

Electronic Supplementary Material (ESI) for

Self doping and surface plasmon modification induced visible light photocatalysis of BiOCl

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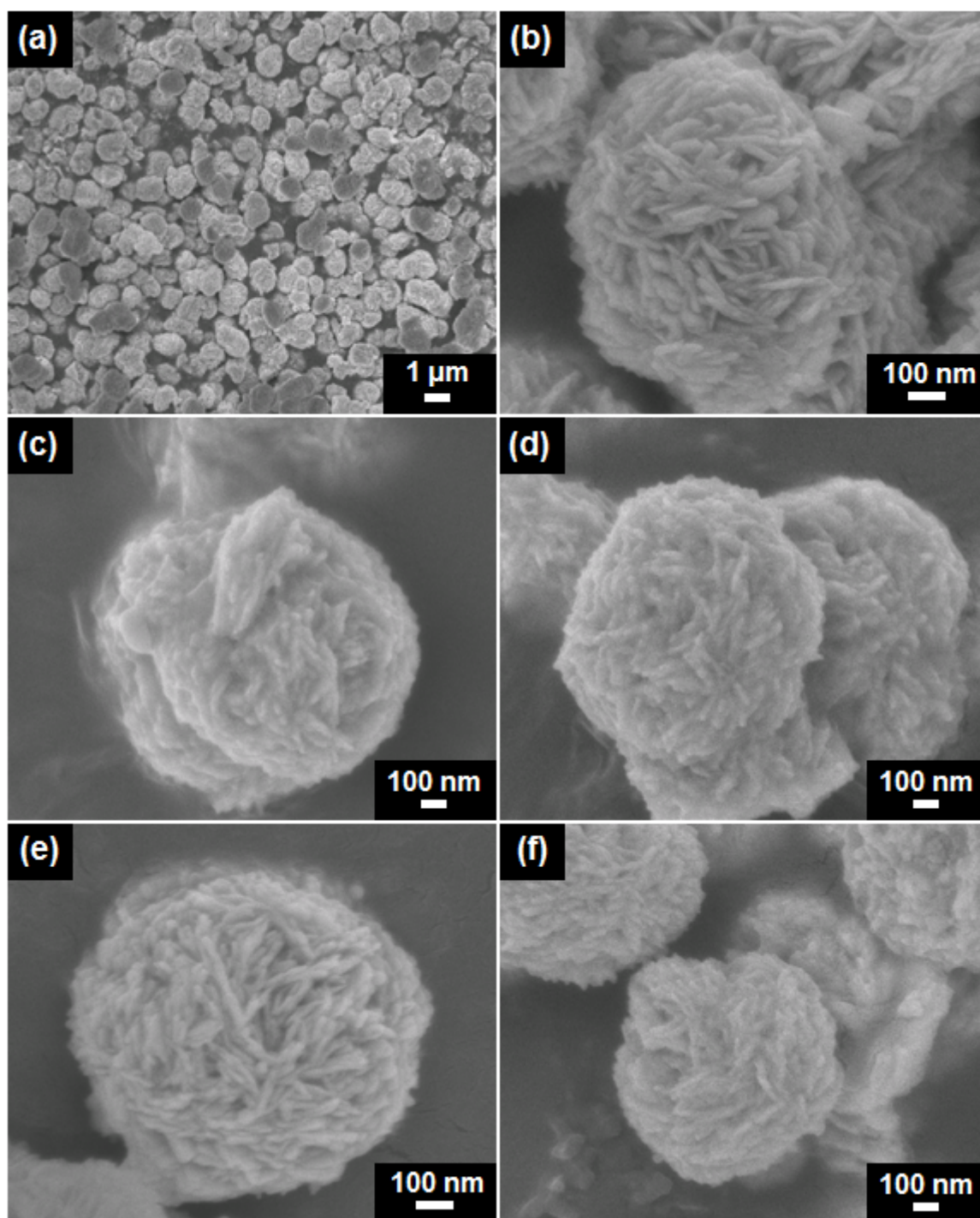


Fig. S1 SEM images of the self-doped BiOCl and Ag/BiOCl: (a, b) AB0, (c) AB1, (d) AB3, (e) AB5 and (f) AB7.

