## NANOCATALYSTS CONTAINING MAGNETITE AND α-IRON FOR LOW-TEMPERATURE OZONE DECOMPOSITION

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Nanostructured compositions containing magnetite (Fe<sub>3</sub>O<sub>4</sub>) and  $\alpha$ -iron ( $\alpha$ -Fe) were obtained by welding of steel using a bare welding wire of Sv08G2S (ER70S-6) type in CO<sub>2</sub> as a shielding gas. The wire had the following composition (wt. %): Fe (97.2), Mn (1.51), Si (0.92), Cu (0.2), and C (0.08). With the help of an electromechanical filter, three fractions of the solid component of welding aerosol were obtained. The first fraction (CD) was picked up from a corona discharge zone, the second fraction (PE) – from a perforated electrode, and the third fraction (GE) – from a grid electrode. The samples of each fraction were characterized by X-ray phase and FT-IR spectroscopy, water vapor adsorption, and pH-metry and also were tested in the reaction of ozone decomposition at the initial ozone concentration,  $C_{O_3}^{in}$ , of 1.0 mg/m³. The phase composition was identical for the three fractions: their diffraction patterns showed peaks at the interplanar spacing, d (Å), of 2.964, 2.528, 2.097, 1.712, 1.614, 1,483, and 1.091 characteristic of magnetite and 2.025, 1.432, and 1.169 characteristic of  $\alpha$ -iron. However, for CR, PE, and GE fraction sequence, Fe<sub>3</sub>O<sub>4</sub> content increased from 61 to 92 wt. % whereas  $\alpha$ -Fe content decreased from 31 to 5 wt. %. In this sequence, sizes of Fe<sub>3</sub>O<sub>4</sub> crystallites varied as 58, 58, and 45 while sizes of  $\alpha$ -Fe varied as 59, 28 and 42.

Figure shows time dependences of a final ozone concentration in the course of ozone decomposition by samples of CD, PE, and GE fractions. All kinetic curves have portions where ozone

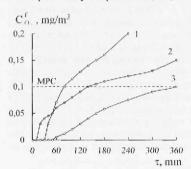


Figure. Time dependences of C for ozone decomposition with samples of different fractions obtained with the help of the electrochemical filter: CD (1), PE (2), and GE (3)

is not detected (a gas analyser sensitivity was 0.025 mg/m³). Then final ozone concentrations go up and become equal to maximum permissible concentration, MPC, in different periods of time,  $\tau_{MPC}$ , depending on the fraction tested.  $\tau_{MPC}$  Characterizes ozone protection intrinsic to the samples and increases in the order CD < GE < PE from 80 to 360 min. A further increase in the final ozone concentration,  $C_{O_3}^f$  is very slow: in 1800 min,  $C_{O_3}^f \approx 0.25$  mg/m³ for all three fractions (not shown in Figure).

It has been found from the experimental data that magnetite phase is responsible for the activity of the two-component, Fe<sub>3</sub>O<sub>4</sub> +  $\alpha$ -Fe, composition.