

Supporting information

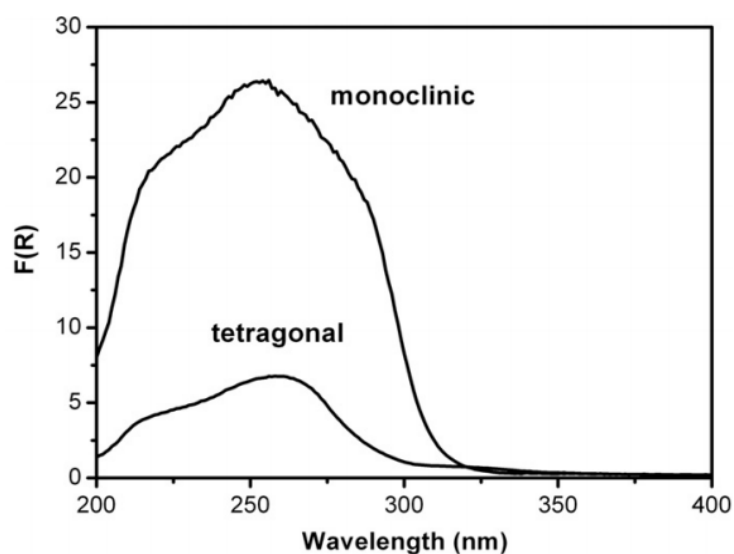


Figure S1. UV-vis diffuse reflectance spectra for both polymorphs of monoclinic and tetragonal CdWO_4 .

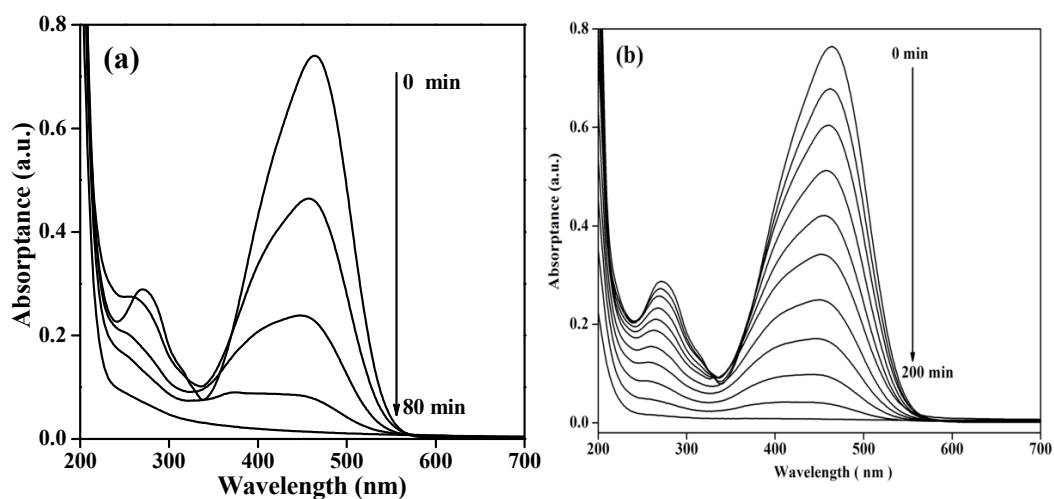


Figure S2. Time dependent absorption spectra of MO solutions in the presence of (a) $m\text{-CdWO}_4$ and (b) $t\text{-CdWO}_4$ under UV light irradiation. ($m\text{-CdWO}_4$ was prepared in water at 200 °C for 10 h. $t\text{-CdWO}_4$ was prepared in 1,2-propanediol at 200 °C for 10h.)

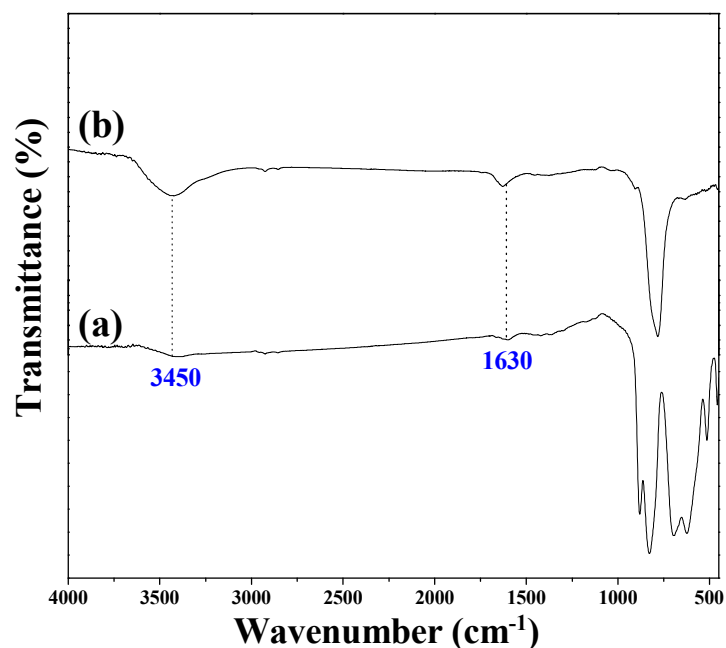


Figure S3. FT-IR spectrum for the as-prepared: (a) *m*-CdWO₄, and (b) *t*-CdWO₄.

Infrared spectra of the samples were measured on a Perkin-Elmer IR spectrophotometer at a resolution of 4 cm⁻¹ by KBr pellet technique. The band observed at 3450 cm⁻¹ is attributed to the vibrations of O–H bonds for the water absorbed on sample surfaces and the band centered at 1630 cm⁻¹ is associated with the deformation vibration for H–O–H bonds of the physisorption water. The bands appeared at 400~1000 cm⁻¹ are the characteristic of the intrinsic vibrations for *t*-CdWO₄ and *m*-CdWO₄, which are apparently different.

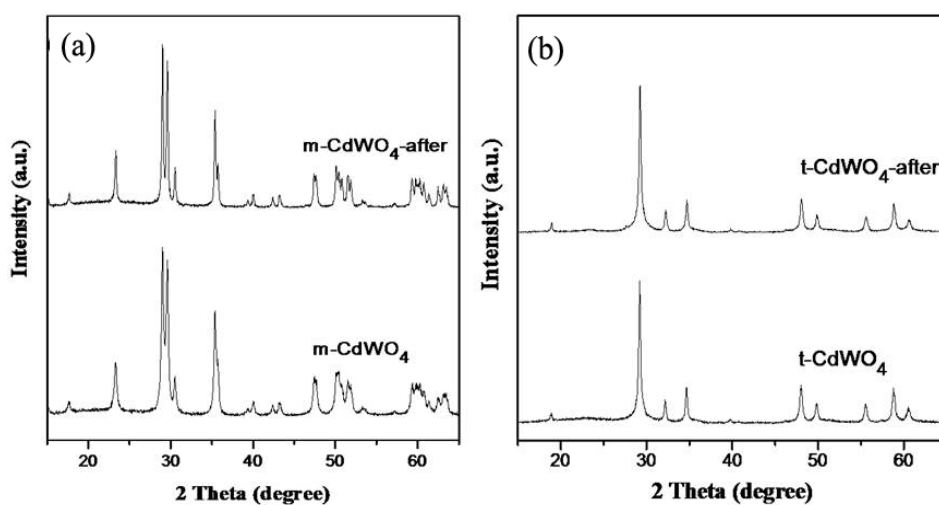


Figure S4. XRD patterns of the as-prepared: (a) *m*-CdWO₄, and (b) *t*-CdWO₄ before and after photocatalytic reaction.