

Comparative study of BiVO₄ and BiVO₄/Ag₂O on their properties and photocatalytic degradation mechanism

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Supplementary Experimental Section:

2.1.1 Synthesis of BiVO₄

BiVO₄ particles were synthesized by the solvothermal method. Bi(NO₃)₃·5H₂O (1.8 g) was dissolved into the mixture solution of HNO₃ (1.8 mL) and 1,3-PDO (15 mL). NH₄VO₃ (0.4 g) was dissolved into NaOH (2 mol/L, 15 mL) solution. Then, the NH₄VO₃ solution was slowly dropped into Bi(NO₃)₃ solution. The pH of the mixture solution was adjusted to 7. The resulting mixture was transferred into teflon-lined autoclave and heated at 180°C for 12 h. Subsequently, the mixture was then centrifuged, washed with water/methanol solution, and dried at 60°C for 8 h. Finally, the solids were calcined at 450 °C for 1 h to obtain the final BiVO₄ product.

2.2 Photocatalytic experiment

BiVO₄ or BiVO₄/Ag₂O (50 mg) was added to 18 mg/L DC solution (250 mL). The solution was stirred in the darkness for 30 minutes and then exposed to visible light. A certain volume of solution (1 mL) was taken out at the set time and then passed through 0.22 μm filters. The residual DC concentration was detected by ultra-performance liquid chromatography (UPLC) at a wavelength of 345 nm. The solar illumination was simulated by using Xenon lamp (PLS-SXE300 Perfectlight), and visible light was obtained through a cutoff filter. The actual wavelength range was 420 nm to 780 nm. The degradation intermediates of DC were measured by UPLC-QTOF (agilent).

2.3 Characterization of BiVO₄ and BiVO₄@Ag₂O particles

The crystalline phases of BiVO₄ and BiVO₄@Ag₂O particles were analyzed by XRD (D8 Advance, Bruker, Germany). The surface chemical composition and

chemical states of the samples was performed by XPS (Escalab 250Xi ThermoFisher). The morphology and particle sizes of the samples were observed by SEM (Sigma 500, Zeiss, Germany). UV-vis DRS was recorded on a Shimadzu UV-2550 spectrometer. The photogenerated electron-hole formation and separation of solid semiconductor materials were analyzed through the SS-SPS technique. The SS-SPS measurements of the samples were carried out with a home-built apparatus, equipped with a lock-in amplifier (SR830) synchronized with a light chopper (SR540).

