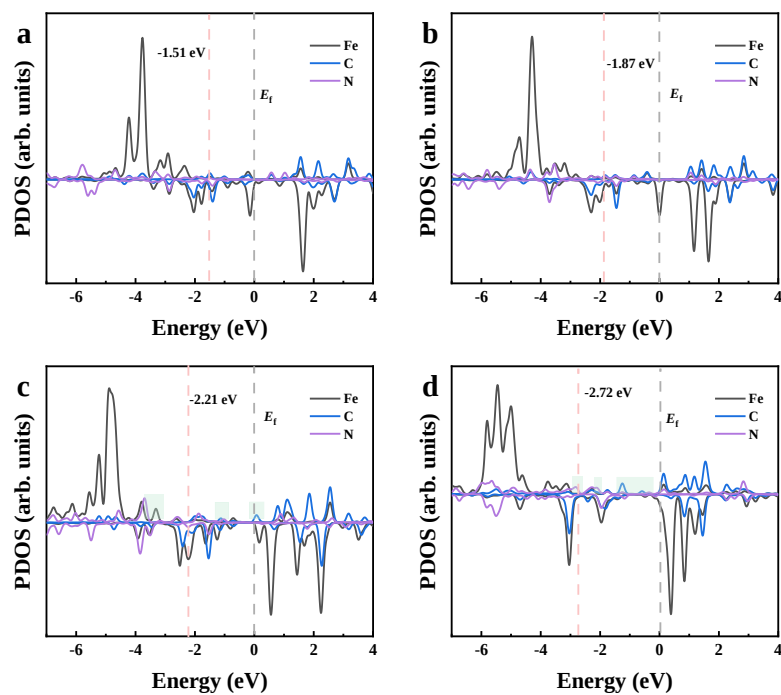


Supplementary Fig. 16 | The scaled-up structure model of Fe-SAC-41.31. The core computing unit cells used in DFT calculations were the green area.

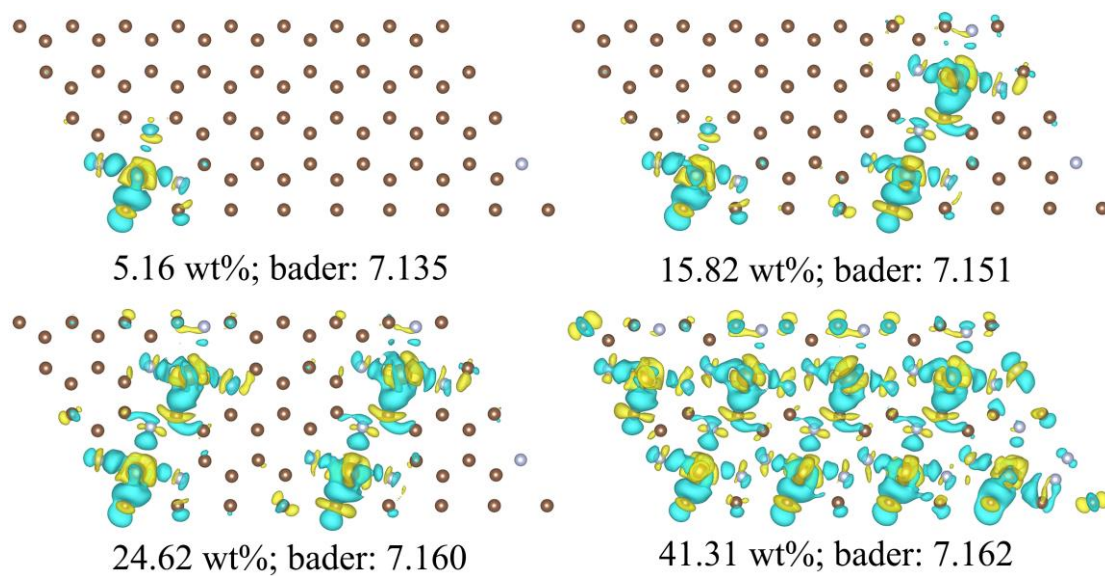
Sample: Fe-SAC-41.31

Number of atoms: 200 Fe, 600 N, 661 C

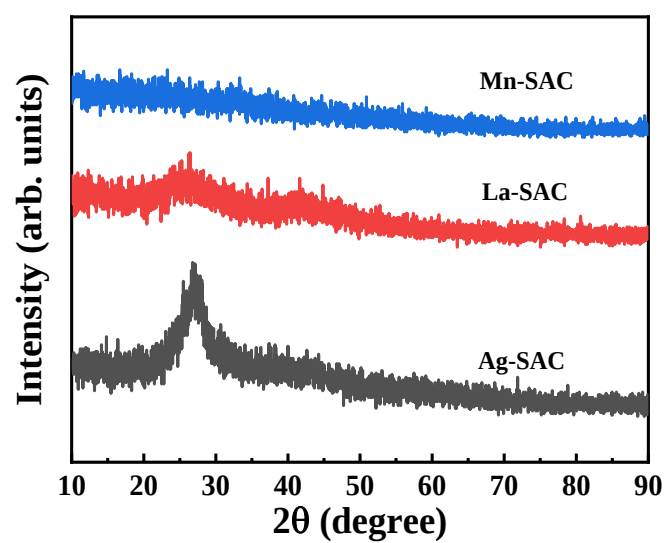
Theoretical Fe loading: $\text{Fe}(\text{wt}\%) = \frac{200 \times 56}{200 \times 56 + 600 \times 14 + 661 \times 12} = 40.68\text{wt}\%$