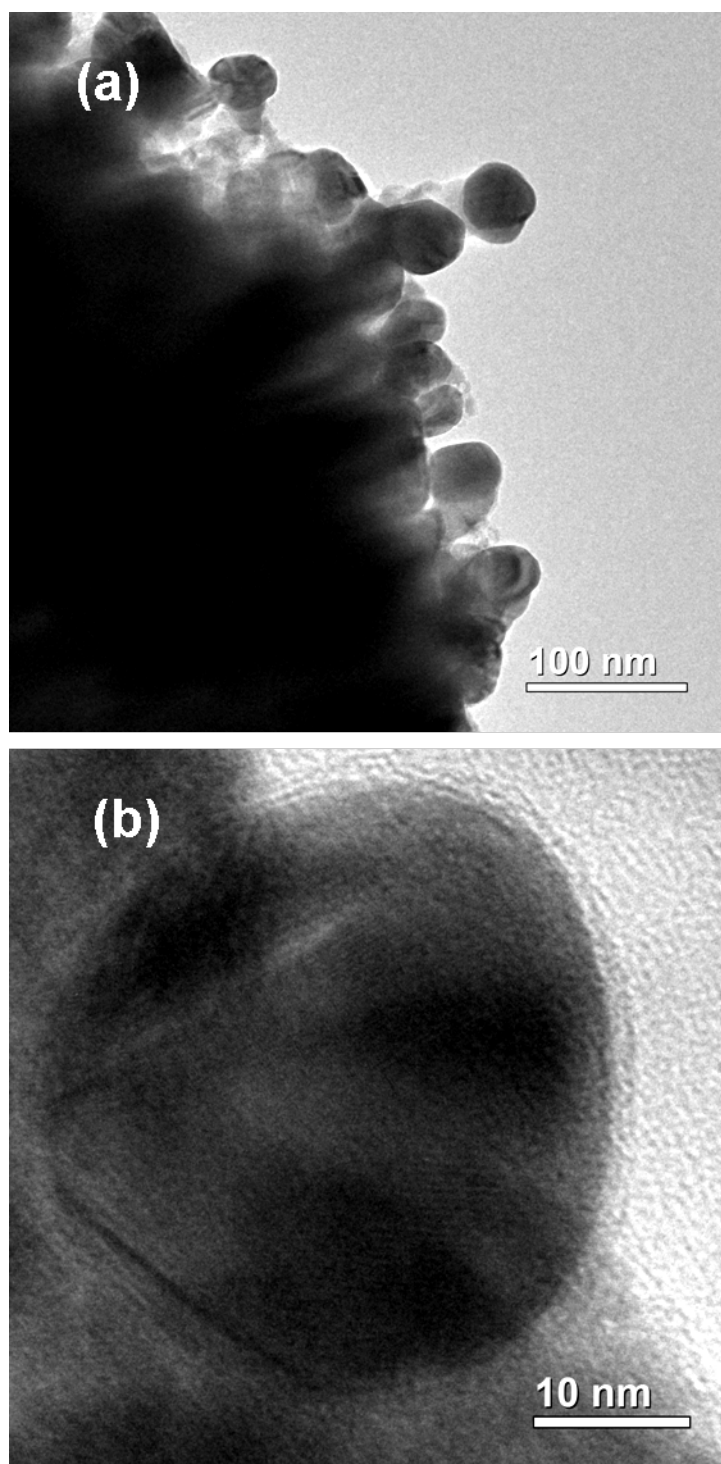
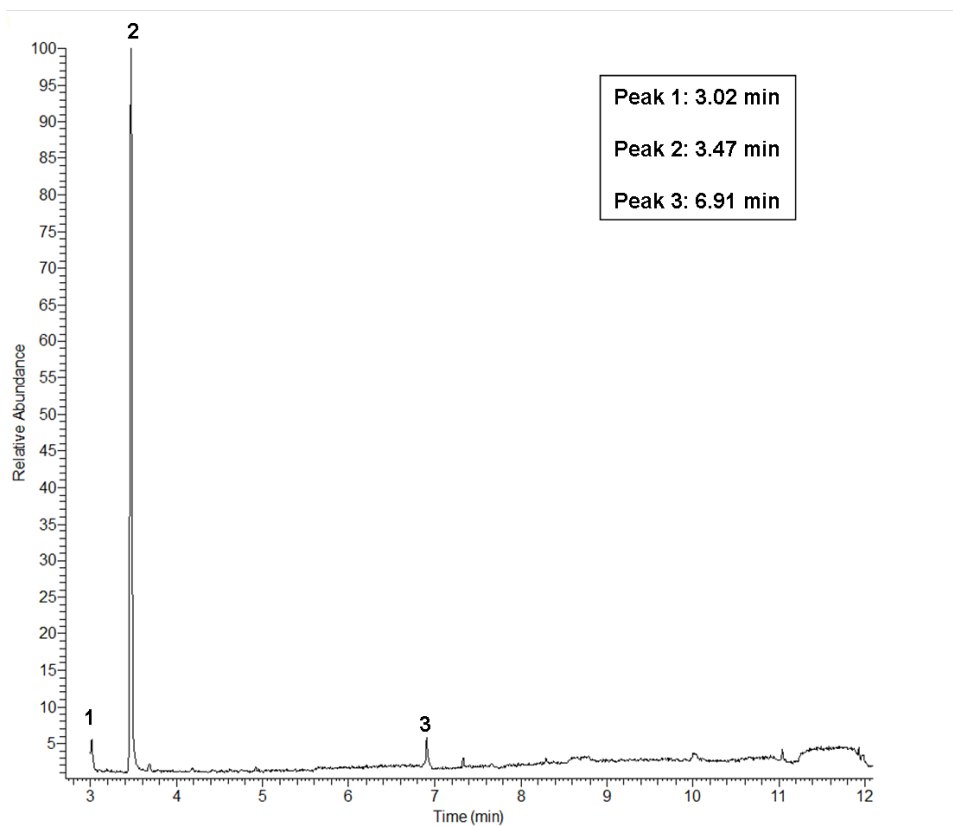
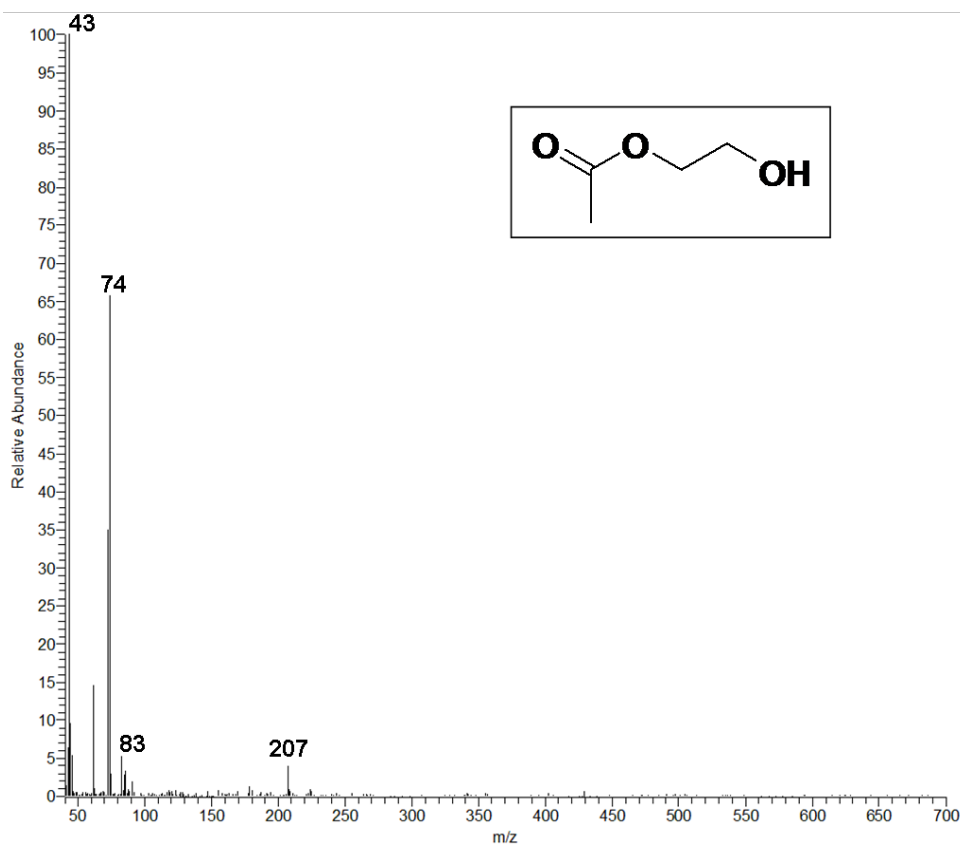