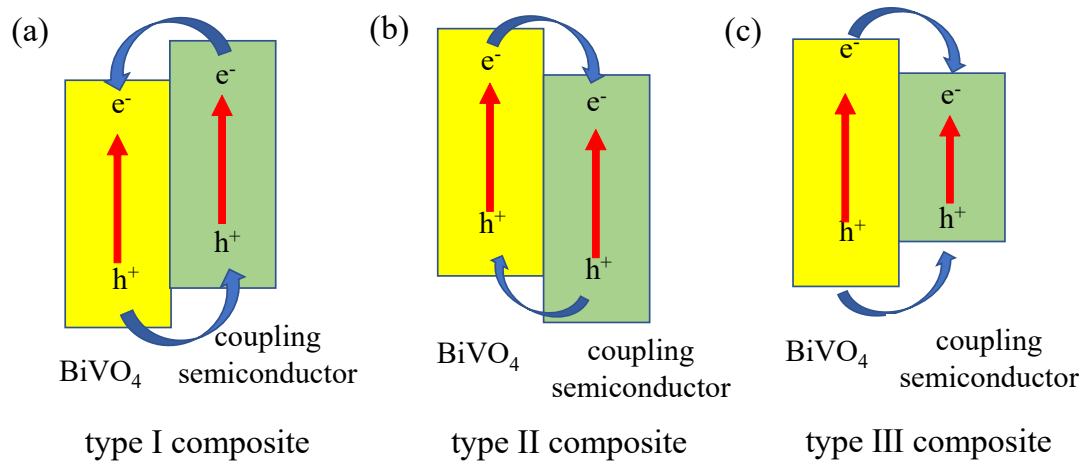
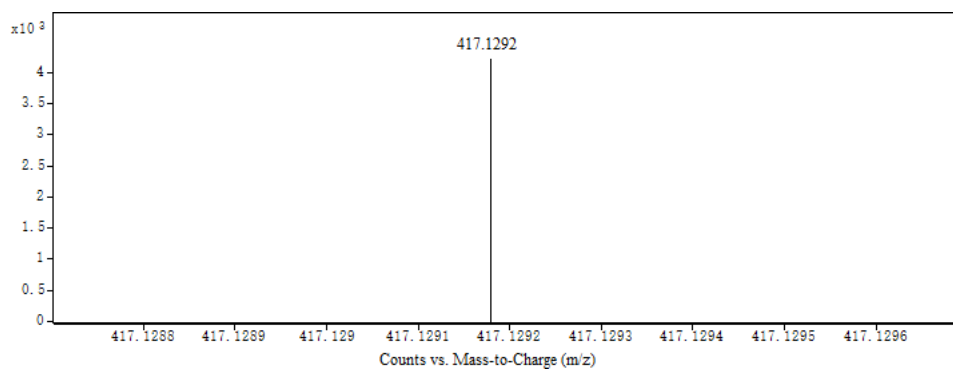
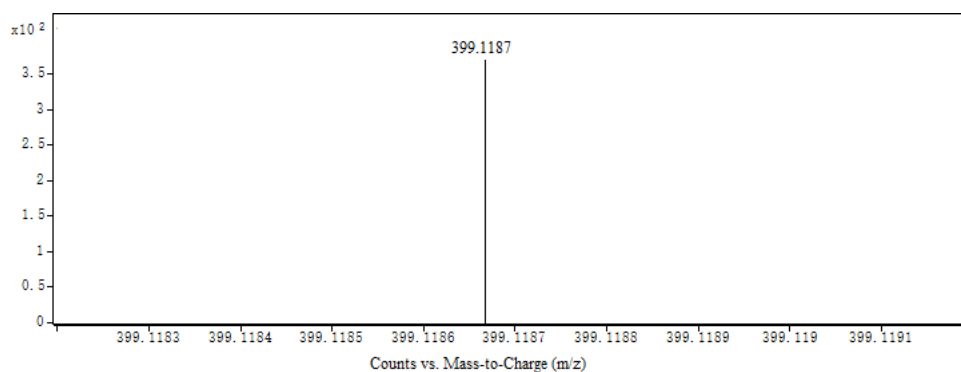
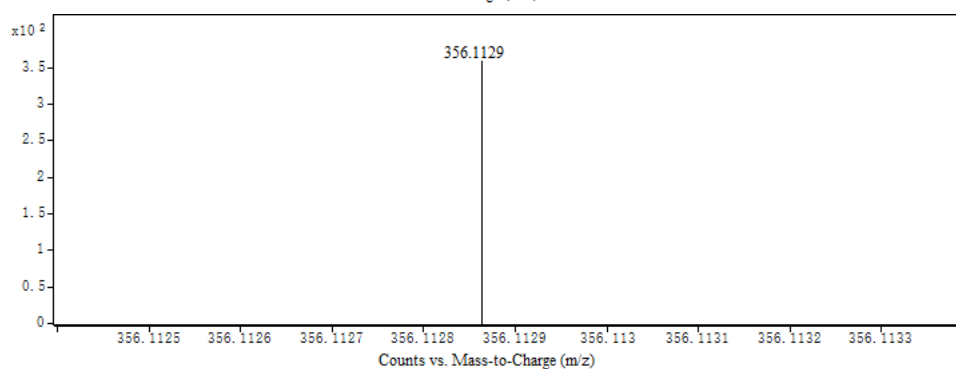
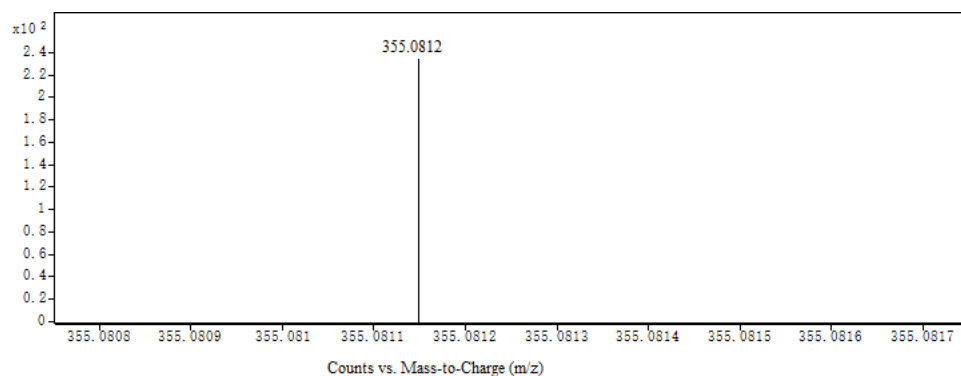
