Supporting Information File

Insights on the use of Carbon Additives as Promoters of the Visible-Light Photocatalytic Activity of Bi₂WO₆



Figure S1. Emission spectrum of the lamp used in the photocatalytic experiments.



Figure S2. SEM images of the carbon materials used as additives (samples CS, CB and CL).



Figure S3. X-ray diffraction patterns of A) BWO and BWO/carbon catalysts and B) the carbon materials used as additives. Diffractograms have been shifted for clarity.



Figure S4. Nitrogen adsorption/desorption isotherms at 77 K of the studied photocatalysts.

	Conversion at 120 min	TOC initial	TOC final	Mineralization ([1-(Final TOC/InitialTOC)])
	(%)	(mgC/L)	(mgC/L)	(%)
RhB Photolysis	6	6.8	6.5	4.4
BWO	89	7.4*	0.3	95.9
BWO/CS-2	99	6.3	0.1	98.4
BWO/CL-2	96	8.3	0.3	96.4
BWO/CB-2	98	6.8	-	-
BWO/CS-5	99	8.3	0.14	98.3
BWO/CL-5	94	7.4	0.42	94.3
BWO/CB-5	95	6.3	-	-

Table S1. Rhodamine B conversion and Total Organic Carbon (TOC) values upon 2 h of irradiation of the studied catalysts.

*Measured at 180 min



Figure S5. Rhodamine B conversion upon exposure to simulated solar light of the catalysts with 2 (A) and 5 wt. % (B) of carbon additive.



Figure S6. Evolution of RhB photooxidation intermediates (Rhodamine, R; N-ethylrhodamine, ER; N,N-diethylrhodamine, DR; N-ethyl-N''-ethylrhodamine, EER; N,N-diethyl-N''-ethylrhodamine, DER) upon irradiation of photocatalysts BWO, BWO/CS-2 and BWO/CL-2.

Table S2. Surface concentration of carbon species obtained by fitting the C 1*s* core level peak of the XPS spectra of composites BWO/CL-2 and BWO/CS-2 as received (fresh) and after irradiation of an aqueous dispersion to explore the stability of the carbon component.

Bond assignement (energy, eV)	Fresh	Irradiated
BWO/CL-2		
C–C (graphitic carbon - 284.6 eV)	59.3	62.7
C–O (phenolic, alcoholic, etheric - 286.1 eV)	13.8	13.1
C=O (carbonyl or quinone - 287.1 eV)	17.7	18.2
O–C=O (carboxyl or ester - 288.7 eV)	5.8	6.0
π-π* (291.0 eV)	3.4	3.5
BWO/CS-2		
C–C (graphitic carbon - 284.6 eV)	66.5	65.1
C–O (phenolic, alcoholic, etheric - 286.1 eV)	15.9	17.5
C=O (carbonyl or quinone - 287.1 eV)	6.8	7.8
O–C=O (carboxyl or ester - 288.7 eV)	7.5	8.2
π–π* (291.0 eV)	3.3	1.3



Figure S7. (left) Characteristic ESR signals corresponding to DMPO-OH adducts obtained upon 20 min of irradiation of an aqueous suspension of BWO and BWO/carbon photocatalysts in the presence of DMPO as trapping agent; (right) Quantification of O-radical species by integration of the second peak in the 1:2:2:1 quartet profile of the DMPO-OH adducts of selected photocatalysts.



Figure S8. Luminescence inhibition of *Vibrio Fischeri* bacteria upon exposure to Rhodamine B aqueous solutions for 15 min. The toxicological parameter EC_{50} determined as the concentration of RhB for a 50% inhibition was ca. 2.5 ppm.