ECS Transactions, 25 (2) 2625-2630 (2009) 10.1149/1.3205820 ©The Electrochemical Society

The Electrochemical and Thermal Performances of $Ca_3Co_4O_{9-\delta}$ as a Cathode Material for IT-SOFCs

K. Nagasawa^a, O. Mentre^b, S. Daviero-Minaud^b, N. Preux^b A. Rolle^b and H. Nakatsugawa^a

^a Graduate School of Engineering, Yokohama National University
79-5 Tokiwadai, Hodogaya Word, Yokohama, Kanagawa, 240-8501, Japan
^b UCCS - Unite de Catalyse et de Chimie du Solide-UMR CNRS 8181
Ecole Nationale Superieure de Chimie de Lille
Batiment C7a – BP.90, 108, 59652, Villeneuve d'Ascq cedex, France

With the aim of application for IT-SOFC cathode, two dimensional (2D) misfit layered calcium cobalt oxide $Ca_3Co_4O_{9-\delta}$ was studied using XRD, SEM, thermal dilatometer, and impedance spectroscopy. Chemical reactivity of $Ca_3Co_4O_{9-\delta}$ with electrolyte indicated good compatibility with CGO against generating impunity phase $CaZrO_3$ with YSZ. Thermal expansion coefficient of pure $Ca_3Co_4O_{9-\delta}$ exhibited $10.0-11.0\times10^{-6}$ K⁻¹, which is similar to that of CGO electrolyte. Indeed, we verified a good adherence between electrode thin layer and the electrolyte. Electrochemical property of $Ca_3Co_4O_{9-\delta}$ pellet showed good performances, i.e. ASR values are 3.0 and 24.5 $\Omega \cdot cm^2$ at 700 and 600°C in ambient air respectively, in our experimental conditions.

Introduction

Solid oxide fuel cells (SOFCs) have received considerable attention due to high conversion efficiency in several fuel cell types, without adding of catalytic agent to improve the kinetics. Recently, the intermediate temperature SOFCs (IT-SOFCs) operating at 600-800°C are of interest for improvement of temperature controllability and longer operation time of materials. In the background, it put forward that gadoliniumdoped ceria (CGO) for electrolyte material shows higher ion conductivity at intermediate temperature (under 800°C) in comparison with that of conventional yttria-stabilized zirconia (YSZ) used in the high temperature region (900-1000°C) (1-2). Previous studies on cathode materials for SOFC have already recently focused on perovskite cobalt-based oxide such as La_{1-x}Sr_xMnO₃ (LSM) (3-5), La_{1-x}Sr_xCoO₃ (LSC) (6-8) and Sm_{1-x}Sr_xCoO₃ (SSC) (9-11), showing promising performances. At the same time, it is known that perovskite cobalt oxide has incompatible aspect with electrolytes such as thermal expansion coefficient, and chemical reactivity with zirconia. Layered calcium cobalt oxide $Ca_3Co_4O_{9-\delta}$ is most known for thermoelectric material due to the high figure of merit ZT (ZT= $S^2 \sigma T/\kappa$ where S, σ , T and κ are Seebeck coefficient, electrical conductivity, absolute temperature and thermal conductivity respectively) (12-14). The crystal structure is constituted of stacking rock-salt type [Ca₂CoO₃] layers and CdI₂ type pseudo hexagonal CoO₂ layers alternately. The constitutional formula can be represented $[Ca_2CoO_3]_pCoO_2$ (p value represents b_H/b_{RS} , where b_H and b_{RS} are b axis of pseudo hexagonal and rock-salt layer respectively with p ~0.62) (15-16). Furthermore, the misfit

between each subset along the b axis results in an incommensurate modulated structure between elements of each seat adequately described using the 4D superspace formalism. From the oxygen deficient nature of the rock-salt layers and from the excellent electric transport of similar [CoO₂] layers in superconducting sodium-cobalt bronzes (17), Ca₃Co₄O_{9- δ} could be considered as a natural intergrowth between electronic and potential O²⁻ conducting units. We surmise that this new candidate for IT-SOFC cathode is well from the point of view of electrochemistry, thermal expansion and chemical stability. Those several points constitute the purpose of this study.