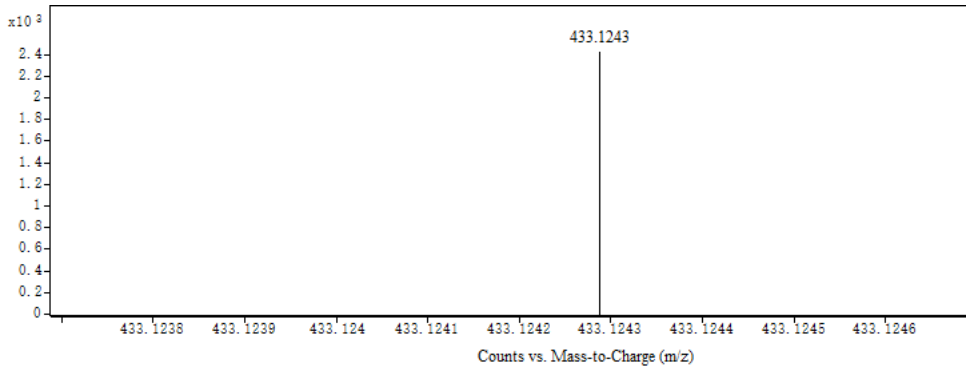
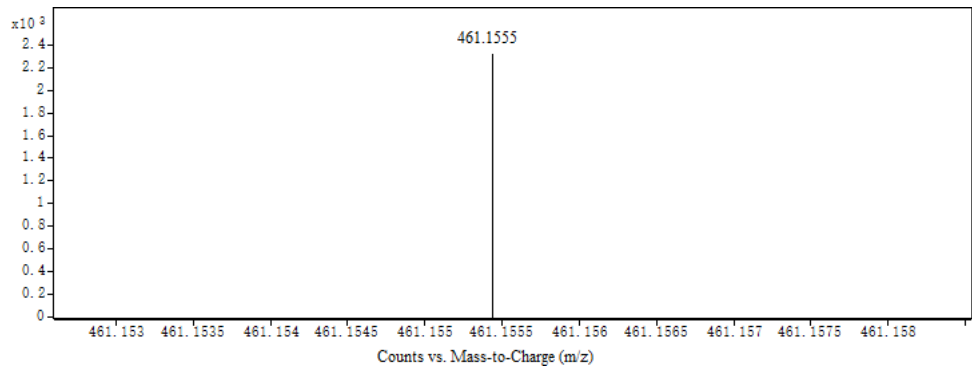
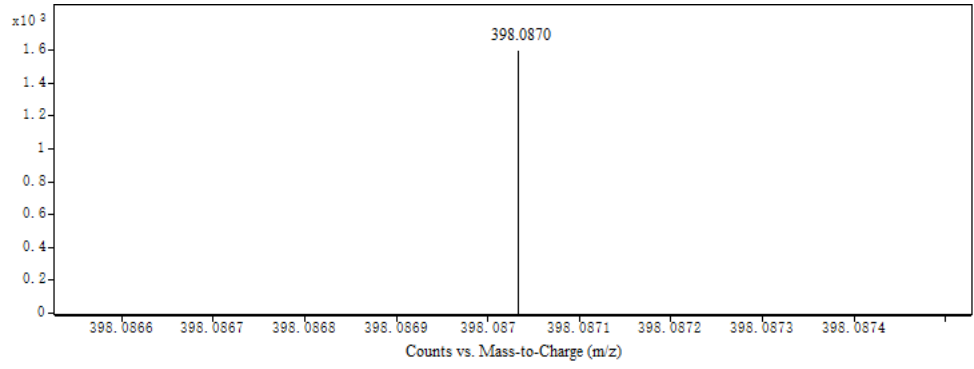
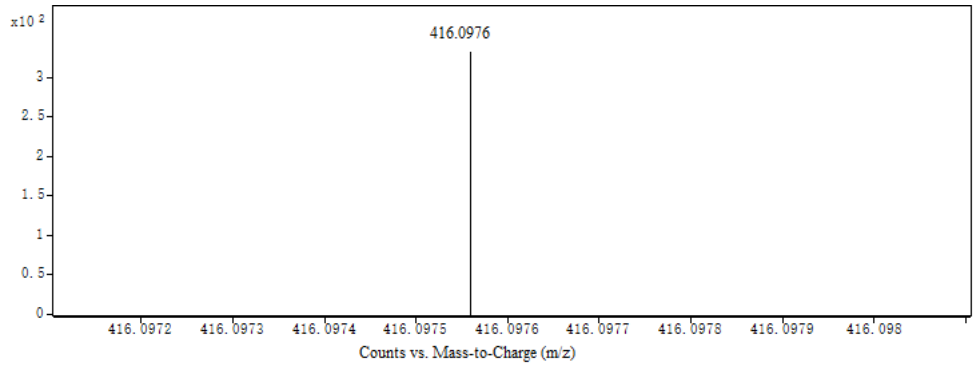


