SUPPORTING INFORMATION

Tuning Ceria Catalysts in Aqueous Media at the Nanoscale: How Do Surface Charge and Area Determine Peroxidase- and Haloperoxidase-Like Reactivity

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Fig. S1. X-ray powder diffraction pattern of $Ce(C_2O_4)(HCO_2)$ formed after 120 h in a flask reaction. Red ticks indicate the calculated reflection positions for $Ce(C_2O_4)(HCO_2)$ based on structural data from the ICSD data bank (ICSD# 43450).



Fig. S2. SAXS measurements (A) of mesoporous CeO_{2-x} nanoparticles formed after 2.0 h, 2.5 h, 3.0 h and 3.5 h of reaction time, with (B) and (C) showing the red and blue regions marked in (A) in more detail. Arrows are a guide for the eye to show the shift to larger values of q for 2.0 h, 2.5 h and 3.0 h samples.



Fig. S3. Average radial dimension of the four populations observed from fits of the SAXS for mesoporous CeO₂. _x nanoparticles formed at different reaction times. Fit regions refer to the population distributions and are set to the ranges of (*P1*) 0.5 - 2.5 nm, (*P2*) 2.5 - 10 nm, (*P3*) 10 - 30 nm and (*P4*) 30 - 100 nm, though for 2.0 h reaction times regions *P3* and *P4* are set between 10 - 20 nm and 20 - 100 nm respectively.



Fig. S4. Particle radial histograms derived from small angle X-ray scattering data with cumulative abundance (right y-axis) for mesoporous CeO_{2-x} nanoparticles formed after (A) 2.0 h, (B) 2.5 h, (C) 3.0 h and (D) 3.5 h of reaction time.



Fig. S5. Digital images of the reaction solutions after 2.0 h, 2.5 h, 3.0 h and 3.5 h.



Fig. S6. Scanning electron microscopy images of mesoporous CeO₂ nanoparticles formed after 2.0 h.