

Enhanced electrochromic switches and tunable green fluorescence based on terbium ion doped WO₃ film

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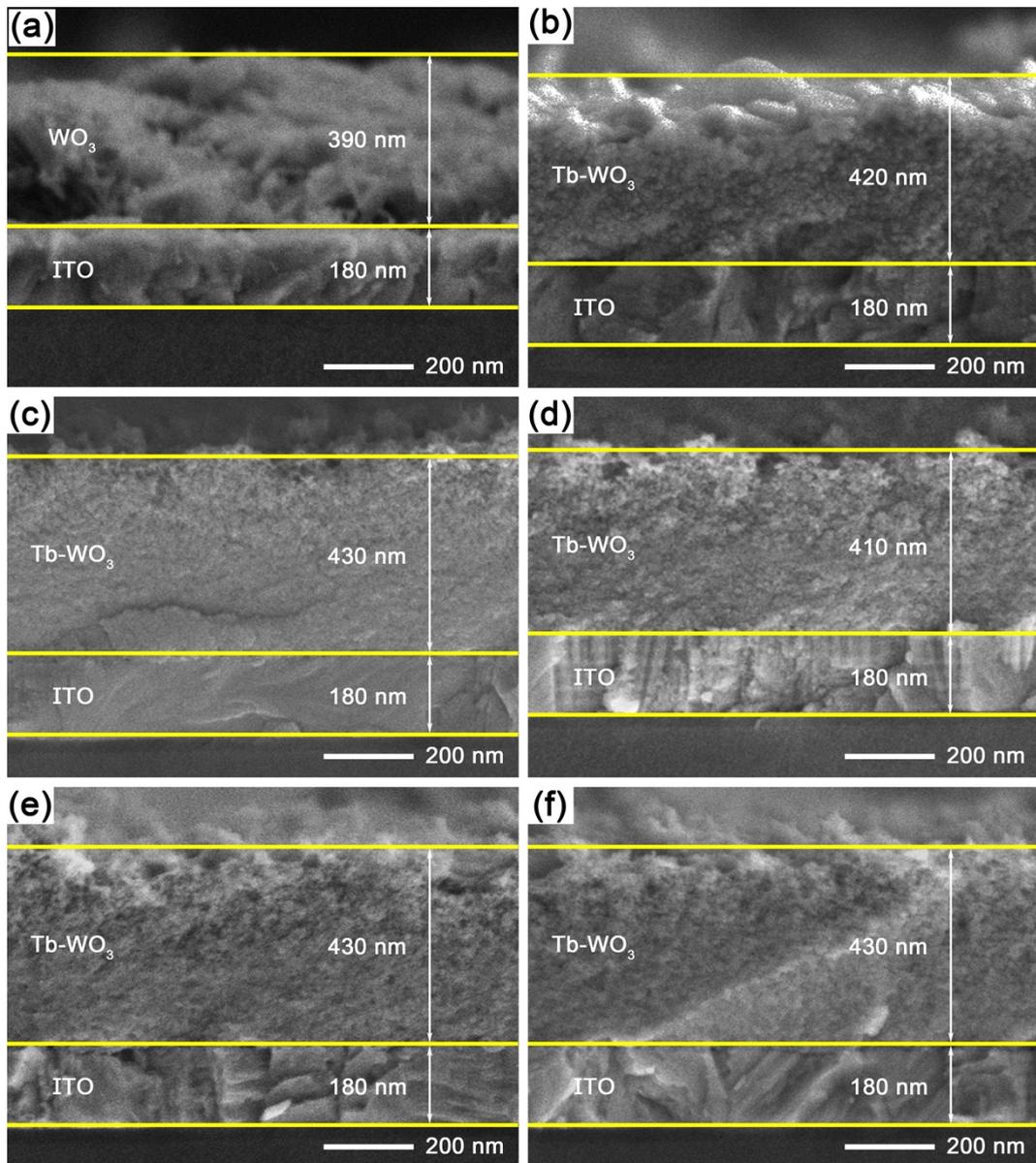


Fig. S1. SEM cross-sectional views of Tb-WO_3 films with different doping contents: (a) 0%, (b) 5%, (c) 10%, (d) 12%, (e) 13% and (f) 15%.