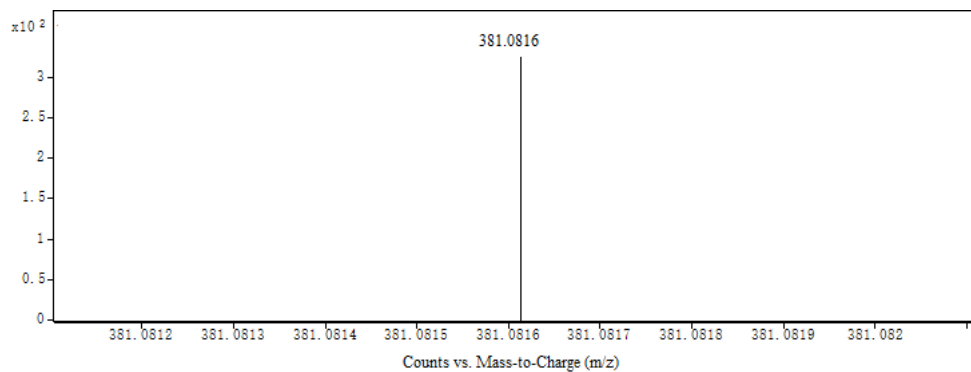
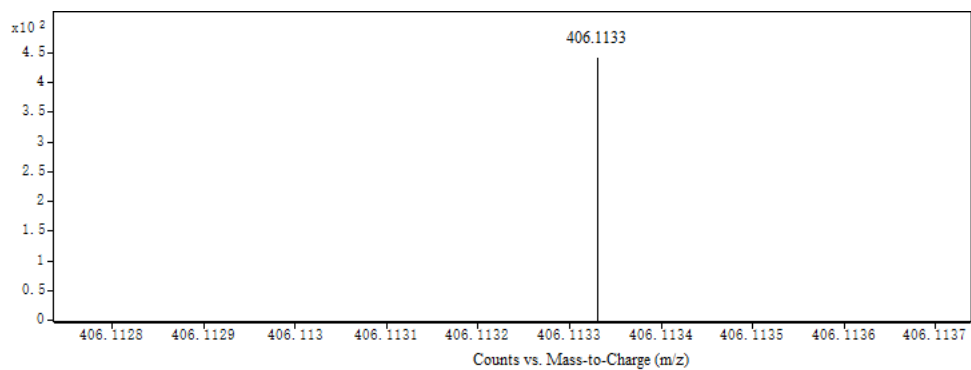
Fig. S1. Separation mechanism of BiVO_4 composites

