Self-organized formation of spherical porous granules only by one-step heat-treatment in MgO Fe₂O₃ Nb₂O₅ system

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Self-organized formation of spherical porous granules only by one-step heat-treatment in
MgO-Fe₂O₃-Nb₂O₅ system

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Abstract

In this letter, a new process to obtain spherical porous granules (SPGs) is presented, namely self-organized formation only by a one-step heat-treatment. Commercial MgCO₃ (basic), α-Fe₂O₃ and Nb₂O₅ powders of Mg:Fe:Nb = 1:1:1 in mole fraction were wet-ball milled, and were heat-treated in air at 800-1350°C. The samples heated at ≥1100°C, most of the samples became SPGs with the diameter of ~50-100 µm, containing 3-D network structure. The main constituent phase of porous spheres was pseudobrookite-type MgFeNbO₅. The formation mechanism of porous microspheres was discussed by using SEM observation and high-temperature XRD analysis.

Keywords: porous spheres, granular materials, self-organization, pseudobrookite, MgFeNbO₅

1. Introduction

Porous ceramics are widely used for fluid filters, catalysts, catalyst supports, lightweight structural materials, and so on. Among various shapes of the porous ceramics, spherical porous granules (SPGs) are favorable for heterogeneous catalysts, photonic materials, chromatography supports, drug delivery carriers, and healing of defective bones [1-4]. A variety of processes, mostly liquid-based, have been developed to synthesize SPGs [1, 5-15]. To synthesize SPGs, solid-based processes have seldom been realized, except the mechanical granulation method with rotating fluidized bed [16].

We developed uniformly porous ceramics with 3-D network structure (UPC-3D) by using pyrolytic reactive sintering of powder mixtures including gaseous spices such as carbonates and hydroxides [17,18]. Stimulated by Bayer's [19] early report in 1971 on pseudobrookite-type ceramics (generally expressed as Me₃O₅), uniformly porous MgTi₂O₅ [20-21], and its derivative, bulk MgFeNbO₅ (Fe³⁺ + Nb⁵⁺ substitute 2 Ti⁴⁺) have recently been synthesized [22]. In this previous report on bulk MgFeNbO₅ [22], mold-pressed compacts of mortar-mixed powders of MgCO₃ (basic), α-Fe₂O₃ and Nb₂O₅ were sintered in air to obtain relatively dense materials. However, when the ball-milled powder mixture was simply heated in air, we have found the self-organized formation of

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SPGs, mainly composed of MgFeNbO$_5$ phase. Here we present the new process to synthesize SPGs only by one-step heat-treatment in the MgO-Fe$_2$O$_3$-Nb$_2$O$_5$ system.

2. Experimental

MgCO$_3$ (basic) (Mg$_5$(CO$_3$)$_4$(OH)$_2$•4H$_2$O, 99.9%, Kojundo Chemical Laboratory), $\alpha$-Fe$_2$O$_3$ and Nb$_2$O$_5$ (99.9% each, Wako Pure Chemical Ind.) powders were used as the starting materials, similar to the previous report [22]. The MgCO$_3$ (basic), $\alpha$-Fe$_2$O$_3$ and Nb$_2$O$_5$ powders (Mg:Fe:Nb = 1:1:1 in mole fraction) were wet-ball milled for 24 h in ethanol with ZrO$_2$ balls, dried and sieved through 100 mesh screen (<150 $\mu$m). The mixed powder was heated at the ramp rate of 10°C/min, up to the maximum temperatures from 800 to 1350°C, and then cooled to room temperature without holding time. For the precise temperature control, TG-DTA apparatus is useful, but normal air furnaces can also be used.

The microstructure of the heat-treated products was characterized by scanning electron microscopy (SEM, SU-70, Hitachi High-Tech.) and digital optical microscopy (VHX-1000, Keyence Co.). The reaction behavior of the MgCO$_3$ (basic)-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder (without compressing) was analyzed by high-temperature X-ray diffraction (HT-XRD, Multiflex, Cu-K$_\alpha$, 40 kV and 40 mA, XRD, Rigaku), from room temperature to 1350°C in air. Diffraction patterns were acquired at 100°C each up to 1000°C, and 50°C each up to 1350°C. The ramp rate of the furnace was 10°C/min. XRD patterns were acquired after 5 min holding at each temperature. The scanning rate was $2\theta$=4°/min and the scanning range was $2\theta$=10-60°, hence, 5 min holding + 15 min scanning at each temperature.

3. Results and discussion

Fig. 1 shows representative SEM images of (a) the MgCO$_3$ (basic)-$\alpha$-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder and (b) its heat-treated-only product (SPGs) at 1350°C (ramp speed: 10°C/min, holding time: 0 min), and (c) a digital optical microscope image (real color) of SPGs. The microstructure was drastically changed only by the simple heating. In order to verify the microstructure development during the heating, Fig. 2 shows lower magnification SEM images of the heat-treated products at 800-1300°C. To prepare the SEM samples, heat-treated products were spread on a carbon tape using a spatula, and coated with Pt by sputtering. At 800°C, the surface of fine particles reflected the sample preparation with a spatula (i.e., like spread butter with a butter knife), but special structural feature was not observed. At 900°C, a part of fine particles started to form SPGs with the diameter of ~ 50 $\mu$m. At 1000°C, formation of SPGs became frequent. The spherical granules seem to be soft due to the weak bonding among primary particles, because the surface of some granules was slightly flattened by the sample preparation. At >1100°C, most of the sample became SPGs, with typical diameter of ~50-100 $\mu$m.
Fig. 1

