Supporting Information

Solvent-Induced Surface Disorder and Doping-Induced Lattice Distortion in Anatase TiO₂ Nanocrystals for Enhanced Photoreversible Color Switching

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Fig. S1 Rietveled refined XRD patterns for TiO₂/H₂O (a), TiO₂/EtOH (b), TiO₂/EG (c), TiO₂/DEG (d), and TiO₂/PEG200 (e). Black cross stand for the observed value, red line represents the fitting line, blue curve is the difference of observed and the calculated curve and black bar is the peak position. (f) Boiling point of the solvent dependences of the lattice parameter *a*, *c* values for TiO₂ NCs.



Fig. S2 (a) Enlarged XRD patterns and (b) FWHM value of (101) peak of all the samples.



Fig. S3 (a) Raman spectra of TiO_2 NCs synthesized in different solvents. (b) FT-IR spectra of EG, DEG, PEG200.



Fig. S4 The decoloration and recoloration process of the TiO₂/MB/H₂O system: (a, b) TiO₂/H₂O, (c, d) TiO₂/EtOH, (e, f) TiO₂/EG, (g, h) TiO₂/DEG, and (i, j) TiO₂/PEG200.



Fig. S5 (a) FT-IR spectra of TiO_2/EG , TiO_2/DEG , $TiO_2/PEG200$ after cycling. (b) Schematic illustration of the change of surface functional groups of TiO_2 NCs.

Table S1 Parameters related to the lattice of TiO_2/H_2O , $TiO_2/EtOH$, TiO_2/EG , TiO_2/DEG , $TiO_2/PEG200$ NCs determined by linear fit of the Halder-Wagner plot.

| Samples | Crystalline size (nm) | Linear fit of the Halder-Wagner plot | | |
|------------------------------------|--------------------------|--------------------------------------|-------------------|---------|
| | | Slope | Y intercepts | |
| TiO ₂ /H ₂ O | 11.4 | 1.1598 ± 0.39314 | 0.01389 ± 0.02205 | 0.96501 |
| TiO ₂ /EtOH | 9.3 | 1.22686 ± 0.11524 | 0.01273 ± 0.00743 | 0.99273 |
| TiO ₂ /EG | 19.8 | 0.6701 ± 0.128 | 0.00479 ± 0.00447 | 0.98409 |
| TiO ₂ /DEG | 19.7 | 0.65286 ± 0.11237 | 0.00516 ± 0.00392 | 0.98773 |
| TiO ₂ /PEG200 | 4.4 | 2.67788 ± 0.24131 | 0.00295 ± 0.02972 | 0.99351 |



Fig. S6 Rietveled refined XRD patterns of (a) 1%Sn²⁺:TiO₂, (b) 3%Sn²⁺:TiO₂, (c) 5%Sn²⁺:TiO₂, and (d) 10%Sn²⁺:TiO₂. Black cross stands for the observed value, red line represents the fitting line, blue curve is the difference of observed and the calculated curve and black bar is the peak position.



Fig. S7 Raman spectra of Sn^{2+} : TiO₂ NCs having different doping concentrations.



Fig. S8 Low-magnification TEM patterns of (a) $TiO_2 NCs$, (d) 3% Sn^{2+} : $TiO_2 NCs$ and (g)10% Sn^{2+} : $TiO_2 NCs$. Intensity line profile (c), (f) and (i) extracted from the HRTEM images of (b) $TiO_2 NCs$, (e) 3% Sn^{2+} : $TiO_2 NCs$, and (h) 10% Sn^{2+} : $TiO_2 NCs$.



Fig. S9 The decoloration and recoloration process of Sn²⁺ doped TiO₂/MB/H₂O system: (a, b) 1%Sn²⁺:TiO₂, (c, d) 3%Sn²⁺:TiO₂, (e, f) 5%Sn²⁺:TiO₂, and (g, h) 10%Sn²⁺:TiO₂.



Fig. S10 Diagram of the relationship between the change of Zeta potential and the solution system with different pH values of 3%Sn²⁺:TiO₂ and 10% Sn²⁺:TiO₂.