Order and Disorder Effects in Nano- ZrO₂ Investigated by Micro- Raman and Spectrally and Temporarily Resolved Photoluminescence

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Supplemental Information



Figure S1. STEM images of calcined pure zirconia samples (ZE) synthesized in o/w microemulsions. ZE-500: (a) bright field and (b) dark field. ZE-1000: (c) bright field and (d) dark field.



Figure S2. STEM images of calcined europium- doped zirconia (ZEB) samples synthesized in o/w microemulsions. 500/ZEB-500: (a) bright field and (b) dark field. 1000/ZEB-1000: (c) bright field and (d) dark field.



Figure S3. TGA and DTA curves for pure (ZE, a) and europium doped zirconia (ZEB, b) samples.

Textural characterization. Pure zirconia, ie ZE sample, after calcining at 350 °C exhibited a surface area of 253 m²g⁻¹ with an average pore size of 26.9 Å. Further calcinations at 500 and 1000 °C led to a strong decrease of the surface areas to 24 and 4 m²g⁻¹, respectively. Doping with europium (ZEB sample) was not generating important differences in the texture of these materials. Thus, the surface areas of the samples calcined at 350, 500 and 1000 °C corresponded to 248, 31 and 5 m²g⁻¹, that are in fact very similar.



Figure S4. PL excitation spectra of europium in ZEB-500 and ZEB-1000.



Figure S5. (a) Steady state PL spectra of ZEB-500 and ZEB- 1000; (b) Time-resolved PL spectra of ZEB- 1000 upon excitation at 240 nm.



Figure S6. From Left to Right: PL decays of ZEB- 750, ZEB- 900 and ZEB- 1000. Emission and excitation wavelengths are indicated on the Figures.



Figure S7. Evolution with delay time of the asymmetry ratio, R, of ZEB- 500 and ZEB- 1000.