## **Supplementary Information**

## Water-based synthesis and cleaning methods for high purity ZnO nanoparticles – comparing acetate, chloride, sulphate and nitrate zinc salt precursors

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**Fig. S1** Scanning Electron micrographs and size distributions (counting number = 250) of commercial ZnO particles a) NanoTek b) MKnano



**Fig S2.** Scanning Electron micrograph of octahedron ZnO particles showing twin petals. The polygons (twin petals) were synthesized under stoichiometric conditions using the zinc nitrate salt at reaction yield 4g/L (60°C).



Fig S3. Scanning Electron micrograph of ZnO particles showing intermediate early steps in the formation of flower-shaped particles by growing spikes on the octahedron polygones. The particles were synthesized under stoichiometric conditions using the zinc nitrate salt at reaction yield 4g/L ( $80^{\circ}C$ ).



**Fig S4.** High-resolution transmission electron micrograph of ZnO particles showing the basal plane of prism with an inter-planar distance of 0.28 nm.

## Video 1

**Video 1.** Transformation of facetted salt entities from ion suspension of nitrate-containing precipitation medium.



**Fig S5.** Scanning Electron micrograph of ZnO particles showing more extensive formation of larger flower-shaped particles by reducing the concentration of precursors and reducing the influence of nitrate counter-ions on material migration during synthesis.

## Video 2

**Video 2.** Evaporation characteristics of sulphate ion containing supernatant on TEM grid in the vicinity of ca. 150 particles. Note the adsorption of the salt residuals to the surfaces of the particles.



**Fig S6.** EDX spectra for chloride- and sulphate-derived ZnO nanoparticles a)ZC-8g b)ZS-8g after UC cleaning. These examples were chosen to emphasize the absence of counter-ions traces on the particle surfaces. The EDX characterization of acetate-derived particles showed no counter-ion removal efficiency because carbon in the acetate group was interfering with carbon from the TEM grid.