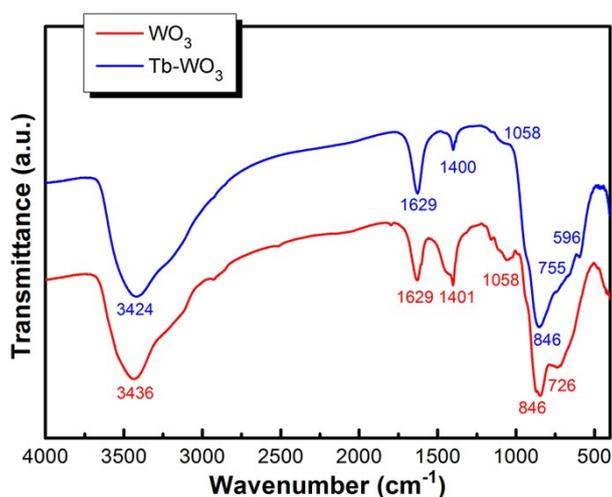


Fig. S2. FT-IR spectra of pristine and typical 13% Tb doped WO_3 films in the 4000-400 cm^{-1} wavenumber region.

FT-IR spectra in Fig. S1 were collected to further elucidate the structure changes induced by Tb ions doping. The spectrum of pristine WO_3 film can be roughly divided into two regions: 500 cm^{-1} to 1100 cm^{-1} with vibrations of the W-O-W and O-W-O bonds, and above 1300 cm^{-1} with vibrations of the H-O-H bonds. The additional peak at 846 nm can be attributed to W=O bonds. Tb doped WO_3 film exhibits the similar characteristic absorption of WO_3 species, indicating that the Tb ions may have entered into the WO_3 lattice without the obvious change of chemical band.