Experimental

Polycrystalline Ca₃Co₄O_{9- δ} was synthesized using the conventional solid state reaction. The stoichiometric mixture sample was prepared from reagent grade CaCO₃ and Co₃O₄ powders as precursors and calcined at 880°C for 12 h in air. The obtained sample was pelletized by uniaxial pressing at 10 MPa and sintered at 880°C for 24 h in air. Impedance measurements were performed on symmetrical cells with CGO electrolyte. CGO pellets were formed by uniaxial pressing at 2.5 MPa, and sintered at 1200°C for 2 h in air. Ca₃Co₄O_{9- δ} ink was deposited on both side of the pellet by brush and dried at room temperature. The samples were heat treated in three steps: the step was heated up from room temperature to 500°C during 12 h, next was up to 700°C during 2 h and last was down to room temperature at 6 h. The cylindrical cell after heat treating was 10.57 mm in diameter and 1.73 mm in thickness.

XRD measurements (Siemens, D8) were carried out the powder or pellet samples at the condition of 2θ =6-70° and 0.02° step using Cu-K_a radiation. The lattice parameters refinements were carried out using JANA2006 program (18). Thermal dilatometry was performed for examining thermal expansion coefficient (TEC) of cathode materials. The measurements were carried out the temperature range from 150°C to 820°C. Crosssection scanning electron microscopy (SEM, Hitachi Ltd, S-4700) was performed for observation of the electrode/electrolyte interface, at 15 kV. Impedance spectroscopy measurements were performed using the Solartron SI 1260 analyzer. Here, the cells were slightly pressed between golden plates used as current collectors. Measurement temperature and oxygen partial pressure conditions were carried out from 760 to 540°C at 50-60 K step and from 1 to 3.7×10^{-5} atm respectively. The applied AC-amplitude was 50mV, under zero DC polarization, in a frequency range from 1.0 mHz to 1.0 MHz.

Results and Discussion

The XRD pattern of powder Ca₃Co₄O_{9- δ}, which was refined by Rietveld analysis, is shown in Figure 1. The superspace group was supposed to be X2/m(0,p,0)s0, and cell parameters resulted in high fitting at a ~ 4.8249 Å, b ~ 2.8247 Å, c ~ 10.831 Å, β ~ 98.065° and p ~ 0.6182, which are in good agreement of R_p ~ 3.57% and R_{wp} ~ 5.22 %. About the two unidentified peaks at 2 θ = 12.9 and 18.0°, Grebille *et al.* had mentioned that they correspond to new satellite reflections in RS layer (19). On the other hand, Ling et al. suggested that they might be small-unidentified impurities (20). Previous studies have already shown the incompatibility between zirconia-based electrolyte and perovskite cobaltite/manganite, La_{1-x}Sr_xCoO₃ (LSC) or La_{1-x}Sr_xMnO₃ (LSM), leading to the formation of low conductive La₂Zr₂O₇ or SrZrO₃ by reaction with YSZ (21). Then, we carried out preliminary chemical reactivity tests between layered Ca₃Co₄O_{9- δ} and several electrolytes. Ca₃Co₄O_{9- δ} powders were mixed in 50 mass% with CGO or YSZ, pelletized by uniaxial pressing at 10 MPa and heat-treated for 100 h at 750°C. Figure 2 shows corresponding XRD patterns. In the case of YSZ electrolyte, XRD pattern clearly showed CaZrO₃ and Co₃O₄ impurity peaks in addition to Ca₃Co₄O_{9- δ} and other cobaltites. In other words, the reaction with YSZ electrolyte leads to the decomposition of Ca₃Co₄O_{9- δ}. On the other hand, no impurity phase was observed between Ca₃Co₄O_{9- δ} and CGO electrolyte, confirming a good compatibility between them.



