## Electronic Supplementary Information

Table S1. Physical and chemical properties of oxytenacyclin	Table	S1.	Physical	and	chemical	properties	ofox	ytetracyclin
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Antibiotic	Molecular	Chemical structure	Molecular	$\lambda_{max}$	Water	pKa	Existing
	formular		weight	(nm)	solubility		
			(g/mol)		(mg/L)		
Oxytetracycline	C <sub>22</sub> H <sub>24</sub> N <sub>2</sub> O <sub>9</sub>	H <sub>3</sub> C、,CH <sub>3</sub> H0 CH-OH N	460.4	276 and	200	pKa <sub>1</sub> =3.37	OTC <sup>+</sup> , OTC <sup>zwitterion</sup>
(OTC)		ОН		354 nm		pKa <sub>2</sub> =7.49	OTC <sup>-</sup> , OTC <sup>2-</sup>
		OH O OH O O				pKa <sub>2</sub> =9.88	

Catalyst	BET surface area (m²/g) <sup>(a)</sup>	Pore volume (cm <sup>3</sup> /g) <sup>(b)</sup>	Pore size (nm) <sup>(c)</sup>
Natural clinoptilolite	36.21	0.097	12.18
Activated	75 76	0 105	11.62
clinoptilolite	75.70	0.105	11.02
g-C <sub>3</sub> N <sub>4</sub>	34.41	0.127	15.83
Bi <sub>2</sub> MoO <sub>6</sub>	37.14	0.169	16.21
CNBC30	48.55	0.130	11.55

Table S2. The pore and surface characterization of different samples

<sup>(a)</sup> The specific surface area was calculated by BET method

<sup>(b)</sup> The pore volume was obtained from the BJH desorption cumulative volume of pores between 1.70 nm and 300.00 nm width

<sup>(c)</sup> The BJH Desorption average pore width

Catalyst and fabricated	Organic	Reaction conditions	Results	Remarks	References
method	pollutants				
MoS <sub>2</sub> /TiO <sub>2</sub> /clinoptilolite	Sodium isopropyl	[catalyst] = 0.4  g/L,	SIPX: 97.3 % (3 h)	- Complicated synthesis process	1
(MTC3 – 1mmol	xanthate (SIPX)	[SIPX] = 10 mg/L,	Main ROS: $h^+ < e^- <$	< - Long degradation duration	
Na <sub>2</sub> MoO <sub>4</sub> .2H <sub>2</sub> O)	$(\lambda_{max} = 301 \text{ nm})$	400W Xe lamp	$OH^{\bullet} < O_2^{\bullet-}$		
2-step hydrothermal					
TiO <sub>2</sub> /NCP	Atenolol	[catalyst] = 3 g/L,	Atenolol: 75 % (1h)	- High energy requirement due to	2
(96.6 % NCP)	$(\lambda_{max} = 284 \text{ nm})$	$[\text{atenolol}] = 10 \text{ mg/L}^{-1},$	Main ROS: h <sup>+</sup> and	calcination at high temperature	
Heating		UV lamp 60 W,	OH•	- Low degradation efficiency	
			Mineralization	- High cost due to UV light	
			efficiency: 74 %	- Great amount of catalyst	
			Final products: CO <sub>2</sub>	- Degradation efficiency decreased	
			and $H_2O$	at pH > 6.0	
ZnO/Fe <sub>2</sub> O <sub>3</sub> /Clinoptilolite	Metronidazole	[catalyst] = 1 g/L,	MNZ: 99 %	- Narrow pH range, good	3
(44 % Clinoptilolite,	(MNZ)	[MNZ] = 60  mg/L,	(90 min)	performance at base medium	
$Fe^{3+}/ZnO = 0.06)$	$(\lambda_{max} = 254 \text{ nm})$	$[H_2O_2] = 40 \text{ mg/L},$		- High cost due to UV light	
Sol-Gel		pH = 10		- Great amount of catalyst	
		UV lamp 8 W,			
		reaction time = 90 min			

Table S3. Comparison of the degradation of organic pollutants by various photocatalysts supported on clinoptilolite

Catalyst and fabricated	Organic	Reaction conditions	Results	Remarks	References
method	pollutants				
TiO <sub>2</sub> /Fe <sub>2</sub> O <sub>3</sub> /Clinoptilolite	Diphenhydramine	[catalyst] = 0.5 g/L,	80 % DPH	- High energy requirement due to	4
$(\text{Fe}^{3+/}\text{TiO}_2 = 0.6, 25\%)$	(DPH)	рН=5,	(120 min)	calcination at high temperature	
Clinoptilolite)	$(\lambda_{max} = 258 \text{ nm})$	[DPH] = 50  mg/L,		- Narrow pH range, good	
Hydrothermal		$[H_2O_2] = 50 \text{ mg/L},$		performance at acid medium	
		UV lamp 6 W		- High cost due to UV light	
ZnO/Fe <sub>2</sub> O <sub>3</sub> /Clinoptilolite	Diphenhydramine	[catalyst] = 0.5 g/L,	95 % DPH	- Narrow pH range, good	5
$(Fe^{3+/}ZnO = 0.6, 25\%)$	(DPH)	pH=10,	(100 min)	performance at base medium	
Clinoptilolite)	$(\lambda_{max} = 258 \text{ nm})$	[DPH] = 50  mg/L,		- High cost due to UV light	
Sol-Gel		$[H_2O_2] = 50 \text{ mg/L},$			
		UV lamp 6 W			
BiOCl/TiO <sub>2</sub> /clinoptilolite	Sodium isopropyl	[catalyst] = 0.2 g/L,	SIPX: > 90 % (3 h)	- Complicated synthesis process	6
$(TiO_2/clinoptilolite = 1.5,$	xanthate (SIPX)	[SIPX] = 20 mg/L,	Main ROS: e <sup>-</sup> <	- Long degradation duration	
BTC1 - 0.25 mmol Bi <sup>3+</sup> )	$(\lambda_{max} = 301 \text{ nm})$	400W Xe lamp	$OH^{\bullet} < h^+ < O_2^{\bullet-}$		
Hydrothermal + water bath					
precipitation					
TiO <sub>2</sub> /CLP/graphene	Nitenpyram	[Nitenpyram] = 80 mg/L,	100 % Nitenpyram	- Complicated synthesis process	7
(30% clinoptilolite, 1 %	$(\lambda_{max} < 400 \text{ nm})$	570 W Xe lamp	(90 min)	- Long degradation duration	
graphene)			Mineralization		
Hydrothermal + Solvothermal			efficiency: 71%		