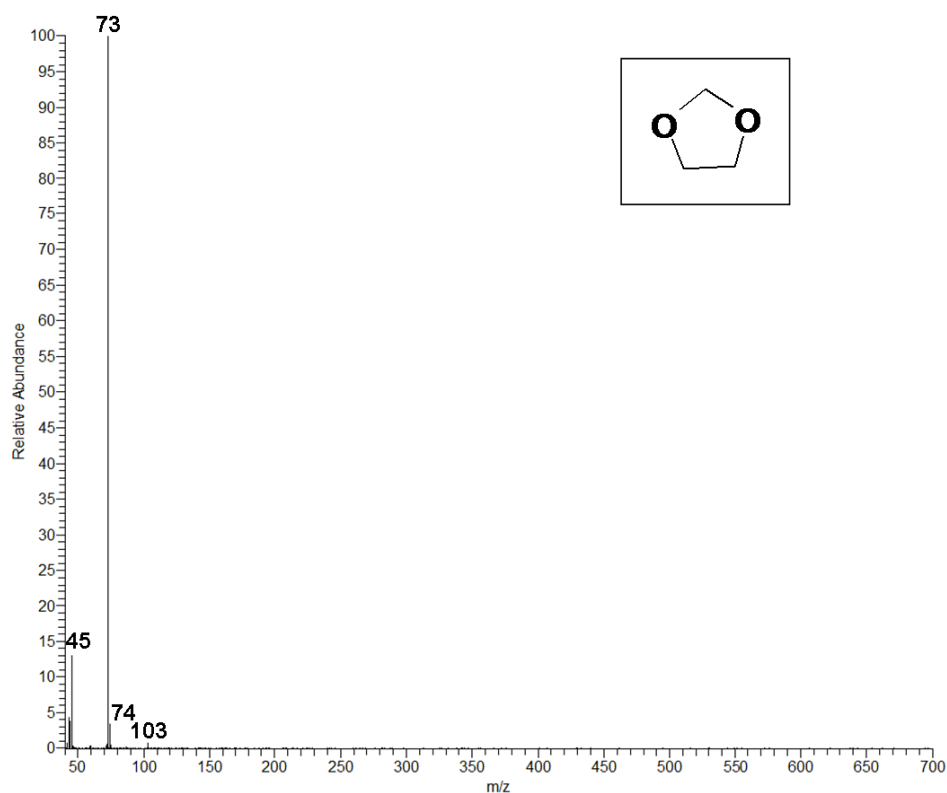
Fig. S2 TEM (a) and HRTEM (b) images of the AB5.