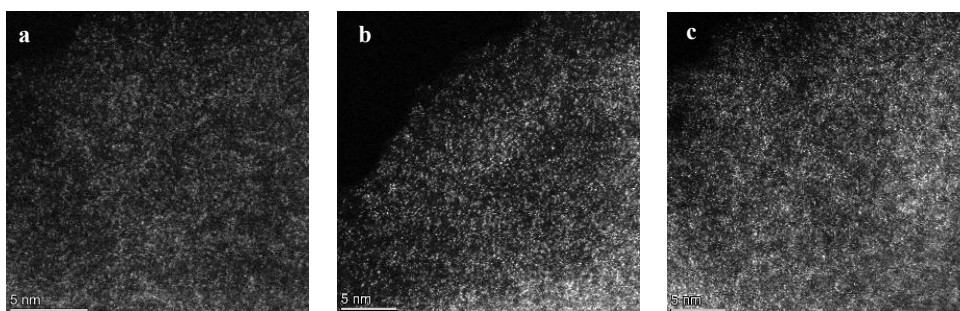
Supplementary Fig. 17 | Projected density of state (PDOS) of Fe 3*d*, N 2*p*, and C 2*p* orbital of Fe-SAC-5.16 **(a)**, Fe-SAC-15.82 **(b)**, Fe-SAC-24.62 **(c)** and Fe-SAC-41.31 **(d)**, respectively.



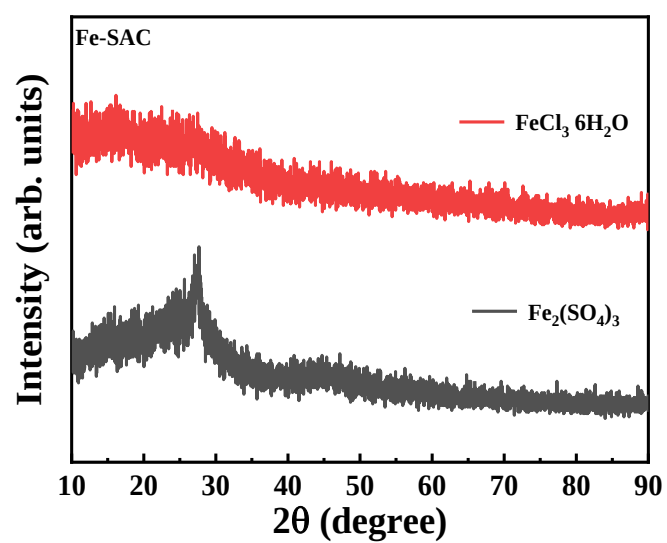
Supplementary Fig. 18 | Differential charge density of Fe-SAC-*x* models.



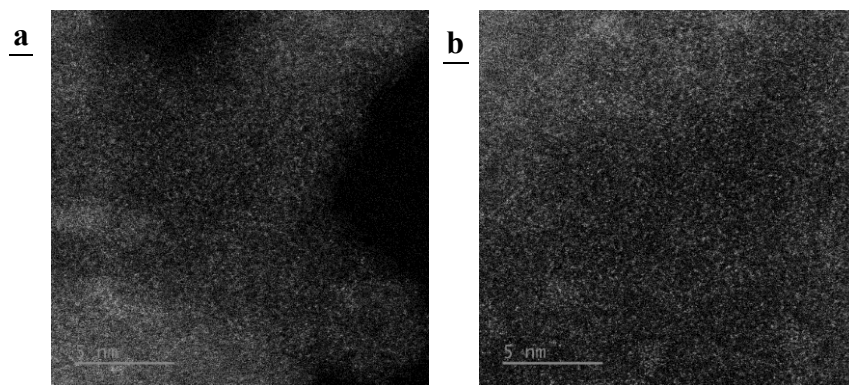
Supplementary Fig. 19 | XRD patterns of as-prepared Mn-SAC, La-SAC and Ag-SAC.



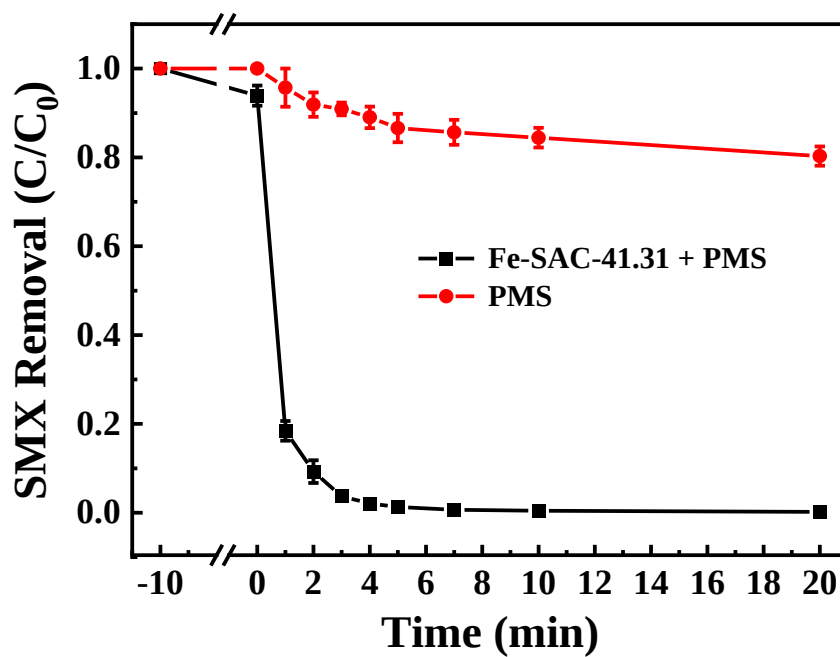
Supplementary Fig. 20 | HAADF-STEM images of **a**, Mn-SAC; **b**, La-SAC and **c**, Ag-SAC.



