In certain embodiments of the present disclosure, a solid oxide fuel cell is described. The solid oxide fuel cell includes a hierarchically porous cathode support having an impregnated cobaltite cathode deposited thereon, an electrolyte, and an anode support. The anode support includes hydrocarbon oxidation catalyst deposited thereon, wherein the cathode support, electrolyte, and anode support are joined together and wherein the solid oxide fuel cell operates at a temperature of 600° C. or less.

5 Claims, 6 Drawing Sheets
OTHER PUBLICATIONS


Chen et al., “Preparation of Ordered Macroporous Sr$_{0.5}$Sm$_{0.5}$CoO$_2$ as Cathode for Solid Oxide Fuel Cells”, Chemistry Letters, vol. 30, Issue 10, 2001, pp. 1032-1033.


Zhang et al., “Sm$_3$Sr$_{0.7}$CoO$_4$ + Sm$_2$Ce$_{0.9}$O$_{1.9}$ Composite Cathode for Cermet Supported Thin Sm$_2$Ce$_{0.9}$O$_{1.9}$ Electrolyte SOFC Operating Below 600° C.”, Journal of Power Sources, vol. 160, No. 2, 2006, pp. 1211-1216.


* cited by examiner
FIG. 2

(a) cathode
electrolyte
anode

(b) Cathode catalyst (electronic conductor)
Electrolyte (oxide ion conductor)
Anode catalyst (electronic conductor)
CH$_4$ + 4O$^{2-}$ = CO$_2$ + 2H$_2$O + 8e$^-$

**FIG. 3**

**FIG. 4**
FIG. 7

FIG. 8
Table 1. Peak power density and interfacial polarization resistance ($R_p$) of anode-supported SOFCs with different cathode tested at 500°C

<table>
<thead>
<tr>
<th>Power density ($\text{Wcm}^{-2}$)</th>
<th>$R_p$ ($\text{Ωcm}^2$)</th>
<th>Cell components</th>
<th>Cathode</th>
<th>Electrolyte</th>
<th>Anode</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.19</td>
<td>1</td>
<td>Cobaltite-SDC composite</td>
<td>SDC</td>
<td>Ni-SDC</td>
<td></td>
</tr>
<tr>
<td>0.25</td>
<td>0.5</td>
<td>Cobaltite-SDC composite</td>
<td>SDC</td>
<td>Ni-SDC</td>
<td></td>
</tr>
<tr>
<td>0.21</td>
<td>0.79</td>
<td>Cobaltite-SDC composite</td>
<td>SDC</td>
<td>Ni-SDC</td>
<td></td>
</tr>
<tr>
<td>0.44</td>
<td>0.21</td>
<td>Impregnated cobaltite</td>
<td>SDC</td>
<td>Ni-SDC</td>
<td></td>
</tr>
</tbody>
</table>

FIG. 9
ELECTRODE DESIGN FOR LOW TEMPERATURE DIRECT-HYDROCARBON SOLID OXIDE FUEL CELLS

CROSS-REFERENCE TO RELATED APPLICATION

The present application is based on and claims priority to U.S. Provisional Application 61/278,381 having a filing date of Oct. 6, 2009, which is incorporated by reference herein.

GOVERNMENT SUPPORT CLAUSE

This invention was made with government support under DE-FG36-08GO88116 awarded by the U.S. Department of Energy and under award NNX07AL01A awarded by NASA. The government has certain rights in the invention.

BACKGROUND

Solid oxide fuel cells (SOFCs) have the potential to convert the chemical energy in fuels, including hydrocarbon fuels, directly to electricity as schematically shown in FIG. 1. However, the cost of current SOFC systems is still prohibitive for widespread commercial deployment. Reducing the operating temperature of the current SOFC systems to 400-600°C can significantly reduce the cost of the SOFC technology since relatively inexpensive metallic components can be used for interconnects, heat exchangers, and other structural components of the SOFC system. Lowering the operation temperature can also offer quick start-up ability, which is essential for applications such as transportable power sources and auxiliary power units for automobiles. A lower operating temperature will also ensure a greater overall system stability and durability due to a reduction in the thermal stresses in the active ceramic structures, leading to a longer expected lifetime for the SOFC system.