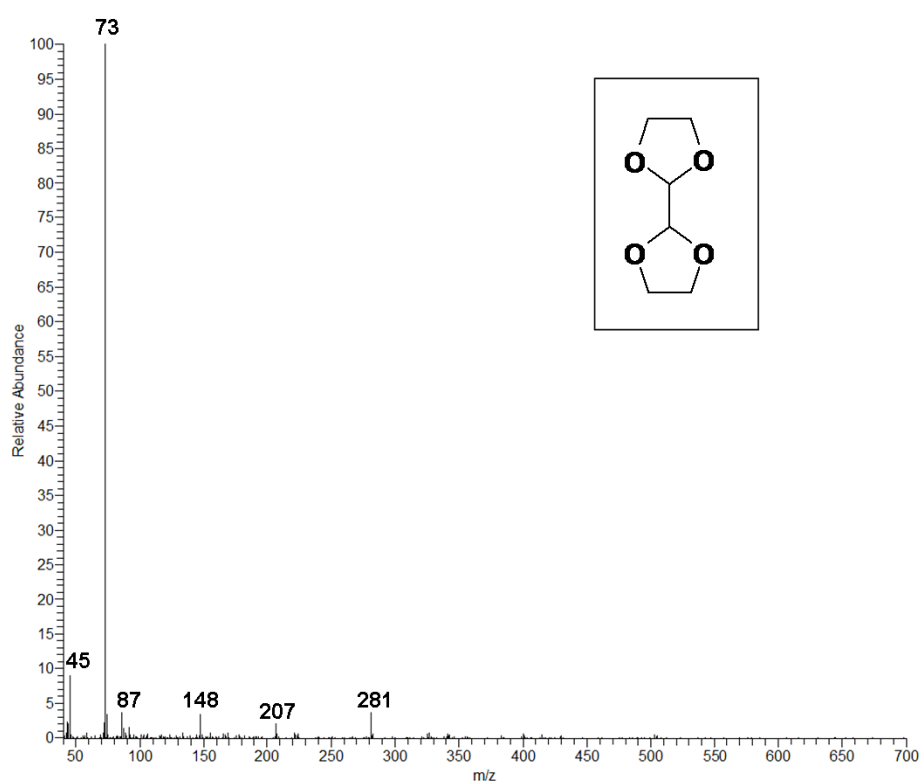
(a)