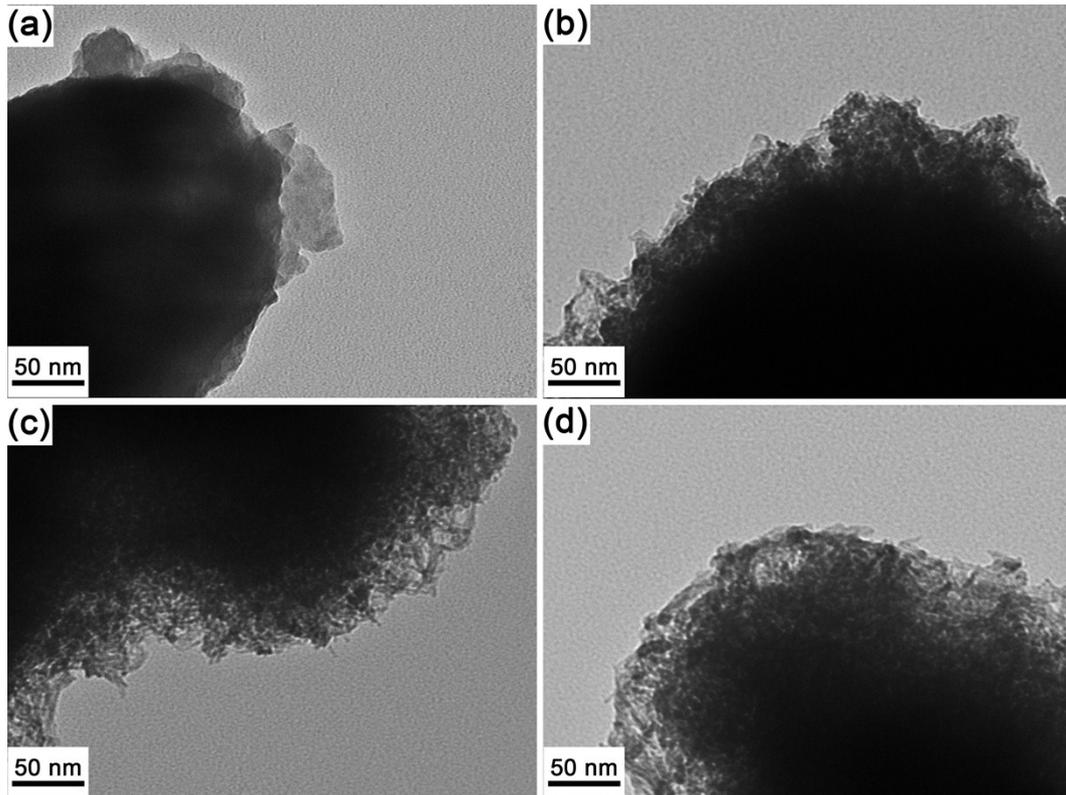


Fig. S3. TEM images of Tb-WO₃ films with different doping contents: (a) 5%, (b) 10%, (c) 12%, (d) 15%.

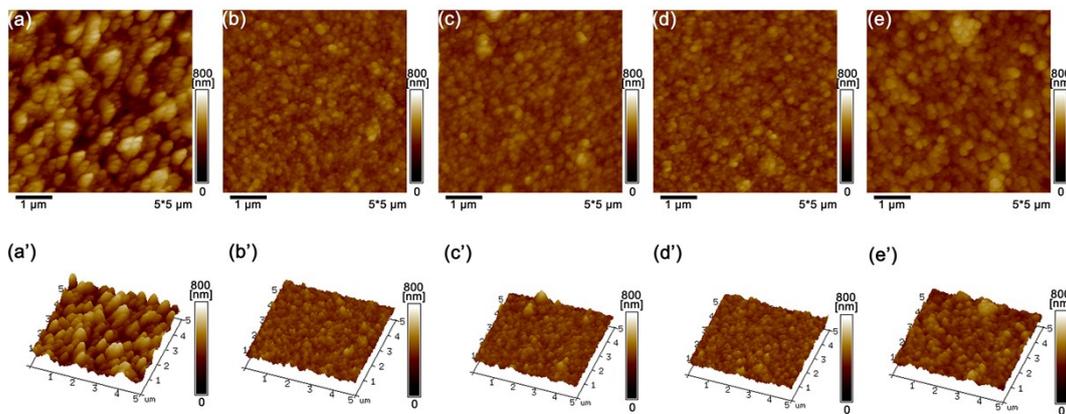


Fig. S4. AFM images in (a-e) surface topography and (a'-e') 3D stereogram of Tb-WO₃ film with various doping concentrations (0%, 5%, 10%, 12% and 15%).

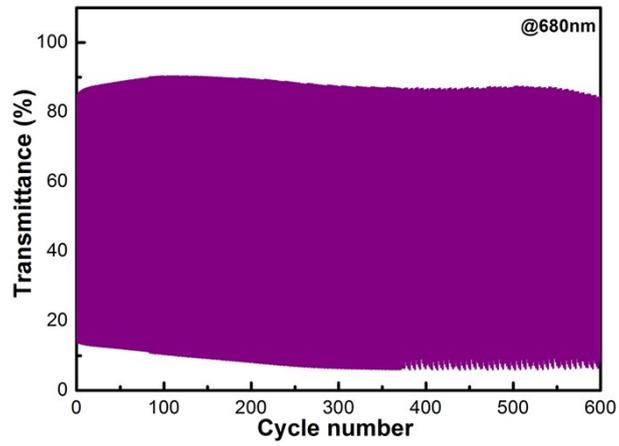


Fig. S5. Cyclic stability of transmittance switching for 13% Tb-doped WO_3 film monitored at 680 nm. The film was cycled in 0.5 M H_2SO_4 aqueous electrolyte under double-potential step between at -1 V for 15 s and 1.5 V for 15 s vs. Ag/AgCl.