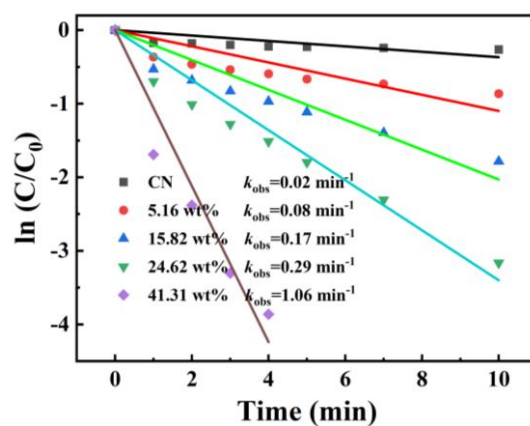
Supplementary Fig. 21 | XRD images of Fe-SAC prepared using FeCl₃·6H₂O or Fe₂(SO₄)₃ as metal sources, respectively.



Supplementary Fig. 22 | HAADF-STEM images of Fe-SAC prepared using $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ **(a)** and $\text{Fe}_2(\text{SO}_4)_3$ **(b)** as metal sources, respectively.

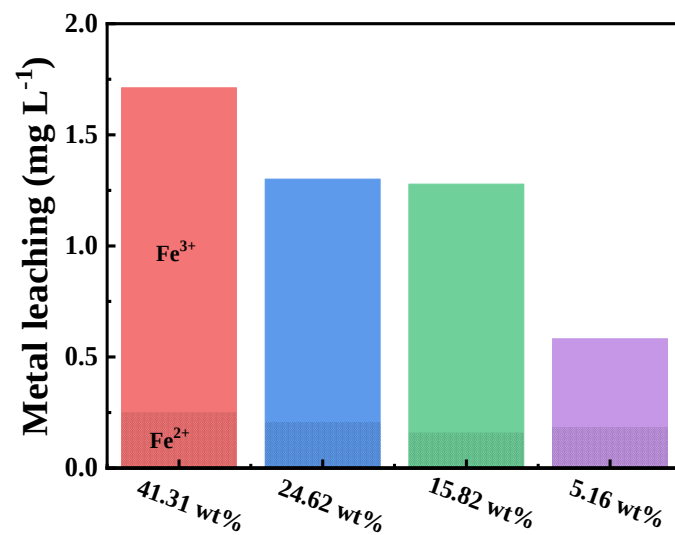


Supplementary Fig. 23 | Degradation effect of PMS on SMX. The error bars in the figures represent the standard deviations from triplicate tests.

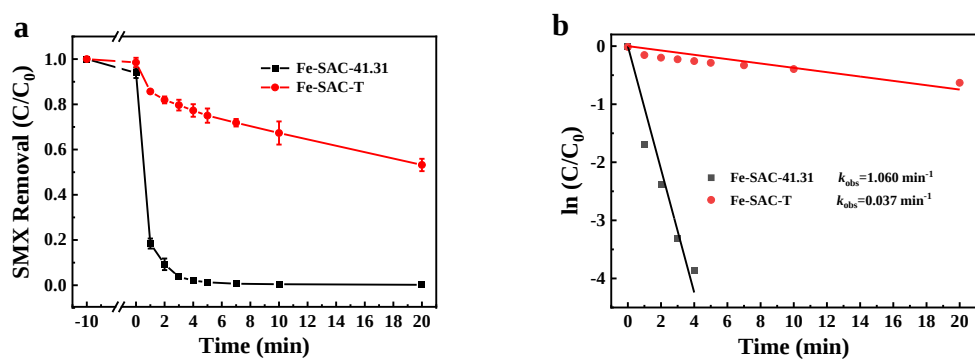


Supplementary Fig. 24 | The degradation kinetics of SMX in Fe-SAC-*x*/PMS system.

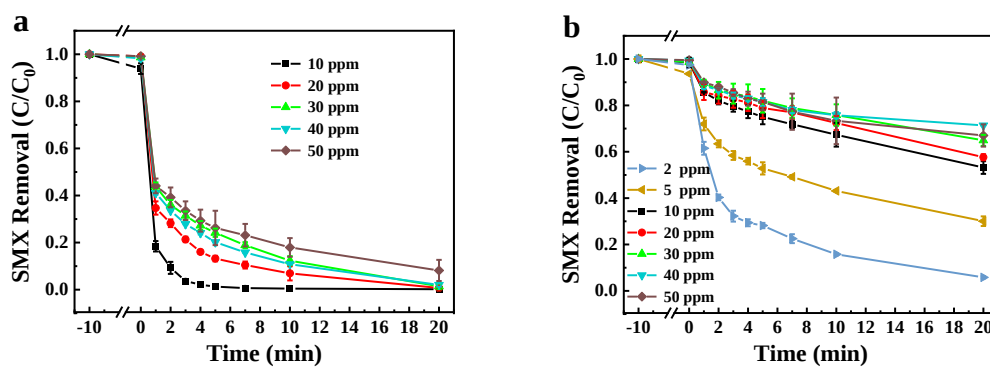
The model used to fit the data was a pseudo-first-order kinetic model.