(b)



(c)



(d)

Fig. S3 (a) GC chromatogram of the final reaction solution, b) Mass spectrum for the product at 3.02 min; c) Mass spectrum for the product at 3.47 min; d) Mass spectrum for the product at 6.91 min.

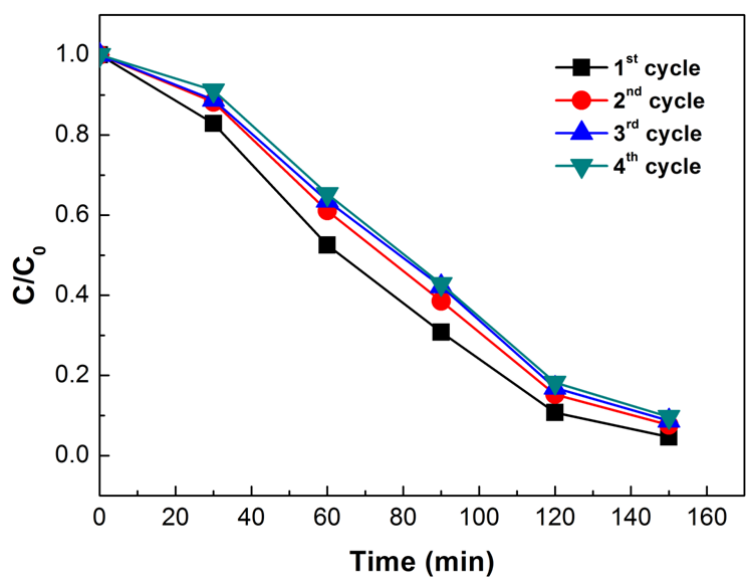


Fig. S4 Recycling test on the AB5 sample for MO photodegradation under visible light irradiation.

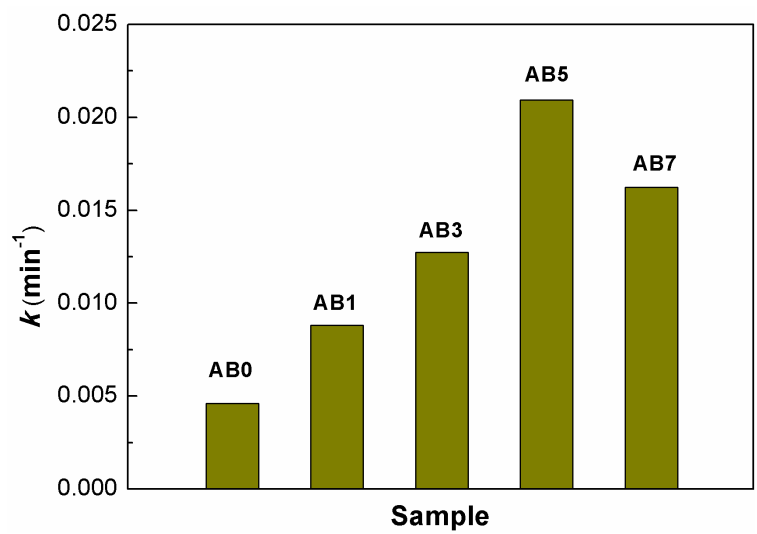


Fig. S5 The apparent reaction rate k (min⁻¹) of the self-doped BiOCl and Ag/BiOCl on the degradation of MO under visible light irradiation ($\lambda > 420$ nm).