Fig. S2. MS spectra of intermediates during the degradation of DC

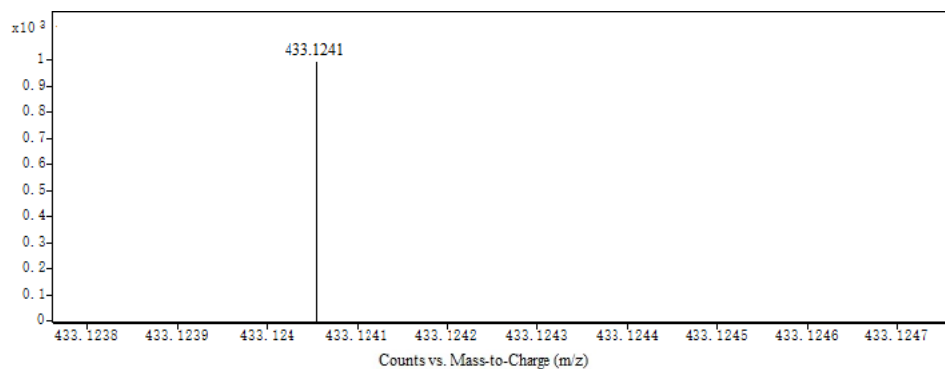
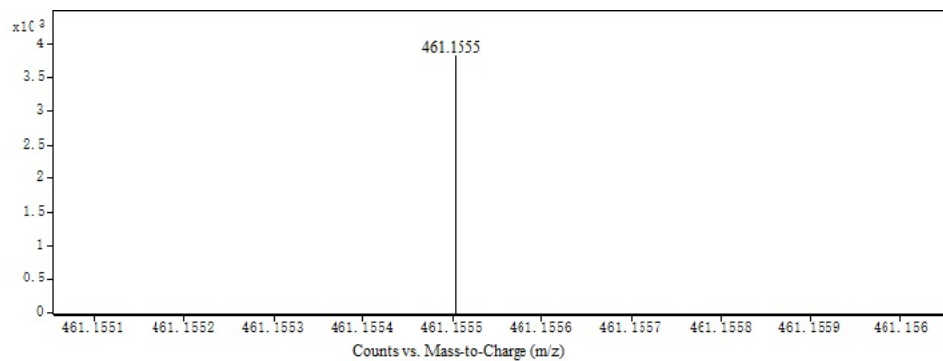
(1) *The intermediates in photocatalytic BiVO₄ system*

