Table I. Comparison of measured and calculated (Eq. 5) diffusion coefficients.

Temperature (°C)	D _{eff} (cm²/s)	
	Measured ^a	Calculated ^b
400	1.95 to 4.0×10^{-10}	1.75 to 3.75 × 10 ⁻¹⁰ (7 to 13D)
450 500	4.3 to 8.1 \times 10 ⁻¹⁰ 1.55 to 2.9 \times 10 ⁻⁹	6.0 to 10.8×10^{-10} (5 to 9D) 1.25 to 2.25 $\times 10^{-9}$ (3 to 5D)
300	1.00 10 2.9 \ 10	$1.00 10 2.20 \land 10 (0.10 0D)$

^a Range calculated from data in Fig. 3.

^b Using D from Fig. 4.

$$D_{\rm eff} = D\left(1 + \frac{2k}{d}\right)$$
 [5]

where *k* ranged from 0.015 at 400°C to 0.005 at 500°C. A comparison of the measured coefficients with calculated values of $D_{\rm eff}$ from Eq.5 using d = 25 to 50 μ m (Table I) shows good agreement given the uncertainty and variability in the grain size.

Also, the apparent activation energy for $D_{\rm eff}$ (~85 kJ/mol) agrees closely with the 92.5 kJ/mol value found for grain boundary etch marker movement.¹³

Manuscript submitted Nov. 18, 1996; revised manuscript received Dec. 19, 1996.

Kaiser Aluminum & Chemical Corporation assisted in meeting the publication costs of this article.

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Reduced-Temperature Solid Oxide Fuel Cell Based on YSZ Thin-Film Electrolyte

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ABSTRACT

A planar thin-film solid oxide fuel cell has been fabricated with an inexpensive, scalable, technique involving colloidal deposition of yttria-stabilized zirconia (YSZ) films on porous NiO-YSZ substrates, yielding solid oxide fuel cells capable of exceptional power density at operating temperatures of 700 to 800°C. The thickness of the YSZ film deposited onto the porous substrate is approximately 10 μ m after sintering, and is well bonded to the NiO/YSZ substrate. Ni-YSZ/YSZ/LSM cells built with this technique have exhibited theoretical open-circuit potentials (OCPs), high current densities, and exceptionally good power densities of over 1900 mW/cm² at 800°C. Electrochemical characterization of the cells indicates negligible losses across the Ni-YSZ/YSZ interface and minor polarization of the fuel electrode. Thin-film cells have been tested for long periods of time (over 700 h) and have been thermally cycled from 650 to 800°C while demonstrating excellent stability over time.

Introduction

The high operating temperatures (~1000°C) of solid oxide fuel cells (SOFCs) can lead to complex materials problems which include electrode sintering, inerfacial diffusion between electrolyte and electrode materials, and mechanical stress due to different thermal expansion coefficients of the cell components. Such problems have limited the commercial development of SOFCs. Accordingly, it may be desirable to operate SOFCs at reduced temperatures (within the practical range for methane reforming), assuming this can be done without sacrificing performance. Lower operating temperatures allow a wider choice of materials to be used as interconnects, including metal alloys. Approaches to minimize resistive losses across the electrolyte have included replacing YSZ by alternative electrolyte materials with higher ionic conductivity, and/or reducing the thickness of the solid oxide electrolyte from that in conventional cells (100 to 200 µm) to approximately 10 µm.

Several methods for depositing thin films onto porous subtrates (CVD/EVD, sputtering, sol-gel deposition among others) have been reported.¹⁻⁵ In this paper we describe an inexpensive approach for thin-film deposition using colloidal techniques, and the exceptional performance attained for SOFCs fabricated with this procedure.

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Experimental

Yttria-stabilized zirconia (YSZ) powder from Tosoh Corporation and nickel oxide (NiO) powder from J. T. Baker was mixed approximately as 50/50 weight percent (w/o), respectively. The anode powders were then compacted under uniaxial pressure. Green NiO-YSZ anode disks were subsequently fired at 950°C for 4 h to coarsen the microstructure. YSZ powders mixed in isopropanol were sonicated with a high intensity ultrasonic probe to form a stable colloidal dispersion of submicron polycrystalline electrolye powders. To avoid cracking of the YSZ thin film due to stress built up by the 2D confinement of the film during sintering, the shrinkage of the substrate was carefully matched to the shrinkage profile of the YSZ thin film. In this way, the film is under compression rather than tension while sintering.⁶ The YSZ electrolyte film was applied with a single casting step, and the electrolyte/electrode bilayer was then fired at 1400°C for 4 h to achieve a fully densified YSZ film.

