THEORETICAL ASPECTS OF MODIFICATION MECHANISM FOR SOL-GEL CERAMIC OXIDE POWDERS OF TECHNICAL APPLICATION

Iryna Lutsyuk1, * , Yaroslav Vakhula1

Abstract. Liquid phase technologies for high-grade ceramic oxide powders of technical purpose and methods of their modification have been developed. On the basis of physico-chemical research methods various mechanisms of modifying processes have been analyzed. Substitution of the main ions by doping ones occurs in complex oxide powders followed by the formation of solid solutions. A powder which was obtained via TiO2 modification by sulfur has a surface layer in which sulfur forms chemical bonds.

Keywords: ceramic powders, structural modification, sol-gel technology, solid solutions, ion implantation.

1. Introduction

Modern ceramics with high performance are obtained using special compositions and microstructures, which require precision control at all technological stages: synthesis of powders with certain sizes, processes consolidation and formation, sintering, final heat treatment and control. The properties of technical ceramics depend on the properties, structure, purity, and granulometry of powders.

Ceramic oxide powders are necessary components of many materials, in particular, solid-state lasers, photocatalysts, plastics and heat-resistant concrete, dental composite materials, sorbents, etc. [1-4]. The main characteristics of ceramic powders are the chemical composition, their homogeneity, shape, particles size and granulometry, the degree of particles crystallinity and the concentration of nonequilibrium defects [5-6].

Despite the high functional indices of powders and ceramics in general, often it is expedient to introduce a small amount (even a fraction of one percent) of a certain additive without changing the basic composition of the powder to change its basic parameters radically. The range of technical characteristics of the powders may be extended due to their modification. Such approach allows to change or strengthen the operational properties of the product without changing the main operating conditions.

The choice of the method and degree of modification, as well as the type of modifying agent is an urgent problem from technological and economical viewpoint. It depends on the kind of technology, powder type and ultimate aim.

From kinetics and energy saving standpoint liquid phase technology and sol-gel technology in particular, is the most promising one [7-8]. This technology allows to modify the powder composition and structure at the stage of solutions preparation on the ionic level and to provide the maximum uniform distribution of the modifier in the mixture. Due to the proceeding of sol-gel process and modification at low temperatures the resulting effect is maximum positive.

The aim of this work is to study the mechanisms of modifying the structure of powders in order to change their properties and to obtain materials with given parameters.

2. Experimental

2.1. Powders Synthesis

Nanocrystalline oxide powders of forsterite Mg2SiO4:Me (Me = Cr3+, Y3+, Zr4+), spinel MgAl2O4:Me (Me = Cr3+, Zr4+) and S-TiO2 were synthesized via sol-gel technology.

Synthesis of MgO-SiO2 powders. Magnesium acetate Mg(CH3COO)2·4H2O (P.A. grade) was dissolved in water and ethyl silicate ETS-40 was slowly hydrolyzed in the presence of chloric acid. The obtained solutions were mixed at the definite molar ratio and sol was formed transformed further into a gel. The gel was dried at 383–388 K for 35–40 h. To form the final powder with the given phase composition the gel was treated at 1173 K for 3 h.
Synthesis of MgO-Al2O3 powders. Magnesium acetate Mg(CH3COO)2·4H2O (P.A. grade), aluminum nitrate Al(NO3)3·9H2O (P.A. grade) and citric acid (CA) were dissolved in water and mixed with the molar ratio of $n(Mg^{2+}+Al^{3+})/n(CA) = 1:1$. The solution was evaporated at the temperature about 363 K until a polymer gel was obtained. Then it was burned out at 1273 K for 3 h.

To vary the properties of forsterite and spinel their structures were modified by Cr3+, Y3+ and Zr4+ ions, which were introduced using water-soluble salts Cr(NO3)3·9H2O, Y(NO3)3·6H2O and ZrOCl2·8H2O. All salts were of pure grade.

Synthesis of S-TiO2 powders. To obtain powders of titanium(IV) oxide modified by sulfur we used titanium butoxide Ti(C4N9O)4 (TBO), thiourea (NH2)2CS (TU) and butanol. TBO was previously hydrolyzed in ethanol, butanol and water. TU was dissolved in water. The obtained solutions were mixed at the definite molar ratio until a sol was formed. Then the sol was dried by two stages: I stage – at 393 K to obtain the paste-like mixture; II stage – at 513 K to obtain the powder.

