

**The structural and magnetic properties of a  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> nanopowder synthesized by atmospheric microwave torch**

**B. David,<sup>a</sup> O. Schneeweiss,<sup>a</sup> E. Šantavá,<sup>b</sup> and O. Jašek<sup>c</sup>**

<sup>a</sup>Institute of Physics of Materials, AS CR, v.v.i., Brno, Czech Republic

<sup>b</sup>Institute of Physics, AS CR, v.v.i., Prague, Czech Republic

<sup>c</sup>Faculty of Science, Masaryk University, Brno, Czech Republic

For the synthesis of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> nanoparticles a microwave torch discharge ignited in Ar at atmospheric pressure has been used. A double-walled nozzle electrode enabled us to introduce separately the gases: Ar flowed in the central channel whereas the mixture of H<sub>2</sub>/O<sub>2</sub> and Fe(CO)<sub>5</sub> vapour was added into the Ar discharge through the outer channel. The composition and properties of the synthesized nanopowders were studied by TEM, XRD, Raman and Mössbauer spectroscopies. For the magnetic measurements in the range 293–1073 K a vibrating sample magnetometer was employed. Heat capacity and ZFC/FC measurements were performed on a PPMS® device from Quantum Design.

Only the cubic  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> phase with the mean crystallite size of 20 nm was identified by XRD in a representative sample. The total spectrum area (*TSA*) of the Mössbauer transmission spectrum measured on the sample in the range 4–293 K strongly decreased with increasing temperature (*TSA* = 0.155 at 5 K, *TSA* = 0.026 at 293 K). This behaviour is attributed to a bimodal particle size distribution and the chain-like morphology of very small particles (observed under TEM), which enables tilting motions of particles. We also present the high-temperature magnetic properties of the representative sample and describe its structural changes and phase transformations up to 1073 K.

9.7 cm

13.4 cm

**Subject category :**

5. Nano-structure, Surfaces, and Interfaces

**Presentation mode :**

poster

**Corresponding author :**

B. David

**Address for correspondence :**

Institute of Physics of Materials

AS CR, v.v.i.

Zizkova 22

CZ-61662 Brno

Czech Republic

**Email address :**

david@ipm.cz