 $La_{0.85}Sr_{0.15}MnO_3$ (LSM) powders were prepared by the glycinenitrate process as described in Ref. 7. LSM was mixed with YSZ in a ratio 50/50 w/o. The LSM/YSZ powder was then mixed with ethylene glycol to form a paste, applied onto an NiO-YSZ/YSZ bilayer, and sintered at 1250°C for ~1 h. The LSM air electrodes were ~100 μm in thickness after sintering.

Platinum mesh was used as the current collector for both the LSM air electrode and the Ni-cermet fuel electrode. Two wires were spot-welded to each current collector so that fuel cell potentials could be measured at the noncurrent carrying leads.



Fig. 1. Fuel cell testing assembly diagram.

Ceramic adhesives were used to fix and seal the fuel cell/current collector structure to an alumina or zirconia tube which was used as a fuel cell testing assembly (Fig. 1). Hydrogen saturated with H₂O at room temperature was used as fuel. Air was used as oxidant and was supplied to the cathode by ambient flow. Since the H₂/H₂O stream was not preheated, there was some cooling of the fuel cell relative to the furnace temperature. A platinum reference electrode was bonded onto the thin electrolyte film, at several millimeters from the working electrode on the air side. Current interrupt and/or impedance spectroscopy was used to measure the ohmic resistance of the fuel cell.

Results and Discussion

The performance of SOFCs, H₂ + 3 volume percent (v/o) H₂O, Ni-YSZ/YSZ/La_{0.85}Sr_{0.15}MnO₃, air, were evaluated under current control. Figure 2 shows current-voltage (*I-V*) curves for a single cell tested at 800°C. The maximum power density achieved was 1935 mW/cm² at a current density of 4.5 A/cm² and cell voltage of 0.43 V (LBNL cells typically exceed 1.8 W/cm² at 800°C). This result illustrates the exceptional performance achievable for YSZ thin-film electrolyte-based SOFCs operating at reduced temperatures. Cells of this type have been operated at power densities of 800 mW/cm² for more than 700 h with no detectable loss in performance.

Figure 3 shows the voltage drop vs. current for a thin-film cell. The total voltage drop was separated into ohmic and nonohmic contributions by the current interrupt method. The use of a reference electrode allowed separation of anodic and cathodic contributions to cell polarization. The YSZ IR drop is estimated from the known conductivity and thickness of the electrolyte (10 μ m). As can be seen from Fig. 3, even at very high current denesities, the voltage drop due to a 10 µm YSZ electrolyte film is almost negligible. The remainder of the ohmic drop is probably due to contact resistance at the LSM/YSZ interface as well as minor losses across the current collectors. The ohmic loss across the Ni-YSZ/YSZ interface is essentially negligible. The polarization of the electrodes (η_a + $\eta_c)$ account for about 40% of the total voltage drop. The cathodic polarization (η_c) was larger than the anodic polarization (η_a) which was almost negligible for current densities below 2.0 A/cm². The YSZ electrolyte IR drop represents about 18% of the total voltage drop.



Fig. 2. *I-V* characteristics of a $(H_2 + v/o H_2O)$, Ni-YSZ/YSZ/LSM, (air) fuel cell at 800°C.



Fig. 3. Separation of cell polarization components by current interrupt techniques; total resistance calculated from the ohmic drop (*IR*) of the cell; η_e , cathode polarization; η_a , anode polarization, and the YSZ electrolyte *IR* drop.

Figures 4 and 5 show the cross sections of the Ni-cermet/ YSZ/Pt and Ni-YSZ/YSZ/LSM fuel cell structures, which have been reduced in hydrogen at 800°C. It is observed that the thin-YSZ electrolyte film is uniformly continuous, pore free, and well adhered to the porous Ni-YSZ fuel electrode. The Ni-cermet is highly uniform with an even distribution of pores and good particle to particle contact. The LSM electrode (Fig. 5) appears to have a less uniform structure when compared to the Ni-cermet which may account for the larger cathodic polarization relative to anodic polarization over most of the *I-V* curve.



Fig. 4. Scanning electron micrograph of a Ni-cermet/YSZ/Pt thin-film fuel cell cross section. YSZ electrolyte film is 9 μ m thick. NiO-YSZ was reduced *in situ* with H₂/H₂O at 800°C to yield final Ni-YSZ cermet structure.