The first stage of sol drying was carried out using three modes:

- mode 1 – natural weathering of the sols ($T = 293 K$, $\tau = 96$ h);
- mode 2 – heat treatment of the sols with simultaneous mixing and heating ($T = 373 K$, $\tau = 24$ h);
- mode 3 – hydrothermal treatment of sols in an autoclave ($T = 383 K$, $\tau = 24$ h).

The dried sols were burned out within the temperature range of 673–973 K for 1 h.

2.2. Investigation Methods

X-ray diffraction analysis (XRD) was carried out using DRON-4 diffractometer with the following mode of operation: Cu Kα radiation; $2\theta = 10^\circ$–$80^\circ$; $U = 34$ kV; $I = 14$ mA. The surface composition was studied by X-ray photoelectron spectroscopy (XPS) using XSAM-800 Kratos spectrophotometer. The surface composition was estimated by the area of obtained bands. The thickness of the analyzed layer was approximately 5 nm.

SEM analysis was carried out using ZEISS EVO 40XVP microscope with the system of X-ray microanalysis INCA Energy. The same system was also used for X-ray emission spectroscopy (XES) to analyze quantitative and qualitative composition of the powders.

The unit cell dimensions, as well as positional and displacement parameters of atoms in the structures were derived by full profile Rietveld refinement technique using software package WinCSD [9].

The particle sizes were determined according to the Scherrer equation.

3. Results and Discussion

MgO-SiO2 powders. The results of XRD analysis confirmed that all pure and modified Mg2SiO4 powders are monophase and are represented by one crystalline phase – forsterite (Fig. 1). Other phases were not observed.

More precise definition of Mg/Cr, Mg/Y and Mg/Zr positions showed that Mg2+ ions are partially substituted by Cr3+, Y3+ and Zr4+ in Mg2 position.

As a result of calculations using WinCSD software the percentage content of chromium, yttrium, and zirconium were found to be 1.18, 0.59 and 2.47 wt % or 0.46, 0.13 and 0.55 at %, respectively. In this case solid solutions are formed corresponding to the formulas: Mg1.97Cr0.03SiO4+δ, Mg1.99Y0.01SiO4+δ and Mg1.96Zr0.04SiO4+δ, respectively.
The size of forsterite powder particles was determined using the expansion of diffraction maxima by the Scherrer method. They were found to be within the range of 35–40 nm.

**MgO-Al₂O₃ powders.** The peculiarity of sols for spinel preparation is the presence of citric acid (CA) as a gelling component in their composition. According to the experimental results the ratio of \( n(Mg^{2+}+Al^{3+}):n(CA) = 1:1 \) was found to be the optimum one to obtain monophase spinel MgAl₂O₄. In such a case we observe the most intensive crystallization and spinel formation begins at lower temperature compared with other ratios. The temperature of 1273 K and isothermal keeping for 3 h were found to be the optimal parameters for the powder synthesis. According to XRD and Rietveld technique data (Table 2) all reflexes at the diffractogram (Fig. 2) correspond to the spinel structure. Other crystalline phases were not observed.

**XRD analyses** the partial introduction of Cr³⁺ ions in both positions (Mg and Al) and formation of the solid solution \( \text{Mg}_{0.96}\text{Cr}_{0.04}\text{Al}_{1.94}\text{Cr}_{0.06}\text{O}_{4+δ} \). The \( \text{Zr}^{4+} \) ion is included in the spinel structure, substituting only Mg²⁺ with formation of the solid solution \( \text{Mg}_{0.98}\text{Zr}_{0.02}\text{Al}_{2}\text{O}_{4+δ} \). The size of the spinel powder particles, calculated according to the Scherrer formula, is within the range of 47–52 nm.

**S-TiO₂ powders.** The results of the research revealed that it is advisable to perform hydrolysis of TBO in butanol, since the solutions obtained are not segregated and remain stable for a long time. According to XRD data the sols dried at 293 and 393 K are close in composition and contain one crystalline compound – thiourea (\( \text{NH}_2\text{CS} \)). The increase in the sol drying temperature to 513 K leads to the disappearance of TU crystals and the appearance of TiO₂, which is crystallized in the form of a significantly deformed anatase-like phase, which is confirmed by the DTA data [10].
XES indicates the presence of sulfur in TiO$_2$ powder in the amount of 0.7 wt %. Thus, we can assume that in the temperature range from 393 to 513 K, the formation of an anatase crystalline lattice and the incorporation of sulfur ions into its structure occurs, which is actually a modification process. It was found that particles of sol based S-TiO$_2$ powder, which was obtained using butanol as a solvent, are characterized by a lower degree of agglomeration and more uniform distribution of particles in size in comparison with ethanol and water.