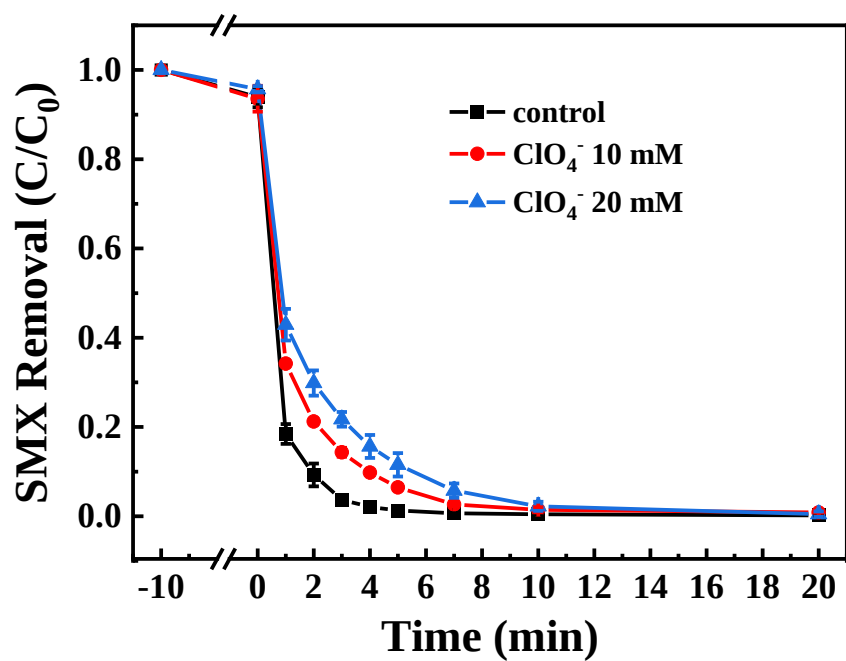
Supplementary Fig. 25 | Iron dissolution of Fe-SAC-x during degradation. The shaded area represents Fe²⁺.



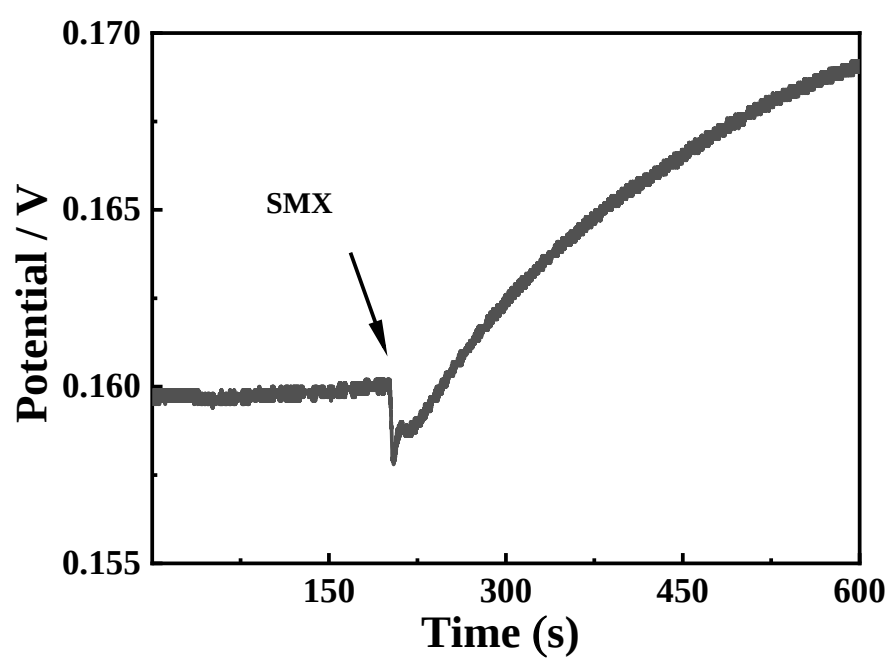
Supplementary Fig. 26 | a, SMX degradation performance. **b**, SMX degradation kinetics in the Fe-SAC-41.31/PMS and Fe-SAC-T/PMS system, respectively. Experimental conditions: $[SMX]_0 = 10 \text{ mg L}^{-1}$, $[PMS]_0 = 2 \text{ mM}$, $\text{pH} = 7$, catalyst: 0.1 g L^{-1} . The error bars in the figures represent the standard deviations from triplicate tests. The model used to fit the data was a pseudo-first-order kinetic model.



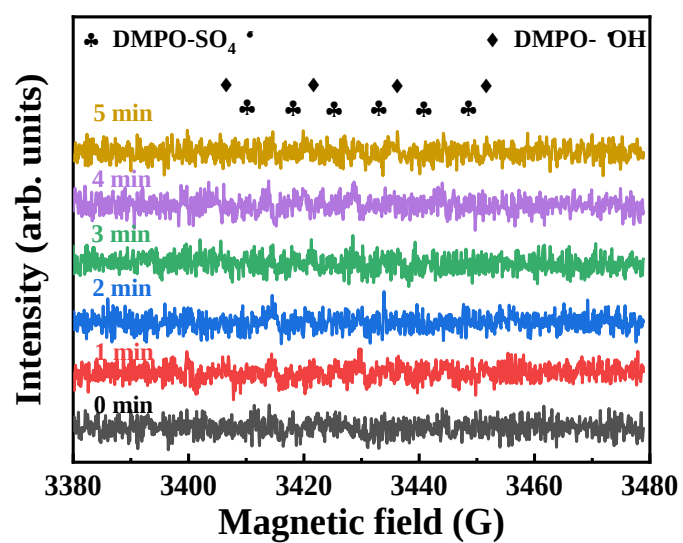
Supplementary Fig. 27 | Degradation performance of SMX at different initial concentrations in the **a**, Fe-SAC-41.31/PMS system **b**, Fe-SAC-T/PMS system, respectively. Experimental conditions: $[PMS]_0 = 2 \text{ mM}$, $\text{pH} = 7$, catalyst: 0.1 g L^{-1} . The error bars in the figures represent the standard deviations from triplicate tests.