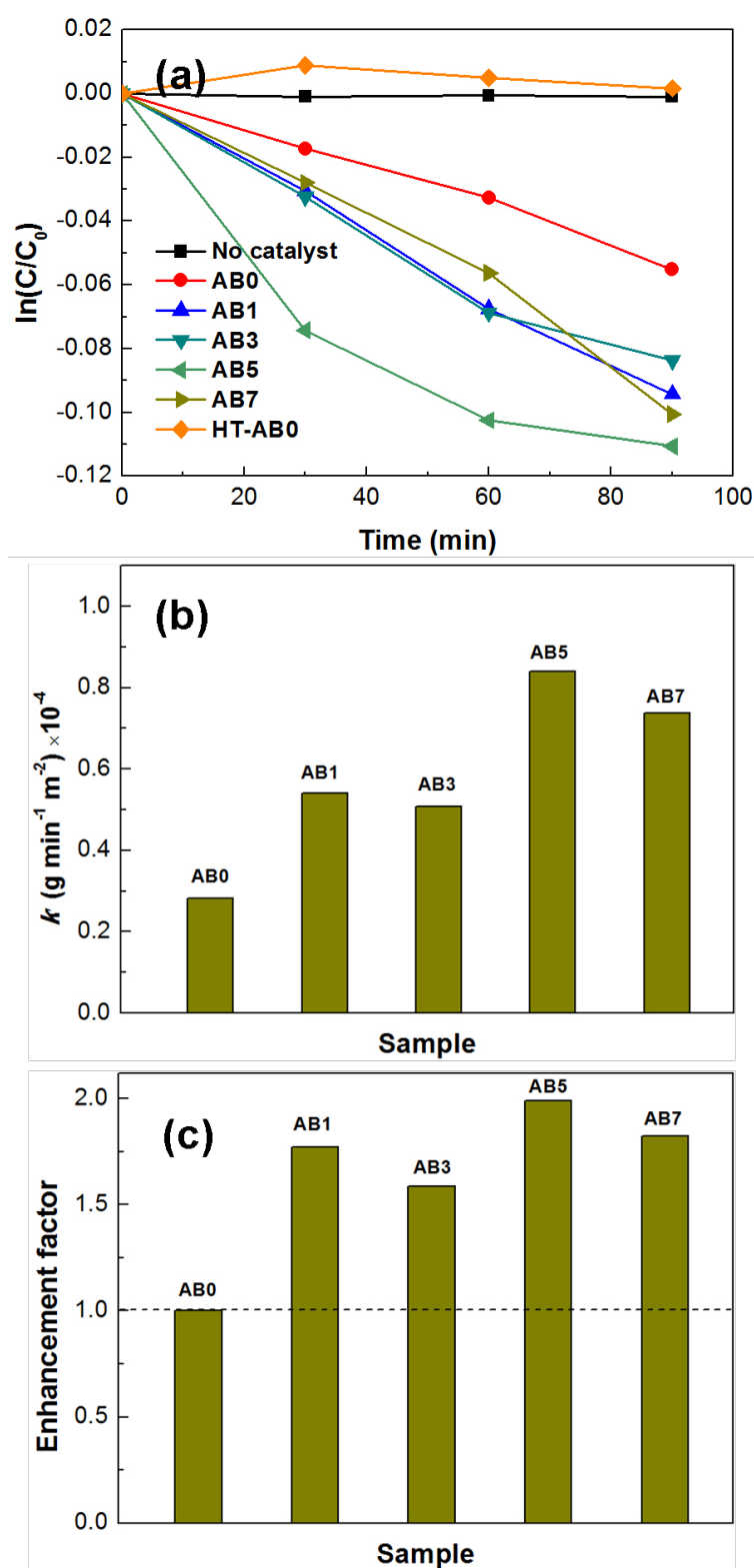


Fig. S6 (a) Photocatalytic activity of the BiOCl and Ag/BiOCl on the degradation of SA (10 mg L⁻¹) under visible light irradiation ($\lambda > 420$ nm). (b) Photoactivity enhancement factor ($k_{\text{Ag/BiOCl}}/k_{\text{BiOCl}}$) of Ag/BiOCl. (c) The apparent reaction rate normalized with the surface areas (k') of the self-doped BiOCl and Ag/BiOCl.