According to the SEM results, the particles of dried sols have spherical form with an average diameter of 290–300 nm (Fig. 3).

Fig. 3. Micrograph (SEM) of sol sample 
\(T_{\text{drying}} = 393 \text{ K, } \tau = 24 \text{ h}\)

Fig. 4. Micrographs of S-TiO$_2$ powders under different modes: 1 (a); 2 (b) and 3 (c)

Fig. 5. Diffractograms of TiO$_2$ (a) and S-TiO$_2$ (b) powders

**Table 3**

<table>
<thead>
<tr>
<th>Atomic concentration of the element, at %</th>
<th>Ti/S</th>
</tr>
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<tbody>
<tr>
<td><strong>surface layer</strong></td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>28.2</td>
</tr>
<tr>
<td>O</td>
<td>17.9</td>
</tr>
<tr>
<td>S</td>
<td>5.59</td>
</tr>
<tr>
<td><strong>layer at the depth of 1–3 (\mu\m)</strong></td>
<td></td>
</tr>
<tr>
<td>–</td>
<td>66.80</td>
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<td></td>
<td>0.41</td>
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The effect of drying conditions on the value and character of the powder specific surface one can see in Fig. 4. The powders treated under modes 2-3 are characterized by more developed specific surface. The sols which were dried at room temperature (mode 1) are agglomerated.

XPS data confirm the presence of sulfur in the form of MeSSMe and (MeOS) agglomerated. The powders treated under modes 2-3 are characterized by more developed specific surface. The character of the powder specific surface one can see in Fig. 4. The powders treated under modes 2-3 are characterized by more developed specific surface. The surface composition of S-TiO2 powder synthesized at 773 K is represented in Table 3 in accordance with the results of XPS and XES analyses. According to XRD data pure and modified titanium oxides are monophase and contain only one crystalline phase with the anatase structure (Fig. 5). The particles size synthesized at 773 K is represented in Table 3 in accordance with the results of XPS and XES analyses. The highest concentration of sulphur was observed in the form of sulfur-containing particle with the diameter of ~10 nm; the formed globules create nanostructured spheres with an average diameter of ~1 µm. According to XRD data pure and modified titanium oxides are monophase and contain only one crystalline phase with the anatase structure (Fig. 5). The particles size synthesized at 773 K is represented in Table 3 in accordance with the results of XPS and XES analyses. The highest concentration of sulphur was observed in the form of sulfur-containing particle with the diameter of ~10 nm; the formed globules create nanostructured spheres with an average diameter of ~1 µm. Thus, TiO2 is modified by sulphur as follows: sulphur-containing particle with the diameter of ~10 nm segregate to the surface of anatase crystals in the form of nanoaggregates and formation of spheres with the diameter of ~1 µm. Implantation of TiO2 structure by sulfur occurs in surface layers of ~5 nm thickness with the formation of bidentate sulfates.

4. Conclusions

A heterovalent isomorphic substitution of Mg2+ and Al3+ ions by Cr3+, Y3+ and Zr4+ ions was established in the structure of forestite and spinel followed by the formation of solid solutions of Mg1.96Cr0.04Al1.96Cr0.06O4.95, Mg1.98Zr0.02SiO4.95, Mg0.96Cr0.04Al1.96Cr0.06O4.95, and Mg0.98Zr0.02Al2O4.96. The formation of the sol-gel powders morphology takes place in accordance with a multi-level scheme, which involves the addition of sulfur-containing particles to anatase crystals in the form of nanoaggregates and formation of spheres with the diameter of ~1 µm. Implantation of TiO2 structure by sulfur occurs in surface layers of ~5 nm thickness with the formation of bidentate sulfates.

References


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ТЕОРЕТИЧНІ АСПЕКТИ МЕХАНІЗМУ МОДИФІКУВАННЯ КЕРАМІЧНИХ ОКСІДНИХ ПОРОШКІВ ТЕХНІЧНОГО ПРИЗНАННЯ

Анотація. Наведено результати розробленої рідкофазової технології високодисперсних керамічних оксидних порошків технічного призначення та способи їх модифікування. На основі фізико-хімічних методів досліджено механізми модифікуючих процесів. У складних оксидних порошках відбувається заміщення основних іонів гелюючими з утворенням твердих розчинів. Модифікування TiO2 Сульфидром дає змогу отримати порошок із поверхневим шаром, в якому Сульфид утворює хімічні зв'язки.

Ключові слова: керамічні порошки, структурні модифікації, золь-гель технологія, тверді розчини, йонна імплантація.