Figure 1. X-ray diffraction patterns of polycrystalline $Ca_3Co_4O_{9-\delta}$ and calculated profile by Rietveld refinement. Filled circle marks indicate unrecognized peaks (see text).



Figure 2. X-ray diffraction pattern in reactivity tests between $Ca_3Co_4O_{9-\delta}$ and YSZ (a) or CGO (b).

After deposition of the electrodes on CGO using the method described in ref. (22), the cross section was analyzed. Then, $Ca_3Co_4O_{9-\delta}$ layer deposited on CGO pellet were fixed by resin after the heat treating and sliced. Typical SEM image of the cross section is shown on Figure 3. It was observed that the adherence between the electrode and electrolyte is good without any evidence for surface cracks. The porosity appears of medium quality. The thickness of cathode cross section is estimated to about 30-40 μ m.

The comparison of TEC between electrode and electrolyte is constitutive attribute along with electrochemical properties. Figure 4 is shown the linear thermal expansion and TEC for $Ca_3Co_4O_{9-\delta}$. It yields TEC values of about $10.0-11.0\times10^{-6}$ K⁻¹ in the investigated region. On the other hand, Du et al. (23) and Kharton et al. (24) mentioned TEC values for CGO electrolyte of 11.1, 11.5 and 12.0×10^{-6} K⁻¹ at 400, 600 and 800°C, respectively. In fact, this apparently good mechanical compatibility is a contributing factor for IT-SOFC cathode application.







Figure 4. Linear thermal expansion and thermal expansion coefficient of Ca₃Co₄O_{9-δ}.

We carried out the impedance spectroscopy measurements for Ca₃Co₄O_{9- δ} electrode deposited CGO electrolyte. Figure 5 shows Nyquist plots in dependence of several oxygen partial pressures or temperatures. The fitting analysis of impedance spectra was used by Zview 3.1 program. Due to the symmetrical cells geometry, the area specific resistance (ASR) was calculated from the total polarization resistance Rp and electrode surface area S using the following: ASR = (Rp*S) / 2. In ambient air, the resulting ASR values are 3.0 and 24.5 $\Omega \cdot cm^2$ at 700 and 600°C respectively. It is remarkable that these resistances values are obtained without any particular effort provided to optimize the microstructure and porosity of the electrode layers. This result indicates that the layered

 $Ca_3Co_4O_{9-\delta}$ has the enough capability to candidate of new cathode material in IT-SOFC. In addition, it was checked by XRD after polarization (Temperature: from room temperature to 800°C; pO₂: from 1 to 10⁻⁵ atm.; total experimental time ~ 1 month) that the electrode remains unchanged.



Figure 5. Oxygen partial pressure (a) or temperature (b) dependence of impedance spectra of $Ca_3Co_4O_{9-\delta}$ deposited CGO pellet.

Conclusions

Electrochemical and thermal properties of layered cobaltite $Ca_3Co_4O_{9-\delta}$ were investigated for applying to cathode material of IT-SOFC. Chemical compatibility with CGO is good, however incompatible with YSZ as well as other perovskite cobaltite. Polarization resistance of $Ca_3Co_4O_{9-\delta}$ cathode showed comparable value to other conventional cathode materials. It was understood that the background of striking impedance spectra was considerable near TEC values between electrode and electrolyte. From a comprehensive standpoint, we revealed that 2D misfit layered calcium cobalt oxide $Ca_3Co_4O_{9-\delta}$ has the possibility of being a cathode material in IT-SOFC.

Acknowledgment

This work was carried out as a collaborative project between Yokohama national university and Ecole nationale superieure de chimie de lille by the program of the consortium japonais du college doctoral franco-japonais (CDFJ) in Ministry of education, culture, sports, science and technology and Embassy of France in Japan.

