

Supporting Information

Enhanced photocatalytic properties and photo-induced crystallization of TiO₂-Fe₂O₃ inverse opals fabricated by atomic layer deposition

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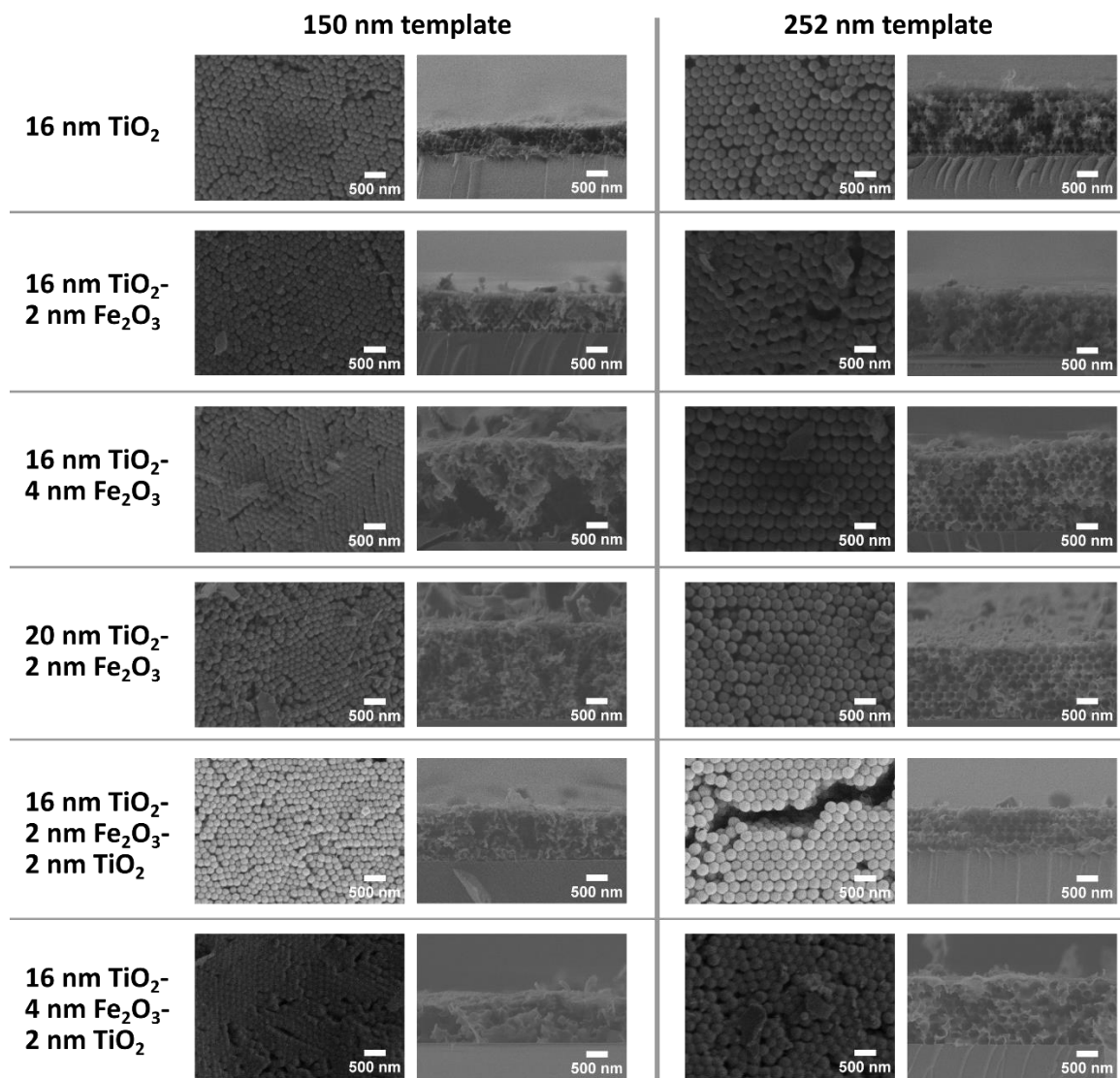


Figure S1. SEM micrographs of TiO₂-Fe₂O₃ IOs and TiO₂-Fe₂O₃ multilayer IOs fabricated with different layer thicknesses and template sizes.