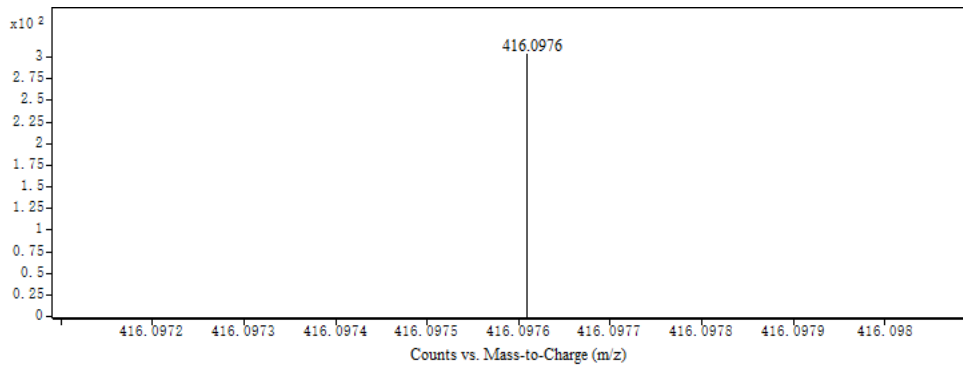
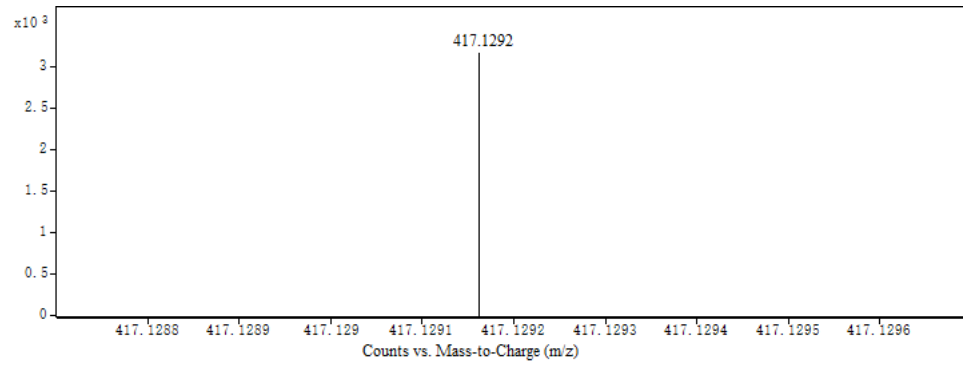
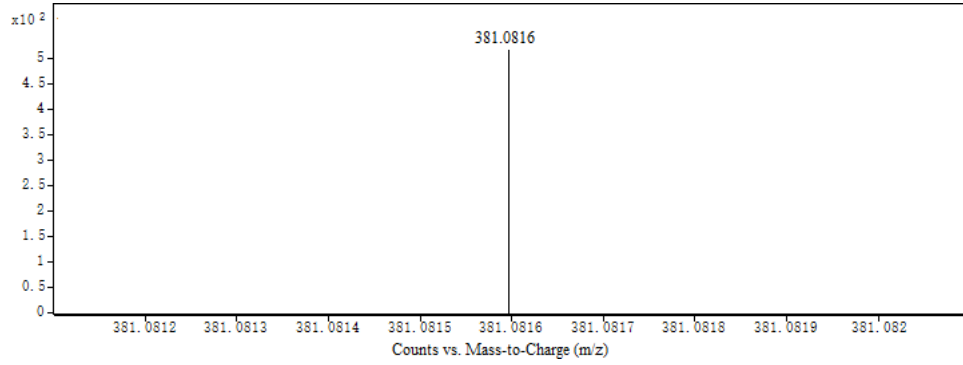
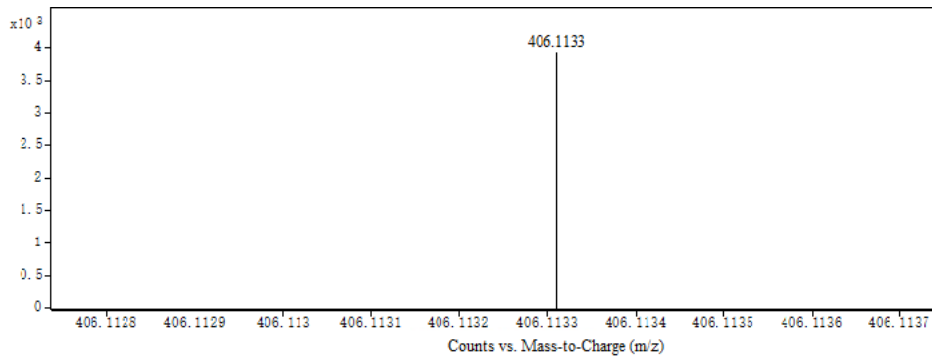