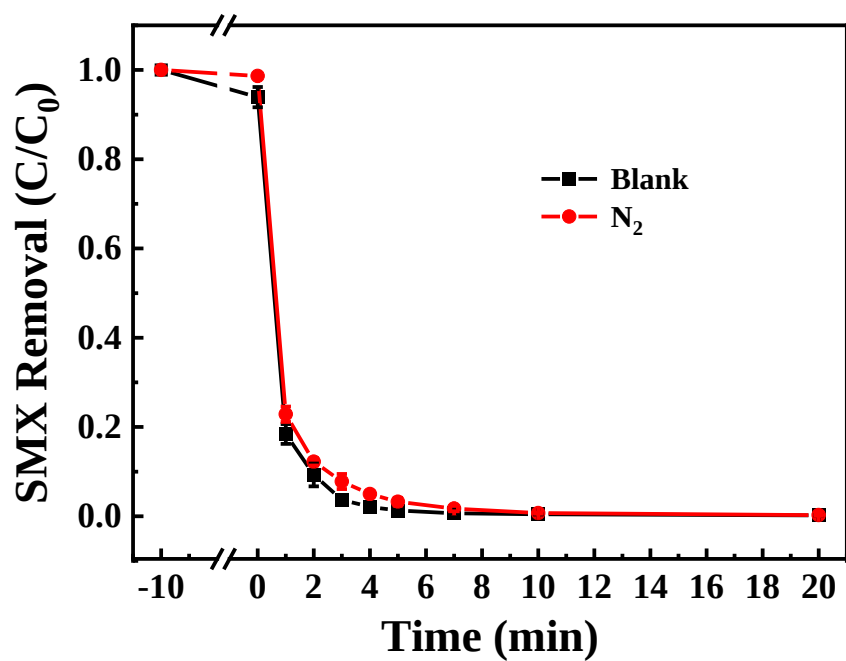
Supplementary Fig. 28 | Effect of ionic strength on SMX removal in Fe-SAC-41.31/PMS system. The error bars in the figures represent the standard deviations from triplicate tests.



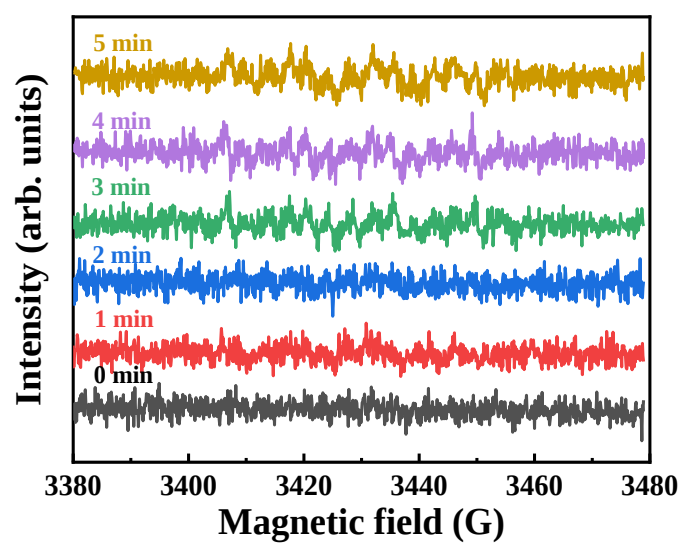
Supplementary Fig. 29 | Open-circuit potential change of Fe-SAC-41.31 with the addition of SMX.



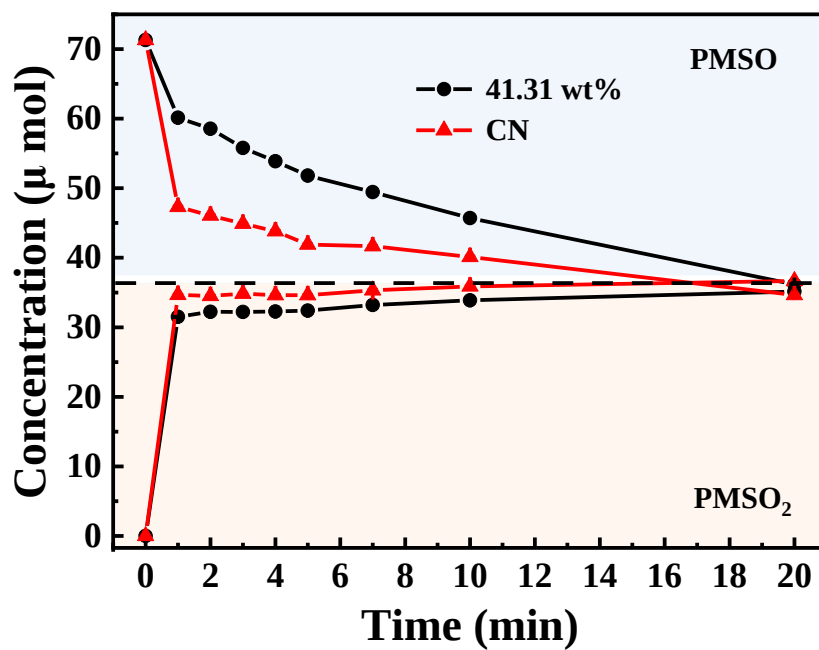
Supplementary Fig. 30 | EPR spectra of $\cdot\text{OH}$ or $\text{SO}_4^{\cdot-}$ in the Fe-SAC-41.31/PMS system.