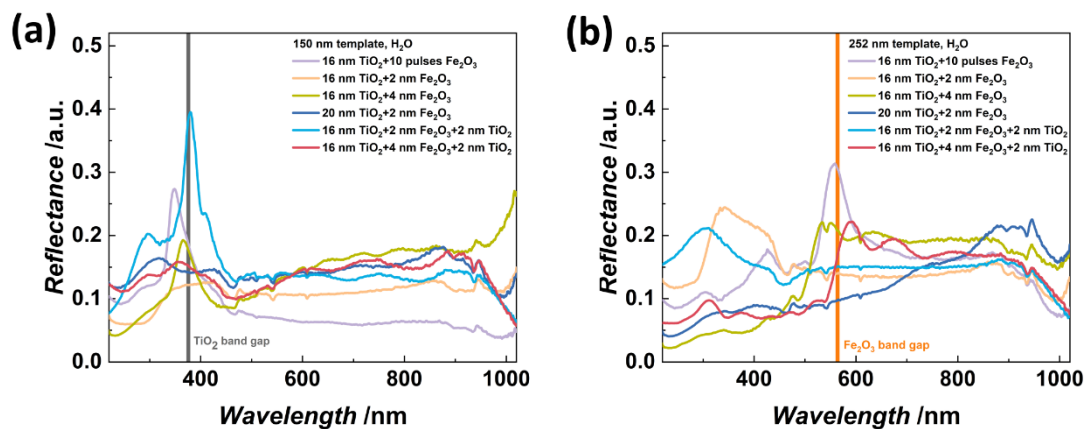


Figure S2. Optical properties of all prepared IOs measured in aqueous environment. The samples were prepared with PS particle template sizes of (a) 150 nm and (b) 252 nm.

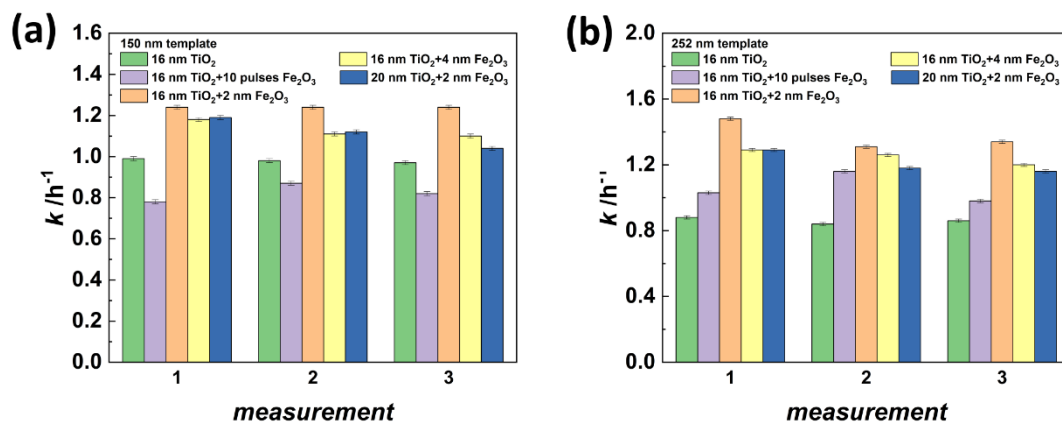


Figure S3. Individual photocatalytic activities of three consecutive measurements for TiO₂ IOs and TiO₂-Fe₂O₃ bilayer IOs. Different TiO₂ and Fe₂O₃ thicknesses were tested for template sizes of (a) 150 nm and (b) 252 nm. The

