

THERMAL DECOMPOSITION OF IRON(III) SULFATE

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Influence of conditions (temperature, particle size, diffusion conditions, layer thickness of the precursor and time of calcinations) on solid-state thermal decomposition of iron(III) sulfate in air towards iron(III) oxides and their polymorphous transformations are discussed. Mechanism of thermal decomposition and polymorphous transformations have been investigated mainly using ^{57}Fe Mössbauer spectroscopy, X-ray powder diffraction (XRD), and thermal analysis (DSC, DTA) methods. $\gamma\text{-Fe}_2\text{O}_3$ (maghemite) and $\beta\text{-Fe}_2\text{O}_3$ were identified as primary products of iron(III) sulfate decomposition. Particle size induced polymorphous transformations from maghemite to hematite via $\varepsilon\text{-Fe}_2\text{O}_3$ and simultaneously from $\beta\text{-Fe}_2\text{O}_3$ to hematite has a direct influence on thermal decomposition of residual iron(III) sulfate. This is related to a diffusion of gaseous SO_3 from the bulk of particle through iron(III) oxide surface layer. The overall decomposition mechanism is an unique exhibition of polymorphous transformations of iron(III) oxide.

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