Fig. 5. SEM micrograph showing cross section of a Ni-Cermet/YSZ/LSM thin-film fuel cell that was thermally cycled and tested over 400 h.

Conclusions

A highly successful procedure has been developed which allows deposition of thin-film ceramics on highly porous substrates. The

methodology is inexpensive and scalable. Thin-film SOFCs fabricated using these techniques demonstrate performances of close to 2 W/cm² at 800°C. Current interrupt techniques indicate the majority of the voltage loss at high current density is due to ohmic losses, most likely associated with cathode/electrolyte contact resistance (0.1 Ω cm²). The exceptional performance of the thinfilm SOFCs implies that reduced temperature operation is possible while still maintaining high power density.

Acknowledgment

This research was supported by the Laboratory Technology Research Program (ER-LTR), Office of Computational and Technology Research, U.S. Department of Energy under a CRADA (Cooperative Research and Development Agreement) between Ernest Orlando Lawrence Berkeley National Laboratory (Berkeley Lab) and The Electric Power Research Institute (EPRI), Palo Alto, CA 94303, under U.S. DOE Contract DE-AC03-76SF00098.

Manuscript submitted Sept. 23, 1996; revised manuscript received Oct. 20, 1996.

Lawrence Berkeley National Laboratory assisted in meeting the publication costs of this article.

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Highly Selective Chemical Etching of Si *vs.* Si_{1-x}Ge_x Using NH₄OH Solution

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ABSTRACT

Highly selective chemical etching of Si vs. epitaxial Si_{1-x}Ge_x in NH₄OH solution has been investigated. It was found the selectivity was better than 80:1 even for a Si_{0.9}Ge_{0.1} in 10 weight percent (w/o) NH₄OH at 75°C. As the fraction x of Ge was increased, higher selectivity was obtained due to the decrease of the etch rate of the Si_{1-x}Ge_x. The achievement of the excellent selectivity in a Si/Si_{1-x}Ge_x/Si heterostructure was clearly demonstrated by scanning electron microscopy. Surfaces of etched Si_{1-x}Ge_x samples were analyzed using x-ray photoelectron spectroscopy. The high etch selectivity obtained in NH₄OH is essentially due to a passivation-film effect at the Si_{1-x}Ge_x surface.

Introduction

Selective chemical etching of Si or Si_{1-x}Ge_x has become a key technique in the fabrication of Si_{1-x}Ge_x/Si heterojunction devices^{1,4} and new kinds of structures.⁵⁻⁷ In the fabrication process of the heterojunction device employing thin Si or Si_{1-x}Ge_x layers, for example, it is often necessary to contact to buried Si or Si_{1-x}Ge_x layers. To make contact to buried Si_{1-x}Ge_x layers one needs to etch Si and vice versa. Thin film bond and etchback silicon on insulator (BESOI) of good quality was fabricated using a strained Si₀₋₇Ge₀₋₃ as an etch-stop layer. An important process of the fabrication was to selectively the Si over the Si₀₋₇Ge₀₋₃. ⁵ Recently, we proposed and successfully fabricated Si quantum wires based on selectively removing Si_{1-x}Ge_x from a Si/Si_{1-x}Ge_x trench array.⁷

During the last several years, therefore, the characteristics of several chemical wet etchants for selectively etching $Si_{1-x}Ge_x$

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and/or Si on Si_{1-x}Ge_x/Si heterostructures have been investigated.^{5:11} Among them, two aqueous etchants to selectively etch Si over Si_{1-x}Ge_x were reported. One is the etchant composed of KOH:K₂Cr₂O₇:propanol:H₂O.^{6:10} A selectivity of 40:1 was obtained recently in etching Si over B-doped Si_{0.7}Ge_{0.3}, but it is completely isotropic and requires the use of a hard mask.¹⁰ Another is the mixture of ethylenediamines, pyrocatechol, and water (EPW), which was reported to etch Si over Si/Si_{1-x}Ge_x (x > 0.2) with a high selectivity.¹¹ For a practical application, however, a selective chemical etching should be compatible with silicon integrated circuit (IC) processes. As a result, IC-compatible, nontoxic, anisotropic, simple, and highly selective etchants become important for processes.¹² Koyama *et al.* investigated the etch characteristics of Si_{1-x}Ge_a alloy in an ammoniac wet cleaning solution NH₄OH:H₂O₂:H₂O.¹³ It was observed that the etch rate of Si_{1-x}Ge_x was in the order of