

## **Effect of particle size on thermal stability of nanoscale minerals**

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The nanoscale minerals can be formed in diverse geological environments through geochemical, biological and mechanical processes. The probing of the effect of particle size on the dehydration and/or temperature-induced phase transition is essential to understand geological processes in the earth. In this study, the thermal stability of nanoscale minerals including alumina, talc, and zeolites are reported with varying nano-scale particle size. High-resolution solid-state <sup>1</sup>H nuclear magnetic resonance (NMR) spectra provides quantitative changes in the atomic structure of hydrous species on the surface of minerals upon *ex-situ* heating. The water content in nanoscale mineral significantly increases with decreasing particle sizes. With increasing heating temperature, these hydrous species significantly decrease in order of physically adsorbed water, chemisorbed water, and hydroxyl groups. The phase transition also can be observed with dehydration as shown in <sup>27</sup>Al NMR spectra and XRD patterns. The lower dehydration temperature and thermal phase transition temperature are observed for the smaller sized minerals. These results indicate that the dehydration of hydrous species on the surface induces the lower phase transition temperature for nanoscale minerals than macroscale and bulk minerals. Higher water content, lower dehydration temperature and thermal phase transition temperature in nanoscale mineral compared to bulk mineral suggests that nanoscale minerals in fault gouge (e.g., mylonite) may play an important role in reducing the fault friction by retaining hydrous species on the surface.