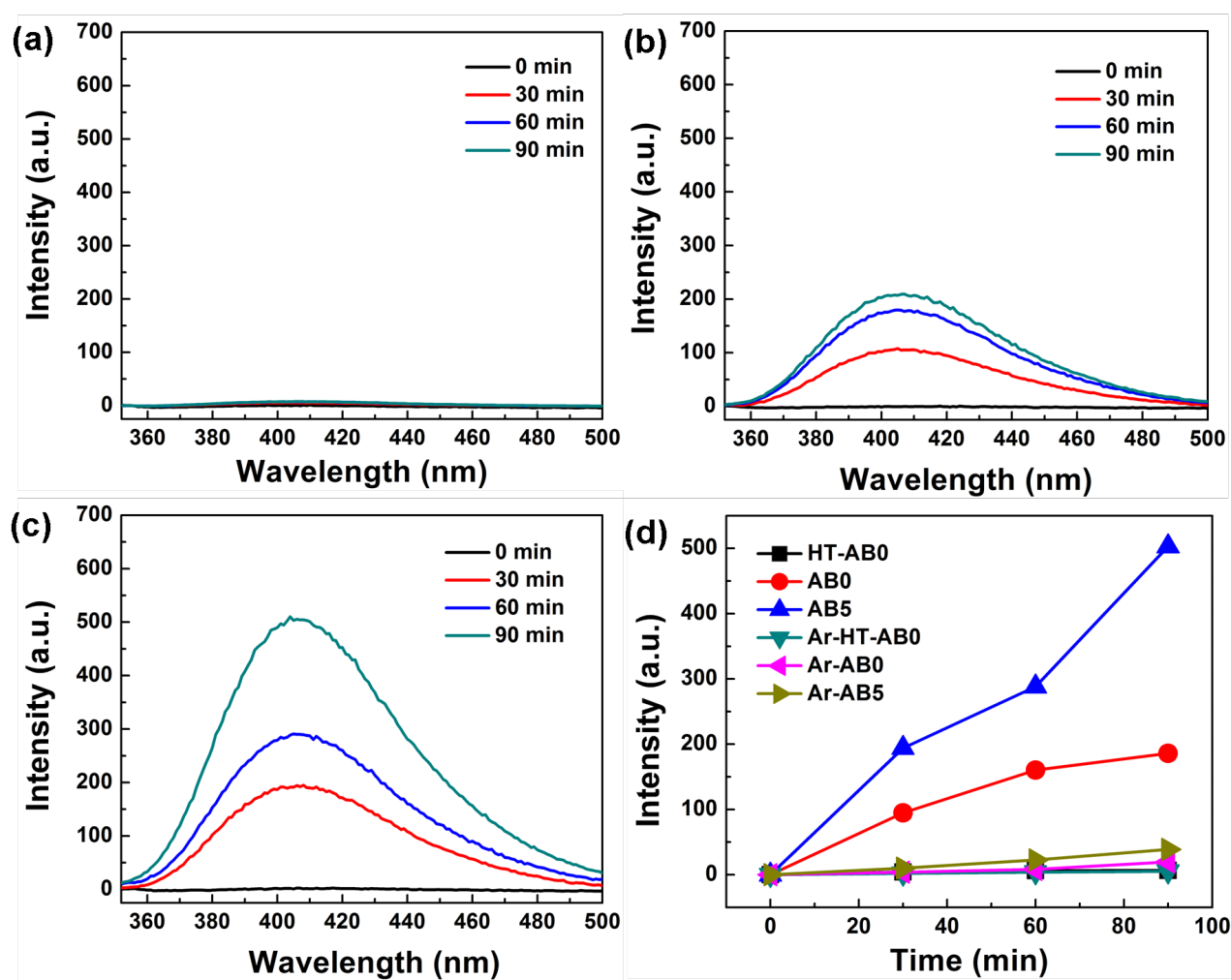


Fig. S7 Detection of H₂O₂ in the HT-AB0 (a), AB0 (b) and AB5 (c) dispersions under visible light irradiation. (d) Comparison of PL intensity for H₂O₂ detection in different catalytic systems.