Eliminating the external hydrocarbon reforming system that generates hydrogen, and directly utilizing hydrocarbon fuels on the anode will increase SOFC efficiency while decreasing the complexity and cost of the SOFC system. However, the current state-of-the-art Ni-based anode deactivates rapidly with direct utilization of hydrocarbon fuels due to carbon deposition on the Ni catalyst surface. Further, SOFC performance becomes unacceptably low at reduced temperature with conventional SOFC technology. However, by controlling the microstructure of the electrode, it is possible to achieve high cell performance at reduced operating temperature and to utilize Ni-based anode materials for direct hydrocarbon oxidation.

In view of the above, a need exists for high performance low temperature direct-hydrocarbon SOFCs.

SUMMARY

In the present disclosure, novel electrode engineering is described to realize high performance low temperature direct-hydrocarbon SOFCs. An innovative SOFC architecture combines highly active cobaltite-based mixed ionic and electronic conducting cathodes with carbon deposition mitigated Ni-based anodes. Further, hierarchically porous microstructure can be created to enhance the electrode performance.

In certain embodiments of the present disclosure, a solid oxide fuel cell is described. The solid oxide fuel cell includes a hierarchically porous cathode support having an impregnated cobaltite cathode deposited thereon, an electrolyte, and an anode support. The anode support includes hydrocarbon oxidation catalyst deposited thereon, wherein the cathode support, electrolyte, and anode support are joined together and wherein the solid oxide fuel cell operates at a temperature of 600°C or less.

Other features and aspects of the present disclosure are discussed in greater detail below.

BRIEF DESCRIPTION OF THE DRAWINGS

A full and enabling disclosure, including the best mode thereof, directed to one of ordinary skill in the art, is set forth more particularly in the remainder of the specification, which makes reference to the appended figures in which:

FIG. 1 illustrates principles of SOFC operation in accordance with certain aspects of the present disclosure.
FIG. 2 illustrates a schematic of a novel (a) and a traditional (b) electrode architecture in accordance with certain aspects of the present disclosure.
FIG. 3 illustrates an anode surface in accordance with certain aspects of the present disclosure.
FIG. 4 illustrates schematic illustrations of the self-rising approach in accordance with certain aspects of the present disclosure.
FIG. 5 illustrates a cross-sectional microstructure for cells made from a freeze-drying process in accordance with certain aspects of the present disclosure.
FIG. 6 illustrates a cross-sectional microstructure for cathode with nano-network structure in accordance with certain aspects of the present disclosure.
FIG. 7 illustrates performance for cells with different SDC coating on Ni-SDC anode substrates tested at 600°C in accordance with certain aspects of the present disclosure.
FIG. 8 illustrates SEM at increasing magnification (a), (b) and macro pore size distribution (c) of the LSCF sample calcined at 900°C in accordance with certain aspects of the present disclosure.
FIG. 9 presents a table of values for power density and interfacial polarization resistance (R_p) of anode-supported SOFCs with different cathode tested at 500°C.

DETAILED DESCRIPTION

Reference now will be made in detail to various embodiments of the disclosure, one or more examples of which are set forth below. Each example is provided by way of explanation of the disclosure, not limitation of the disclosure. In fact, it will be apparent to those skilled in the art that various modifications and variations can be made in the present disclosure without departing from the scope or spirit of the disclosure. For instance, features illustrated or described as part of one embodiment, can be used on another embodiment to yield a still further embodiment. Thus, it is intended that the present disclosure covers such modifications and variations as come within the scope of the appended claims and their equivalents.