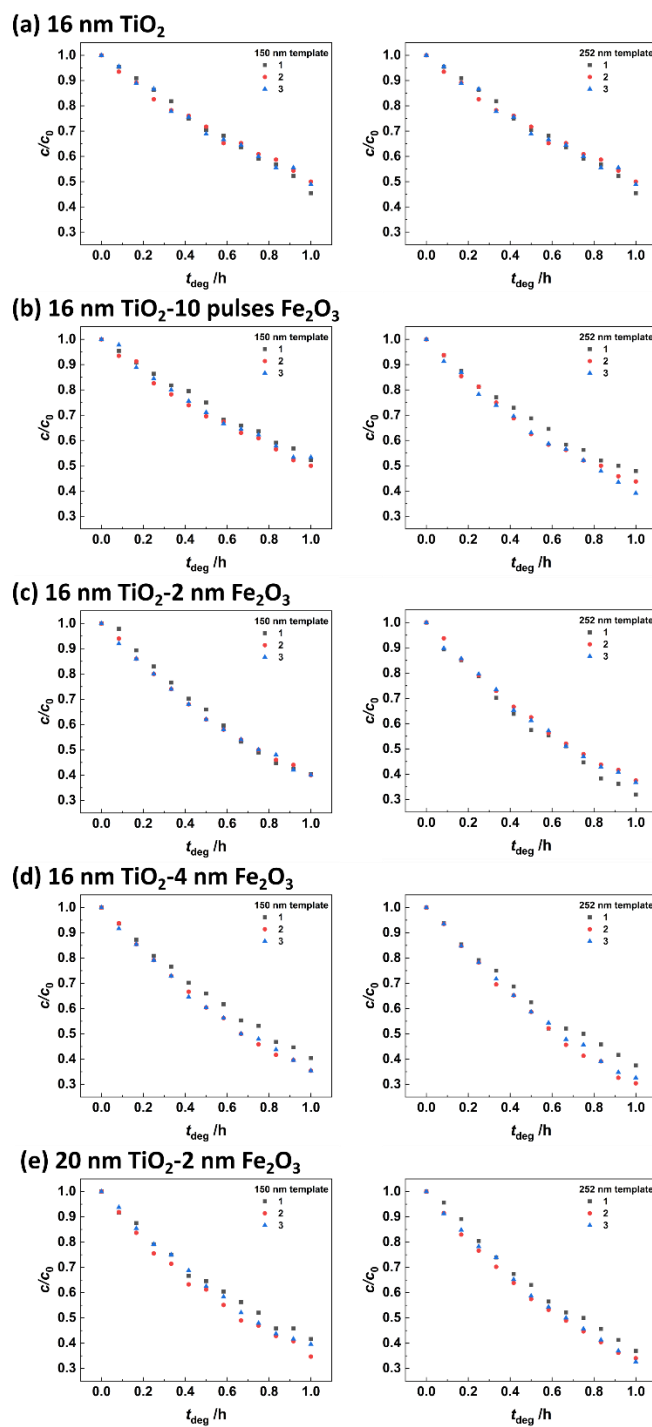


Figure S4. Dye concentration decrease during three consecutive photocatalysis measurements for TiO_2 IOs and $\text{TiO}_2\text{-Fe}_2\text{O}_3$ bilayer IOs. The data represents the MB degradation by (a) 16 nm TiO_2 IOs, (b) 16 nm $\text{TiO}_2\text{-10 pulses Fe}_2\text{O}_3$, (c) 16 nm $\text{TiO}_2\text{-2 nm Fe}_2\text{O}_3$, (d) 16 nm $\text{TiO}_2\text{-4 nm Fe}_2\text{O}_3$, and (e) 20 nm $\text{TiO}_2\text{-2 nm Fe}_2\text{O}_3$.