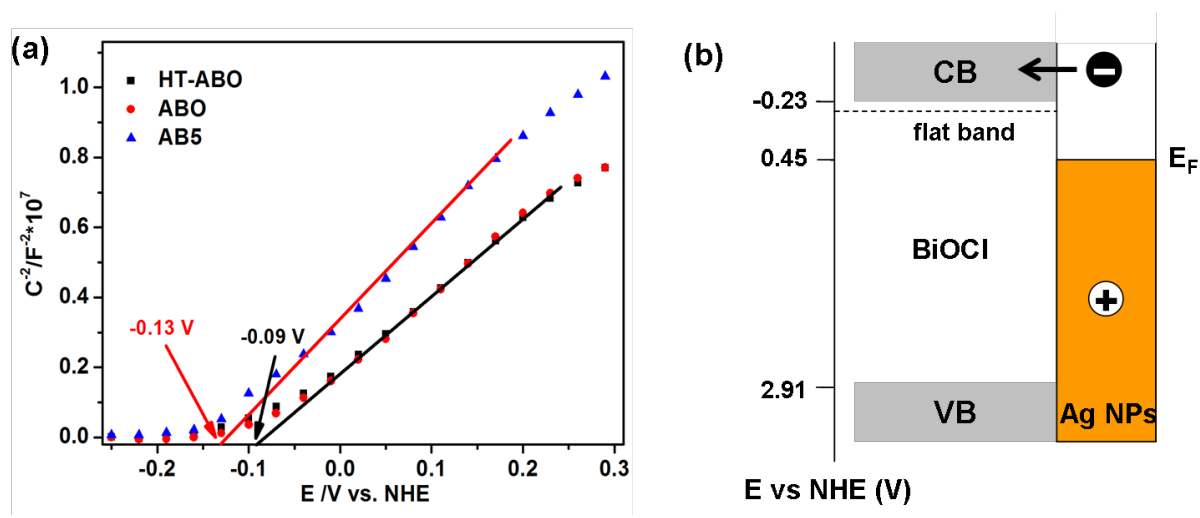


Fig. S8 (a) Mott-Schottky plots of the different sample electrodes and (b) Energy level diagram of the AB5 sample.

Table S1 Textural and photocatalytic properties of the different samples

sample	A_{BET} ($\text{m}^2 \text{g}^{-1}$)	MO degradation in 150 min (%)	k ($\times 10^{-3}, \text{min}^{-1}$)	k' ($\times 10^{-4}, \text{g min}^{-1}$ m^{-2}) ^a
AB0	21.4	51%	4.6	2.1
AB1	19.8	73%	8.8	4.4
AB3	18.9	86%	12.7	6.7
AB5	14.3	96%	20.9	14.7
AB7	14.9	91%	16.2	10.9

^a The k' values were k values normalized with the surface areas.

Table S2 Kinetic analysis of the emission decay of the different samples^[a]

sample	A	B_1	τ_1 (ns)	B_2	τ_2 (ns)	χ^2	$\langle\tau\rangle$ (ns)	k (ns^{-1})
AB0	0.126	0.1247	0.31	1.815×10^{-2}	2.75	6.775	0.63	1.59
AB5	0.144	0.2270	0.17	2.140×10^{-2}	2.40	4.074	0.37	2.70

^[a] The data in each curve was fitted to multiexponential functions ($f(t) = A + \sum_{i=1}^N B_i e^{-t/\tau_i}$), with preexponential factors (B_i), characteristic lifetimes (τ_i) and a background (A). The average lifetime of a multiexponential decay is expressed as $\langle\tau\rangle = \sum_{i=1}^N c_i \tau_i$, where c_i is the relative concentration in a

multiexponential decay ($c_n = \frac{B_n}{\sum_{i=1}^N B_i}$).^{1,2}

Reference:

1. S. Sarkar, A. Makhal, T. Bora, S. Baruah, J. Dutta and S. K. Pal, *Phys. Chem. Chem. Phys.*, 2011, **13**, 12488.
2. J. R. Lakowicz, *Principles of Fluorescence Spectroscopy*, Kluwer Academic/Plenum Publishers, New York, 2nd edn, **1999**.