(2) *The intermediates in photocatalytic $\text{BiVO}_4/\text{Ag}_2\text{O}$ system*





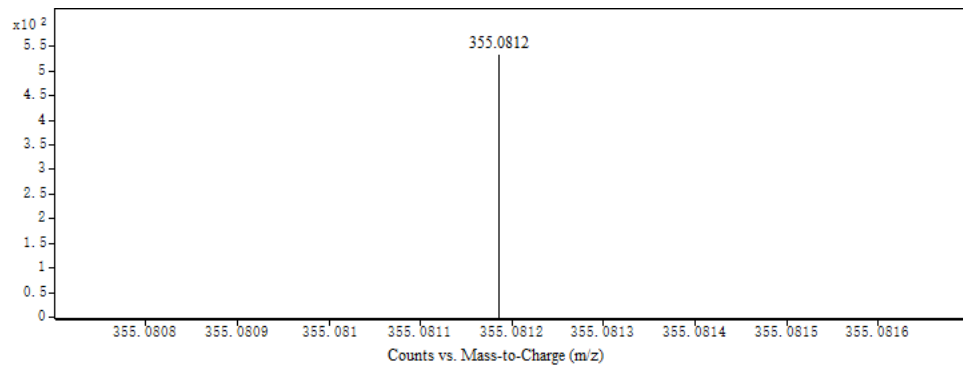
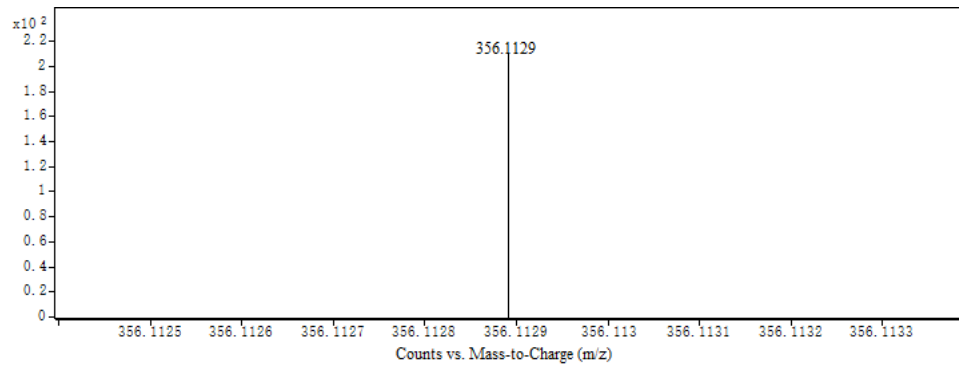
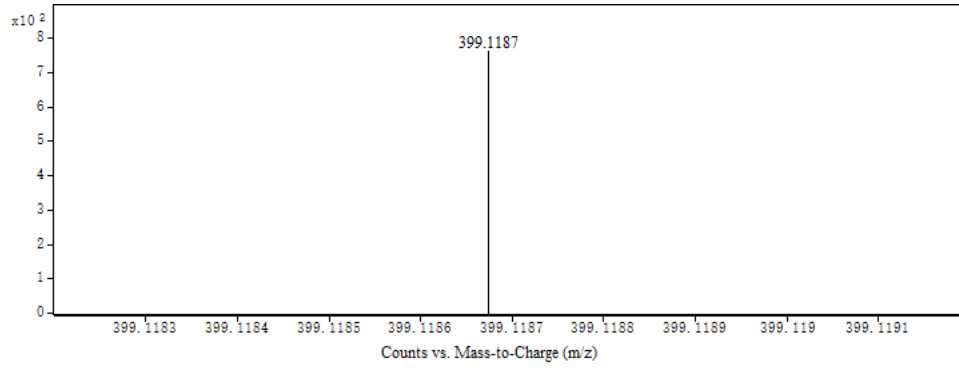
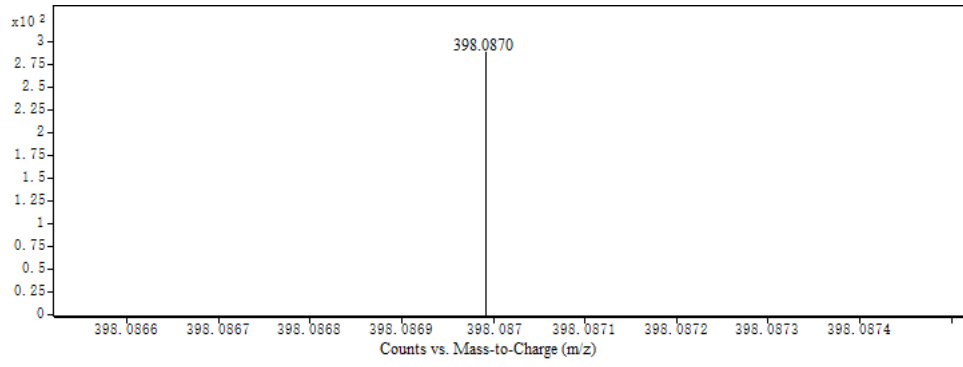