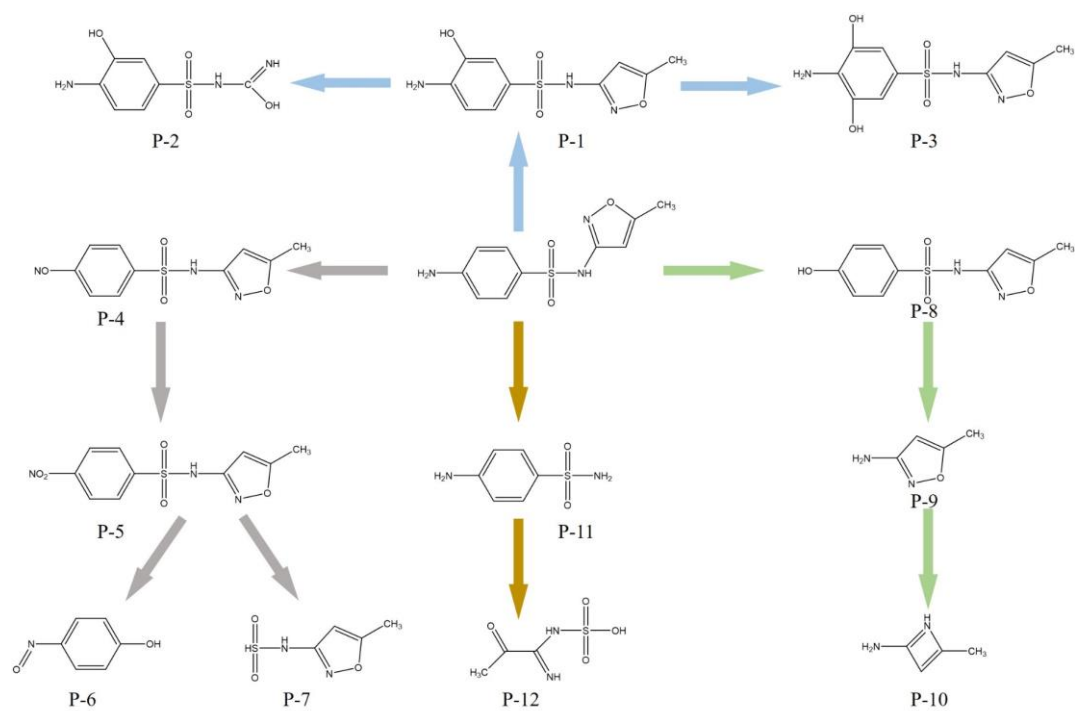
Supplementary Fig. 31 | Degradation performance of SMX with continuous injection of N₂ into the Fe-SAC-41.31/PMS system at a rate of 40 mL min⁻¹. The error bars in the figures represent the standard deviations from triplicate tests.



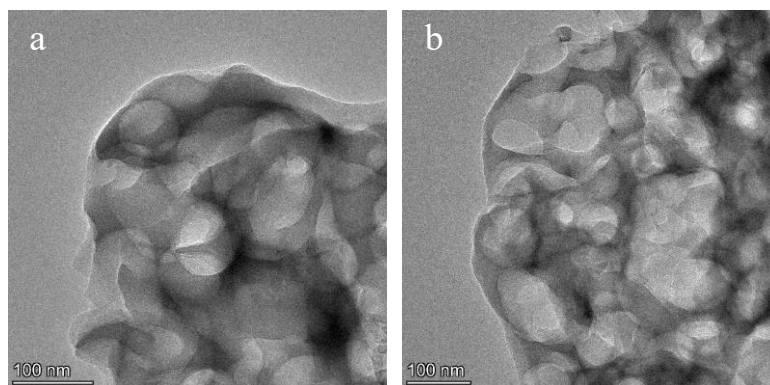
Supplementary Fig. 32 | EPR spectra of $O_2^{\bullet -}$ in the Fe-SAC-41.31/PMS system.



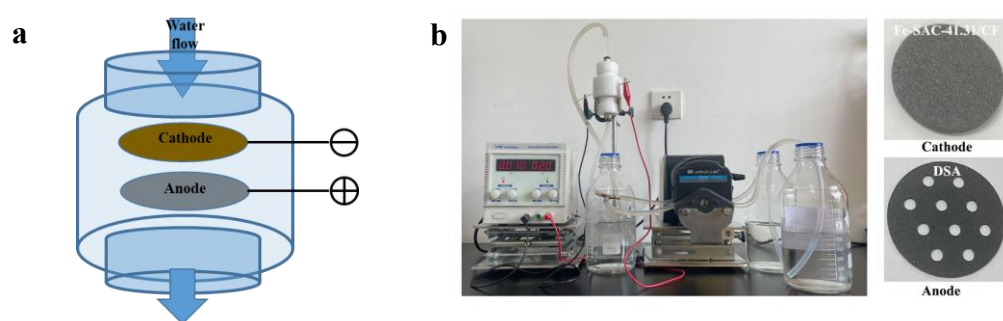
Supplementary Fig. 33 | Concentration changes of PMSO and PMSO₂ over reaction time. Reaction conditions: [PMSO]₀ = 10 mg L⁻¹, [PMS] = 2.0 mM. The error bars in the figures represent the standard deviations from triplicate tests.



Supplementary Fig. 34 | SMX degradation pathways in the Fe-SAC-41.31/PMS system.



Supplementary Fig. 35 | a, b, TEM image of Fe-SAC-41.31 before and after the catalytic reaction.



Supplementary Fig.36 | a, Design principle of continuous flow device; **b,** Physical diagram of continuous flow device.

Supplementary Table 1 | Relationship between metal salts content, SAC precursor yield, and metal loading of as prepared SAC.

Sample	Metal salts (mmol)	SAC precursor yield (g)	Metal contents (wt%)
Fe-SAC-5.16	0.3	1.74	5.16
Fe-SAC-15.82	0.75	1.67	15.82
Fe-SAC-24.62	0.9	1.62	24.62
Fe-SAC-41.31	1	1.53	41.31
Fe-SAC (450 °C)	1	1.55	15.31
Mn-SAC	1	1.55	35.13
Ag-SAC	1	1.51	27.04
La-SAC	1	1.76	22.62
CN	0	3.78	0
Fe-SAC-T	1	3.53	1.61

Supplementary Table 2 | BET surface area, pore diameter and mesoporous pore volume of as-prepared catalysts.