References

- 1. Z. Tianshu, P. Hing, H. Huang and J. Kilner, Solid State Ionics, 148, 567 (2002).
- 2. B. C. H. Steele, Solid State Ionics, 129, 95 (2000).

- 3. Y. Matsuzaki and I. Yasuda, Solid State Ionics, 126, 307 (1999).
- 4. M. J. Jorgensen, S. Primdahl, C. Bagger and M. Mogensen, *Solid State Ionics*, **139**, 1 (2001).
- 5. S. Kening, P. Jinhua, Z. Naiqing, C. Xinbing, X. Shen and Z. Derui, *Rare Met.*, **27**, 278 (2008).
- 6. H. Uchida, S. Arisaka and M. Watanabe, *Electrochem. Solid-State Lett.*, **2**, 428 (1999).
- 7. H. Kishimoto, N. Sakai, T. Horita, K. Yamaji, M. E. Brito and H. Yokokawa, *Solid State Ionics*, **178**, 1317 (2007).
- 8. P. Hjalmarsson, M. Sogaad and M. Mogensen, Solid State Ionics, 179, 1422 (2008).
- 9. H. Fukunaga, M. Koyama, N. Takahashi, C. Wen and K. Yamada, *Solid State Ionics*, **132**, 279 (2000).
- 10. Z. Tang, Y. Xie, H. Hawthorne and D. Ghosh, J. Power Sources, 157, 385 (2006).
- 11. H. Lu, B. Zhao, Y. Wu, G. Sun, G. Chen and K. Hu, *Mater. Res. Bull.*, **42**, 1999 (2007).
- 12. S. Li, R. Funahashi, I. Matsubara, K. Ueno and H. Yamada, *J. Mater. Chem.*, **9**, 1659 (1999).
- 13. P. Limelette, V. Hardy, P. Auban-Senzier, D. Jerome, D. Flahaut, S. Hebert, R. Fresard and Ch. Simon, *Phys. Rev. B*, **71**, 233108 1 (2005).
- 14. Y. Miyazaki, Solid State Ionics, 172, 463 (2004).
- 15. A. C. Masset, C. Michel, A. Maignan, M. Hervieu, O. Toulemonde, F. Suder, B. Raveau, and J. Hejtmanek, *Phys. Rev. B*, **62**, 166 (2000).
- 16. Y. Miyazaki, M. Onoda, T. Oku, M. Kikuchi, Y. Ishii, Y. Ono, Y. Morii and T. Kajitani, J. Phys. Soc. Jpn., 71, 491 (2002).
- 17. K. Takada, H. Sakurai, E. Takayama-Muromachi, F. Izumi, R. A. Dilanian, and T. Sasaki, *Nature*, **422**, 53 (2003).
- 18. V. Petrcek and M. Dusec, JANA2006, Inst. of Phys. Acad. of Science of the Czech Republik, Prague (2006).
- 19. D. Grebille, S. Lambert, F. Bouree and V. Petricek, J. Appl. Cryst., 37, 823 (2004).
- 20. C. D. Ling, K. Aivazian, S. Schmid and P. Jensen, J. Solid State Chem., 180, 1446 (2007).
- 21. J. M. Ralph, C. Rossignol and R. Kumer, J. Electrochem. Soc., 150, 1518 (2003).
- 22. G. Ehora, S. Daviero-Minaud, M. C. Steil, L. Gengembre, M. Frére, S. Bellayer, and O. Mentré, *Chem. Mater.*, **20**, 7425 (2008).
- 23. Y. Du, N. M. Sammes, G. A. Tompsett, D. Zhang, J. Swan and M. Bowden, J. *Electrochem. Soc.*, **150**, A74 (2003).
- 24. V. V. Kharton, A. V. Kovalevsky, A. P. Viskup, A. L. Shaula, F. M. Figueiredo E. N. Naumovich and F. M. B. Marques, *Solid State Ionics*, **160**, 247 (2003).