Catalyst and fabricated	Organic	Reaction conditions	Results	Remarks	References
method	pollutants				
			Main ROS: $h^+ < O_2^{\bullet-}$		
			< OH•		
$g\text{-}C_3N_4/Bi_2MoO_6/clinoptilolite$	Oxytetracycline	[OTC] = 20  mg/L;	87.47 % OTC	- Low dosage of catalysts and	This work
(CNBC-30, 30 % clinoptilolite	) (OTC)	[catalyst] = 500 mg/L;	(120 min)	PDS	
Solvothermal	$(\lambda_{max} = 354 \text{ nm})$	$[Na_2S_2O_8] = 1.26 \text{ mM};$		- Wide range of pH (3-11)	
		pH initial; L4X 40 W		- Environmental friendliness	
		LED lamp		energy (LED light)	
				- High degradation efficiency	

Photocatalyst	Organic pollutants	Reaction conditions	Time	Degradation	Main ROS	References
				efficiency		
TiO <sub>2</sub> /AB	Tetracycline (TC)	[TC] = 30 mg/L	120 min	93.3 %	SO₄•-	8
		[catalyst] = 500 mg/L				
		[PDS] = 3  mM				
$g-C_3N_4$	Bisphenol A (BPA)	[BPA] = 5 mg/L	90 min	99.5 %	$O_2^{\bullet-}, h^+$	9
		[catalyst] = 0.5 g/L				
		[PDS] = 5mM				
CeO2/g-C3N4	Norfloxacin (NOR)	[NOR] = 10  mg/L	60 min	88.6 %	$^{1}\text{O}_{2}, \text{O}_{2}^{\bullet-}, \text{h}^{+} \text{ and }$	10
		[catalyst] = 1 g/L			OH•	
		[PDS] = 5 mM				
Cu/ZnO/CoFe-CLDH	Bisphenol-A (BPA)	[BPA] = 10  mg/L	6 h	99 %	SO4 <sup>•-</sup> , OH <sup>•</sup> , O2 <sup>•−</sup> ,	11
		[catalyst] = 0.5 g/L			$^{1}O_{2}$	
		[PDS] =0.148mM				
Bi <sub>2</sub> MoO <sub>6</sub>	Tetracycline (TC)	[TC] = 20  mg/L	60 min	95.18 %	$SO_4^{\bullet-}, h^+$	12
		[catalyst] = 0.5 g/L				
		[PDS] = 16.8  mM				
ZnFe <sub>2</sub> O <sub>4</sub>	Bisphenol-A (BPA)	[BPA] = 10  mg/L	120min	96.5%	$SO_4^{\bullet-}, OH^{\bullet}, h^+$	9
		[catalyst] = 0.2 g/L				
		[PDS] = 8  mM				

Table S4. The removal of organic pollutants from water by photocatalysts in the presence of persulfate and visible light

Photocatalyst	Organic pollutants	Reaction conditions	Time	Degradation	Main ROS	References
				efficiency		
g-C <sub>3</sub> N <sub>4</sub> /Bi <sub>2</sub> MoO <sub>6</sub> /clinoptilolite	Oxytetracycline	[OTC] = 20  mg/L	120 min	87.47 %	$^{1}\text{O}_{2}, \text{O}_{2}^{\bullet}$ , and $h^{+}$	This work
(CNBC-30)	(OTC)	[catalyst] = 500 mg/L				
		[PDS] = 1.26 mM				

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Fig. S1. Calibration curve of oxytetracycline



Fig. S2 (a) Mott-Schottky of various catalysts, (b) XPS spectra of survey scan



Fig. S3. The k<sub>app</sub> values of (a) different systems, (b) % clinoptilolite, (c) catalyst dosage, (d) PDS concentration, (e) initial pH, and (f) OTC concentration Protection conditions: [OTC] = 10.50 mg/L : [cetalyst] = 0.600 mg/L : [Na S O ] = 0.2.10 mM; pH = 3

(Reaction conditons: [OTC] = 10-50 mg/L; [catalyst] = 0-600 mg/L;  $[Na_2S_2O_8] = 0-2.10 \text{ mM}$ ; pH = 3-11; T=30 °C)



Fig. S4. UV-Vis absorption spectrometry of OTC using CNBC-30 during 120 min (30 min of adsorption and 90 min of photocatalysis)



Fig. S5. (a) Schematic diagram of OTC species distribution according to pH and (b)p $H_{pzc}$  of CNBC-30



Fig. S6. The  $k_{\mbox{\scriptsize app}}$  values of reaction with the presence of different ions





Fig. S7. Mass spectrum of OTC decomposition over time from 0 - 90 minutes