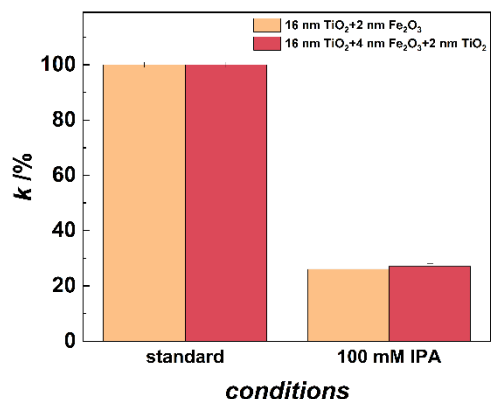


Figure S5. Assessment of the photocatalytic activities with MB solution containing 100 mM IPA as hole scavenger. Both, the 16 nm TiO₂–2 nm Fe₂O₃ bilayer IO and the 16 nm TiO₂–4 nm Fe₂O₃–2 nm TiO₂ trilayer IO demonstrate significant reduction of their photocatalytic activity compared to standard conditions. Each sample was measured three times with IPA containing solution.

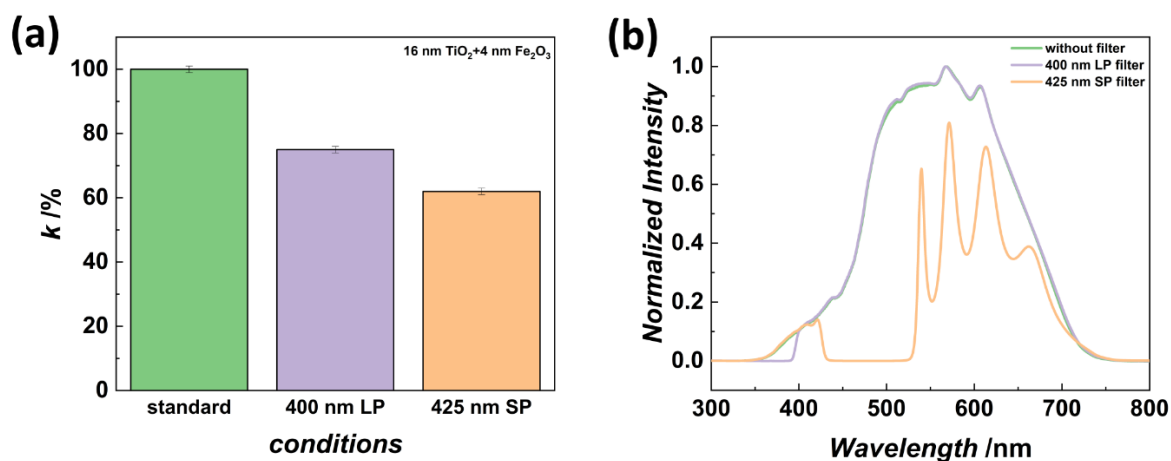


Figure S6. Photocatalytic activities under standard illumination conditions without a filter, with a 400 nm longpass (LP) filter, and with a 425 nm shortpass (SP) filter, respectively. (a) Activities of the 16 nm TiO₂–4 nm Fe₂O₃ bilayer IO normalized to the illumination without filter. The sample was measured once with each illumination spectra. (b) Optical illumination spectra of the light source with and without filters.

Table S1. Comparison of the reaction conditions and photocatalytic performances of Fe₂O₃-functionalized TiO₂ nanostructures.

Sample	Fabrication method	Organic pollutant	Illumination	Photocatalytic activity k	reference
Fe ₂ O ₃ -coated TiO ₂ powder	Coating by ALD	Methyl orange 4 mg/L	300 W Xe lamp with 420 nm shortpass filter	97.4 % removal after 1.5 h	22
Fe ₂ O ₃ -coated TiO ₂ nanocrystals	Hydrothermal method	Rhodamine B 50 μM, H ₂ O ₂	8 W daylight white LED	52 % removal after 1 h	23
Fe ₂ O ₃ -coated TiO ₂ nanoporous structures	ALD	PEC characterization	150 W Xe lamp		26
Fe ₂ O ₃ -coated TiO ₂ microrod powder	Hydrothermal method	Orange II 20 mg/L	500 W Xe lamp with 420 nm shortpass filter	54 % removal after 3 h	28
Fe ₂ O ₃ -coated TiO ₂ IO	TiO ₂ IO by ALD, Fe ₂ O ₃ coating hydrothermal method	PEC characterization	300 W Xe lamp		33
Fe ₂ O ₃ -decorated TiO ₂ IO	TiO ₂ IO by sol-gel method, Fe ₂ O ₃ decoration by chemisorption calcination cycles	salicylic acid	150 W Xe lamp	~0.9 h ⁻¹ , 75 % removal after 1.5 h	34
TiO ₂ -Fe ₂ O ₃ multilayer IOs	ALD	Methylene blue 2.5 mg/L	150 W halogen lamp	1.38 h ⁻¹ ; 65 % removal after 1 h	This study