Table. S1-1 Identified intermediates of DC degradation in photocatalytic BiVO₄ system and photocatalytic BiVO₄@Ag₂O system

Catalyst	Molecular formula	Theoretical value (m/z) [M+H]	Actual value (m/z) [M+H]	Error (ppm)
BiVO ₄	C ₂₀ H ₂₀ N ₂ O ₈	417.1292	417.1292	0.0
	C ₂₀ H ₁₇ NO ₉	416.0976	416.0976	0.0
	C ₂₀ H ₁₅ NO ₈	398.0870	398.0870	0.0
	C ₂₀ H ₁₈ N ₂ O ₇	399.1187	399.1187	0.0
	C ₁₉ H ₁₇ NO ₆	356.1129	356.1129	0.0
	C ₁₉ H ₁₄ O ₇	355.0812	355.0812	0.0
	C ₂₂ H ₂₄ N ₂ O ₉	461.1555	461.1555	0.0
	C ₂₀ H ₂₀ N ₂ O ₉	433.1242	433.1243	0.2
	C ₁₉ H ₁₉ NO ₉	406.1133	406.1133	0.0
	C ₁₇ H ₁₆ O ₁₀	381.0816	381.0816	0.0
	C ₂₀ H ₂₀ N ₂ O ₈	417.1292	417.1292	0.0
	C ₂₀ H ₁₇ NO ₉	416.0976	416.0976	0.0
	C ₂₀ H ₁₅ NO ₈	398.0870	398.0870	0.0
BiVO ₄ @Ag ₂ O	C ₂₀ H ₁₈ N ₂ O ₇	399.1187	399.1187	0.0
	C ₁₉ H ₁₇ NO ₆	356.1129	356.1129	0.0
	C ₁₉ H ₁₄ O ₇	355.0812	355.0812	0.0
	C ₂₂ H ₂₄ N ₂ O ₉	461.1555	461.1555	0.0
	C ₂₀ H ₂₀ N ₂ O ₉	433.1242	433.1241	-0.2
	C ₁₉ H ₁₉ NO ₉	406.1133	406.1133	0.0
	C ₁₇ H ₁₆ O ₁₀	381.0816	381.0816	0.0