

Ce_{0.5}Zr_{0.5}O₂ 固溶体における結晶構造変化, - 高温放射光粉末回折による研究 -
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Crystal structure change in Ce_{0.5}Zr_{0.5}O₂ solid solution, - A high-temperature synchrotron powder diffraction study -

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Introduction

Three-way catalyst has been known to oxidize CO and HC, and to reduce NO_x at the same time. CeO₂-ZrO₂ solid solutions are used as the subcatalysts for purification of automotive exhaust gases. The development of CeO₂-ZrO₂ catalysts requires a better understanding of the crystal structure and structural change. The crystal structure of the CeO₂-ZrO₂ solid solutions has been investigated by Yashima et al. [1-6]. They reported the existence of three metastable tetragonal forms of t, t' and t''. The three tetragonal forms belong to the *P4₂/nmc* space group. However, the crystal change in Ce_{0.5}Zr_{0.5}O₂ at high temperatures has not been investigated *in situ* yet. The purpose of this study is to investigate the structural change in the Ce_{0.5}Zr_{0.5}O₂ solid solution between 1639 K and 1680 K using the synchrotron powder diffraction data.

Experiment

Synchrotron powder diffraction experiment was conducted using a three-axis four circle diffractometer installed at the beam line BL-6C of the Photon Factory, KEK, Japan. Monochromatized 1.55330(7) Å x-ray was used for the diffraction experiment. A furnace with MoSi₂ heaters [7] was attached to a goniometer of the triple-axis/four-circle diffractometer, and used for synchrotron x-ray diffraction measurements at high temperatures. Individual profile fits were performed for the powder data using a profile-fitting program *PRO-FIT* [8].

Results and discussion

Fig. 1 shows the synchrotron x-ray diffraction profiles of the Ce_{0.5}Zr_{0.5}O₂ solid solution measured between 1639 K and 1680 K. All reflections are indexed by a tetragonal cell (*P4₂/nmc*) between 1639 K and 1658 K. The peak splitting between the 004_t and 220_t reflections was clearly observed between 1639 K and 1658 K (Fig. 1). All reflections in the synchrotron x-ray diffraction profile measured between 1664 K and 1680 K are indexed by a cubic fluorite-type cell (*Fm* $\bar{3}$ *m*; Fig. 1). The 400_c reflection exhibits a single feature without splitting between the 004_t and 220_t reflections. The Ce_{0.5}Zr_{0.5}O₂ solid solution was found to transform from the tetragonal t' to cubic phase between 1658 K and 1664 K.

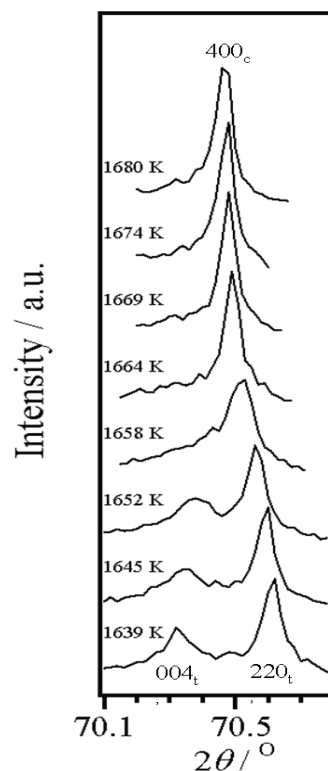


Fig.1 Synchrotron x-ray diffraction profiles from 400 reflection of the Ce_{0.5}Zr_{0.5}O₂ measured from 1639 K to 1680 K.

References

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