Among the perovskite-type cathode materials for SOFCs, cobaltite has the highest surface oxygen exchange coefficient, oxide ion conductivity as well as electronic conductivity at reduced operating temperatures. These are the factors favorable for oxygen reduction reaction for a high performance cathode material. In order to overcome the barriers of thermal
expansion mismatch as well as chemical interdiffusion issues, a novel cathode architecture using a composite of the cobaltite catalyst and the electrolyte material is described herein in accordance with the present disclosure. As shown in FIG. 2(a), the electrolyte material is SDC (samarium doped ceria) and the cathode catalyst is LSC (La$_{0.6}$Sr$_{0.4}$CoO$_3$). At low operating temperatures, SDC has high oxide ion conductivity while LSC has the highest electrocatalytic activity for oxygen reduction. A porous SDC layer (cathode frame) is formed on the dense SDC electrolyte. The SDC substrate layer can be formed by tape casting or any other suitable method as would be known in the art. On top of the SDC electrolyte layer, an SDC-NiO anode substrate layer can be formed by tape casting. After high-temperature co-sintering of the triple layers, porous cathode and anode frames are formed on both sides of the dense SDC electrolyte. Nanosized LSC particles are attached on the surface of the porous SDC cathode frame while SDC nanoparticles are attached to the porous Ni-SDC anode substrate. Both the LSC and the SDC nanoparticles are formed by an impregnation method or any other suitable method as would be known in the art. The proposed novel electrode architecture of the present disclosure has many improved characteristics over conventional electrode architecture.

For instance, the electrode architecture of the present disclosure has superior thermal cycling capability. The porous SDC cathode frame, the dense SDC electrolyte and the porous Ni-SDC anode frame are co-sintered to form an integral part and are well matched in thermal expansion, consequently eliminating cracking and delamination between the electrode and the electrolyte. For comparison, the traditional cathode has a clear physical interface between the cathode and the electrolyte, as shown in FIG. 2(b), resulting in significant thermal stresses and causing poor thermal cycling behavior.

The electrode architecture of the present disclosure also has enhanced oxide ion conductivity, SDC is an excellent oxide ion conductor at reduced operating temperatures (400-600°C). The electrode architecture of the present disclosure provides an uninterrupted oxide ion conduction path from the electrode to the electrolyte and will greatly enhance the oxide ion conduction compared with the composite electrode microstructure currently used for SOFCs, as shown in FIG. 2(b).

In addition, the electrode architecture of the present disclosure has high cathode electronic conductivity and electrocatalytic activity. LSC has the highest electronic conductivity among all the reported perovskite-type oxides. The interconnected LSC nanoparticles form an uninterrupted electron conduction path, minimizing the ohmic losses in the cathode layer. LSC has the highest surface oxygen exchange coefficient which promotes oxygen reduction reaction at the cathode. In addition, nanosized LSC particles have large surface areas which further enlarge the oxygen reduction sites and consequently enhance the oxygen reduction process.

Since LSC nanoparticles are prepared on the surface of the SDC cathode frame through impregnation with low firing temperatures (−700°C) and are uniformly attached to the surface of the porous SDC cathode frame and form only a thin layer, there is a large bonding surface and relatively strong bonding strength between LSC nanoparticles and the surface of the porous SDC cathode frame, consequently preventing delamination induced from the thermal stress during thermal cycling.

The electrode architecture of the present disclosure also has superior electronic and ionic conductivity for the anode. Since the anode substrate is Ni-SDC, the electronic conductivity of the proposed anode architecture shown in FIG. 2(a) is high. Further, SDC nanoparticles form a continuous coating layer on the porous Ni-SDC anode frame, providing an interrupted oxide ionic conducting path from the electrolyte to the anode, as shown in FIG. 3.

The porous Ni-SDC anode substrate is coated with SDC nanoparticles, as shown in FIG. 3. SDC has demonstrated catalytic activity for direct oxidation of hydrocarbon fuels. Nanosized SDC will further enhance the direct hydrocarbon oxidation kinetics. Steam generated from direct oxidation of the hydrocarbons will facilitate internal hydrocarbon reforming on the Ni surface. Even when carbon is formed on the Ni surface, SDC in the porous anode composite substrate will remove the carbon deposit since ceria is a good catalyst to remove carbon through the following reaction:

\[
(CrO_2 + C = \frac{1}{2}Cr_2O_3 + CO)
\]

The present disclosure can be better understood with reference to the following examples.