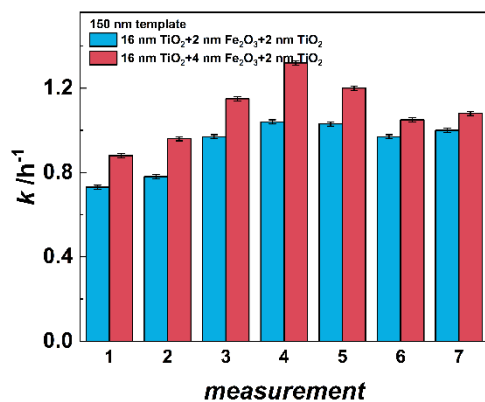


Figure S7. The individual activities during seven consecutive measurements of 150 nm template size 16 nm TiO₂–2 nm Fe₂O₃–2 nm TiO₂ and 16 nm TiO₂–4 nm Fe₂O₃–2 nm TiO₂ multilayer IOs show the same behavior as the 252 nm template size, namely increase during the first four measurements, slight decline in the following two measurements, and stable performance afterwards.

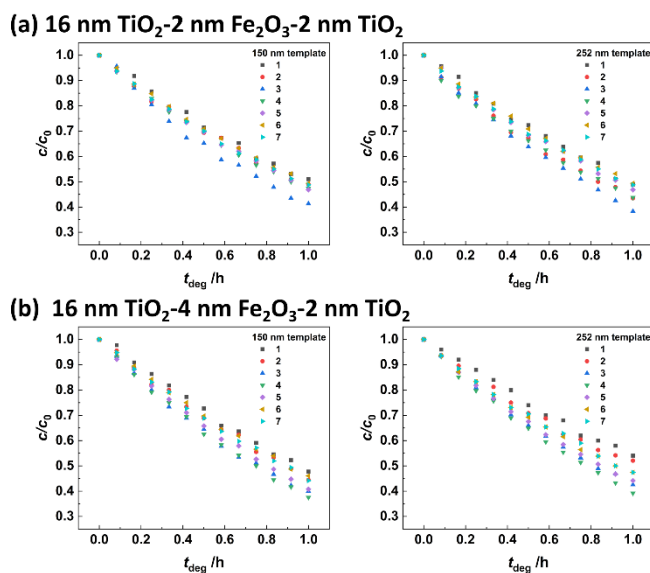


Figure S8. MB concentration decrease during seven consecutive measurements of (a) 16 nm TiO₂–2 nm Fe₂O₃–2 nm TiO₂ and (b) 16 nm TiO₂–4 nm Fe₂O₃–2 nm TiO₂ multilayer IOs.

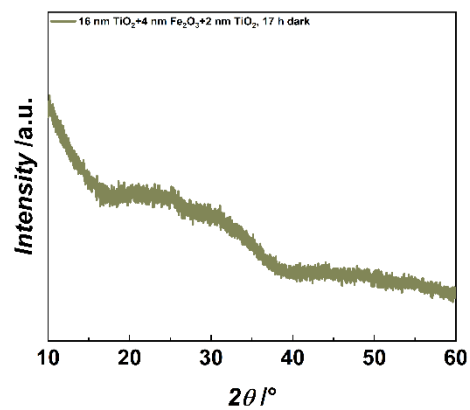


Figure S9. XRD pattern of a TiO_2 - Fe_2O_3 multilayer IO composed of 16 nm TiO_2 , 4 nm Fe_2O_3 , and 2 nm TiO_2 . The sample was kept in the reaction solution for 17 h in darkness and does not show peaks indicating crystalline TiO_2 phases.