Sample	BET surface (m ² g ⁻¹)	Pore diameter (nm)	Mesoporous pore volume (cm ³ g ⁻¹)
C ₃ N ₄ (MA)	18.74	18.55	0.122
CN(MA-OA)	100.05	21.89	0.464
Fe-SAC 5.16 wt%	84.06	27.29	0.344
Fe-SAC 15.82 wt%	57.55	25.36	0.321
Fe-SAC 24.62 wt%	55.44	20.91	0.181
Fe-SAC 41.31 wt%	46.10	24.06	0.176

Supplementary Table 3 | EXAFS fitting parameters for various Fe-SAC-x samples.

Sample	Shell	N^a	$R(\text{\AA})^b$	$\sigma^2(\text{\AA}^2)^c$	$\Delta E_0(\text{eV})^d$	R factor
FePC	Fe-N	4	1.96	0.0068	1.9	0.0124
Fe-SAC-5.16	Fe-N	3.7	2.09	0.0132	4.91	0.0226
Fe-SAC-15.82	Fe-N	4.4	2.04	0.0171	3.4	0.0188
Fe-SAC-24.62	Fe-N	4.2	2.04	0.0195	5.15	0.0199
Fe-SAC-41.31	Fe-N	4.1	2.04	0.0196	4.44	0.0151

N^a : coordination numbers; $R(\text{\AA})^b$: bond distance; $\sigma^2(\text{\AA}^2)^c$: Debye-Waller factors; $\Delta E_0(\text{eV})^d$: the inner potential correction. R factor: goodness of fit. S_0^2 was set to 0.848, according to the experimental EXAFS fit of Fe foil by fixing CN as the known crystallographic value.

Supplementary Table 4 | Relationship between OA content, SAC precursor yield, and metal loading of as prepared SAC.

Number	OA (mmol)	SAC precursor yield (g)	SAC	Metal loads (wt%)
1	1.5	/	/	/
2	3.1	0.5305	No	/
3	4.2	0.8135	No	/
4	5.3	1.1324	Yes	39.8
5	6.5	1.4053	Yes	35.4
6	7.6	1.6483	Yes	31.2
7	15.1	2.0225	Yes	21.4
8	30.2	2.2318	Yes	17.2

Supplementary Table 5 | Element contents and Fe density derived from XPS results.

Sample	Contents (at %)			
	C	N	O _(total)	Fe
NC	39.21	58.06	2.72	0
Fe-SAC 5.16 wt%	40.54	53.29	4.57	1.6
Fe-SAC 15.82 wt%	39.7	45.99	10.91	3.4
Fe-SAC 24.62 wt%	44.83	31.4	19.22	4.55
Fe-SAC 41.31 wt%	38.67	39.18	16.72	5.44

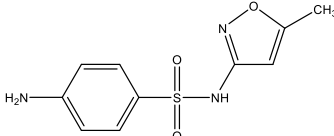
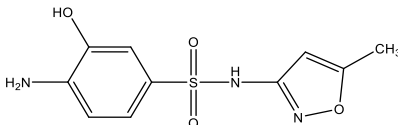
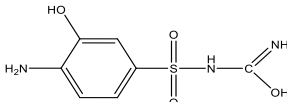
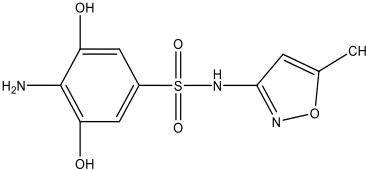
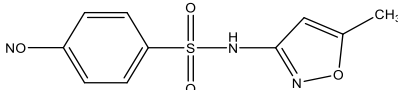
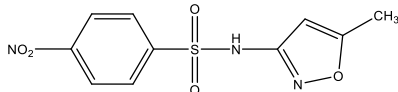
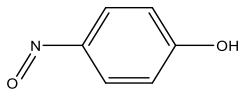
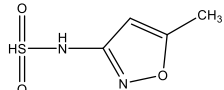
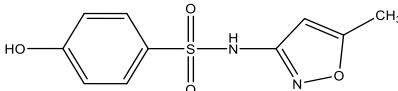
Supplementary Table 6 | Fe coordination bond lengths and bader charges under different loadings.

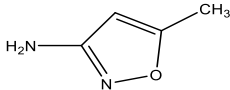
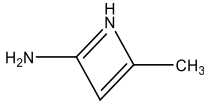
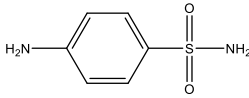
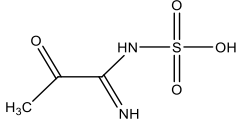
Sample	Fe-C (Å)	Fe-N (Å)	Bader (Fe)
Fe-SAC 5.16 wt%	1.623	2.024	7.135
Fe-SAC 15.82 wt%	1.632	2.033	7.151
Fe-SAC 24.62 wt%	1.640	2.054	7.160
Fe-SAC 41.31 wt%	1.657	2.087	7.162

Supplementary Table 7 | Catalytic oxidation performances of various catalysts for pollutants.