Fig. 2
Figure 3 shows higher magnification SEM images of the heat-treated products from the MgCO$_3$ (basic)-α-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder. At 800°C, the microstructure was similar to that of the as-mixed powder at room temperature (Fig. 1(a)), however, at 900°C, the primary particles start to connect to each other. Somewhat large and rod-like grains observed at 800-1000°C are attributed to the pseudomorphs (remnants of original shape) consisted of fine MgO particles from hydromagnesite. At ≥1100°C, curved and so-called "L-shaped" grains were observed, which are typically found in pseudobrookite-type ceramics with some liquid phase [20-22], and necking between particles became firmer. At 1300°C, although some grains with intermediate clombite-type and spinel-type structures remained at the surface, most of the product converted into pseudobrookite-type MgFeNbO$_5$.

The reaction behavior of the MgCO$_3$ (basic)-α-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder during the heat-treatment was analyzed by HT-XRD as shown in Fig. 4. All peaks before heating were well-indexed with Mg$_5$(CO$_3$)$_4$(OH)$_2$•4H$_2$O (hydromagnesite, ICDD PDF #25-0523), α-Fe$_2$O$_3$ (hematite, #33-0664), and low-temperature phase Nb$_2$O$_5$ (#27-1003). The thermal decomposition of hydromagnesite completed at ~500°C (confirmed by TG-DTA, Fig S1), and MgO nanocrystals was formed from hydromagnesite as shown in Fig. 4. As shown in Fig. 3, MgO nanocrystals kept the pseudomorphs of hydromagnesite at intermediate temperatures (~500-800°C). At ~900-1000°C, the formation of intermediate clombite-type MgNb$_2$O$_6$ (ss) and spinel-type MgFe$_2$O$_4$ (ss) phases started. At ~1100°C, the formation of pseudobrookite-type MgFeNbO$_5$ started. At 1300°C, pseudobrookite-type MgFeNbO$_5$ became preferential, and at 1350°C, the intermediate phases were almost disappeared (corresponding to Fig. 1(b)). After a certain holding time at a high temperature
(e.g., 1350°C for 2 h), the porous spheres started to sinter. Fig. S2 shows a SEM image of the sample after a series of HT-XRD measurements (measurements at each temperature, including final measurement at 1350°C with 2 h holding), corresponding to the sample after cooling in Fig. 4. Spinel grains with octahedral facets were sometimes observed at the surface of semi-sintered porous spheres. That is why spinel 111 peak at 2θ=18.3° is stronger than spinel 311 peak around 35° (usual strongest peak) in the top pattern of Fig. 4.

From the microstructural observation (Figs. 1-3 and S2) and HT-XRD analysis (Fig. 4), the self-organized porous microsphere formation can be attributed to (1) local liquid phase formation by multiphase eutectic reactions at high temperatures and (2) limited sinterability of anisotropic pseudobrookite-type grains. The upper size limitation of the porous spheres might be related to the mesh screen size (<150 µm) used in the powder process, although the agglomerate was very soft as shown in Figs. 1(a) and 2(a). The local liquid-phase formation promoted the granulation and sphericalization of fine particles to reduce the surface energy. In addition, due to the limited sinterability of pseudobrookite phase, it kept the 3-D network structure up to relatively high temperature of 1350°C. Although only a specific system (MgO-Fe2O3-Nb2O5) was presented in this study, similar processing can be extended for the pyrolytic reactive sintering in other systems with satisfying above two conditions. In the future, we will clarify the granulation and sphericalization mechanism in detail by using in situ observation techniques, and will functionalize the microspheres for catalysts and so on.
4. Conclusions

In this report, self-organized formation of spherical porous granules (SPGs) in MgO-Fe₂O₃-Nb₂O₅ system was presented. Only by the heat-treatment of the MgCO₃ (basic)-α-Fe₂O₃-Nb₂O₅ mixed powder, SPGs with 3-D network structure, typically 50-100 µm in diameter, were obtained. The mechanism of porous sphere formation is probably attributable to (1) local liquid phase formation by multiphase eutectic reactions and (2) limited sinterability of anisotropic pseudobrookite-type grains.

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References

Figure Captions

**Fig. 1** SEM images of (a) the MgCO$_3$ (basic)-α-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder and (b) its heat-treated-only product (SPGs) at 1350°C; (c) digital optical microscope image of SPGs.

**Fig. 2** Lower magnification SEM images of the heat-treated products from the MgCO$_3$ (basic)-α-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder.

**Fig. 3** Higher magnification SEM micrographs of the heat-treated products from the MgCO$_3$ (basic)-α-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder.

**Fig. 4** HT-XRD patterns for the MgCO$_3$ (basic)-α-Fe$_2$O$_3$-Nb$_2$O$_5$ mixed powder. Mg : Fe : Nb was 1:1:1 in mole fraction. The top pattern was taken at room temperature after cooling. Patterns at 100°, 200°, 300°, 400°, 1050°, 1150° and 1250°C are not shown here.