Catalyst (g L ⁻¹)	Pollutant (mg L ⁻¹)	ROS	k value min ⁻¹	Reference
High density Fe-SAC (0.1)	SMX (10)	ETP and ¹ O ₂	1.06	This work
Cu-Fe ^{vac} -LDH (0.1)	SMX (10)	Fe(IV)=O/Fe(V)=O	0.13	6
Fe@C-800 (0.4)	SMX (10)	SO ₄ ^{•-} , [•] OH, O ₂ ^{•-} and ¹ O ₂	0.074	7
Fe-NC-4 (0.15)	SMX (5.0)	¹ O ₂ and Fe(IV)=O	0.815	8
CoSiO _x (0.4)	SMX (7.6)	SO ₄ ^{•-} and [•] OH	0.47	9
Fe-Co-O-g-C ₃ N ₄ (0.2)	SMX (10)	SO ₄ ^{•-} and ¹ O ₂	0.085	10
High density Fe-SAC (0.1)	BPA (10)	ETP and ¹ O ₂	1.269	This work
Co-NC ₄ (0.15)	BPA (10)	SO ₄ ^{•-} , [•] OH and ¹ O ₂	0.69	11
Fe-SAC (0.2)	BPA (25)	¹ O ₂	0.104	12
Co -SAC (0.2)	BPA (25)	¹ O ₂	0.083	12
CuO ₄ N ₁ (0.6)	BPA (20)	SO ₄ ^{•-} , [•] OH, ETP and ¹ O ₂	~0.01	13
CuCoNi-NF	BPA (10)	ETP	0.124	14
High density Fe-SAC (0.1)	BP (10)	ETP and ¹ O ₂	1.084	This work
ZnFe-LDH (0.2)	BP (10)	Fe(IV)=O	0.1	15
CuO ₄ N ₁ (0.6)	BP (20)	SO ₄ ^{•-} , [•] OH, ETP and ¹ O ₂	~0.025	13
FeN _x -C (0.2)	BP (10)	SO ₄ ^{•-} , ¹ O ₂ and Fe(V)=O	0.013	16
High density Fe-SAC (0.1)	TC (20)	ETP and ¹ O ₂	1.575	This work
xCCH/CN-Vn (0.04)	TC (20)	Co(IV)=O and ¹ O ₂	0.25	17
Co-N-GC (0.06)	TC (50)	SO ₄ ^{•-} , [•] OH and ¹ O ₂	0.04	18
Cu-In ₂ O ₃ /O _v (0.5)	TC (20)	SO ₄ ^{•-} and [•] OH	0.26	19
CuO ₄ N ₁ (0.6)	TC (20)	SO ₄ ^{•-} , [•] OH, ETP and ¹ O ₂	~0.3	13
High density Fe-SAC (0.1)	RhB (10)	ETP and ¹ O ₂	0.987	This work
g-C ₃ N ₄ @NPC (0.08)	RhB (10)	SO ₄ ^{•-} , [•] OH, O ₂ ^{•-} and ¹ O ₂	0.152	20
High density Fe-SAC (0.1)	MO (10)	ETP and ¹ O ₂	6.11	This work
biochar	MO (10)	SO ₄ ^{•-} and [•] OH	0.127	21

Supplementary Table 8 | Intermediates detected by LC-MS in the Fe-SAC-41.31/PMS system.

Product	m/z	Molecular formula	Proposed structure
SMX	254.05938	C ₁₀ H ₁₁ N ₃ O ₃ S	
P1	270.05430	C ₁₀ H ₁₁ N ₃ O ₄ S	
P2	232.03865	C ₇ H ₉ O ₄ N ₃ S	
P3	286.04921	C ₁₀ H ₁₁ O ₅ N ₃ S	
P4	268.03865	C ₁₀ H ₉ O ₄ N ₃ S	
P5	284.03356	C ₁₀ H ₉ O ₅ N ₃ S	
P6	124.03930	C ₆ H ₅ NO ₂	
P7	163.01718	C ₄ H ₆ O ₃ N ₂ S	
P8	255.04340	C ₁₀ H ₁₀ O ₄ N ₂ S	

P9	99.05528	$\text{C}_4\text{H}_6\text{ON}_2$	
P10	83.06037	$\text{C}_4\text{H}_6\text{N}_2$	
P11	173.03792	$\text{C}_6\text{H}_8\text{N}_2\text{O}_2\text{S}$	
P12	167.01210	$\text{C}_3\text{H}_6\text{N}_2\text{O}_4\text{S}$	

Supplementary Table 9 | The quality parameters of the wastewater.

	Tap water	Medical wastewater	Mati Lake
pH	7.5	7.53	8.02
Conductivity ($\mu\text{S cm}^{-1}$)	330	21.70	1193
COD (mg L^{-1})	3.0	145.0	32.6
TOC (mg L^{-1})	1.56	102.5	31.7
Cl^{-} (mg L^{-1})	2.11	502.1	420.6
HCO_3^{-} (mg L^{-1})	108	48.3	83.1
NO_3^{-} (mg L^{-1})	0.3	18.3	50.8

Supplementary Table 10 | The quality parameters of the coal reverse osmosis water.

	Detection value
pH	9.41
Conductivity ($\mu\text{S cm}^{-1}$)	18400
TOC (mg L^{-1})	69.725
TDS (mg L^{-1})	9200
TN (mg L^{-1})	194.75
Suspension (mg L^{-1})	0.045
Saturation	D=30, buff
UV254 (cm^{-1})	0.675
Cl ⁻ (mg L^{-1})	4108.65
SO ₄ ²⁻ (mg L^{-1})	604.223
NO ₃ ⁻ (mg L^{-1})	789.543
NO ₂ ⁻ (mg L^{-1})	2.828
NH ₄ ⁺	8.31

Supplementary Table 11 | 3DEEM response and their associated characteristics.

Region	I	II	III	IV
Excitation Wavelength	200-250 nm	200-250 nm	250-300 nm	250-500 nm
Emission Wavelength	280-380 nm	380-600 nm	280-380 nm	380-500 nm
Characteristics	Fluorophores associated with aromatic-like molecules (Tyrosine)	Fluorophores associated with fulvic acid-like molecules	Fluorophores associated with SMP-like molecules (Tryptophane)	Fluorophores associated with humic acid-like molecules

Supplementary Table 12 | The HPLC analysis conditions for different substrates.

Substrates	Flow	λ	CH ₃ OH	Acetonitrile	H ₂ O
	(ml min ⁻¹)	(nm)	(%)	(%)	(%)
BPA	0.5	280	70	/	30
TC	0.5	275	/	18	82(0.1% H ₃ PO ₄)
SMX	0.5	265	35	/	65
SMT	0.5	280	35	/	65

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