EXAMPLES

Novel Self-rising Approach to Prepare Hierarchically Porous Electrode

Hierarchically porous materials displaying multiple length scales in pore size are desirable for their improved flow performance coupled with high surface areas. Macropores (pore size>50 nm) provide channels for gas and mass transport while mesopores (pore size in the range of 2-50 nm) provide large surface areas for reaction and reaction. Macropores can be prepared by templating methods such as silica or polystyrene beads (see, e.g., Kim, S. W.; Kim, M.; Lee, W. Y.; Hyeon, T. “Fabrication of Hollow Palladium Spheres and Their Successful Application to the Recyclable Heterogeneous Catalyst for Suzuki Coupling Reactions”, Journal of the American Chemical Society 2002, 124, 7642 and Chen, F.; Xia, C.; Liu, M. “Preparation of Ordered Macroporous Sr$_{0.6}$Sm$_{0.4}$CoO$_3$ as Cathode for Solid Oxide Fuel Cells”, Chemistry Letters 2001, 10, 1032, both incorporated by reference herein). However, the limitations of such methods are obvious: “hard templates” which require complicated preparation sequences are always needed, thus significantly increasing the cost for producing hierarchically porous materials at large scale. A novel self-rising approach to make hierarchically porous material has been developed as illustrated in FIG. 4. The method of the present disclosure utilizes a similar mechanism to that of self-rising flours used in a bakery. Urea has been chosen as the leavening agent to obtain macropores because urea will decompose to only gases (NH$_3$ and CO$_2$), consequently avoiding impurities and contamination of the final product. The preparation of mesopores is achieved through the vaporization-induced self-assembly (EISA) method (see, e.g., Brinker, C. J.; Lu, Y. F.; Sellinger, A.; Fan, H. Y. “Evaporation-Induced Self-Assembly: Nanostructures Made Easy”, Advanced Materials 1999, 11, 579 and Liu, Q.; Chen, F.; Song, W. “Synthesis, Characterization and Application of Nanostructured Porous Metal Oxides”, ECS Transactions 2009, both incorporated by reference herein). In the EISA method, during solvent evaporation, the self-assembly process is triggered when the concentration of
the surfactant in the solution begins to exceed the critical micelle concentration. Upon high temperature calcination, the template is removed and the ordered mesoporous materials can be obtained.

Preparation of the Electrolyte/electrode Frame

Electrode substrate supported cells are first fabricated using a freeze-drying approach. Although the freeze-drying process has long been used for ceramic processing, its application in SOFC fabrication is relatively new (see, e.g., Cable, T. L.; Sofie, S. W. “A Symmetrical, Planar SOFC Design for NASA’s High Specific Power Density Requirements”, Journal of Power Sources 2007, 174, 221, incorporated by reference herein). The porous electrode support is fabricated using the freeze-tape casting process while the electrolyte is screen-printed on the electrode support. The electrode/electrolyte assembly is co-sintered. The freeze-drying method renders better control of the porous electrode frame microstructure, as shown in FIG. 5. The SDC electrolyte film is about 10 µm thick. Porous electrode frame is formed on both sides of the electrolyte layer, and the electrode frame and the electrolyte are an integral part.

The microstructural characteristics of the electrode substrate layer, including the thickness, porosity, surface area, pore size and pore size distribution are correlated with the particle size of the starting SDC and NIO powders, organic additives, the drying and heat-treatment processes. The electrode substrate frame is the platform where the electrode catalyst is deposited via impregnation. Therefore, the microstructure of the electrode substrate frame has direct influences on the effective catalyst loading and consequently the electrocatalytic performance of the electrode.

Preliminary Results

Shown in FIG. 6 is an SEM image of cobaltite nanoparticles on the surface of a porous SDC cathode frame. The cathode catalyst precursor is deposited onto the surface of the porous SDC cathode frame by impregnating porous SDC cathode frame with the respective metal nitrate solution, drying, and subsequent firing at about 700°C. The nano-network structure is formed by heating the impregnated precursors. The nano-sized cobaltite particles are well-connected, forming straight conducting paths for oxygen ion and electron conduction. In addition, the nano-network has high porosity for oxygen transport. The performance of a single anode-supported SOFC using the impregnated cobaltite cathode deposited on the surface of porous SDC cathode frame (see, e.g., Zhao, F.; Wang, Z.; Liu, M.; Zhang, L.; Xiu, C.; Chen, F. “Novel Nano-network Cathodes for Solid Oxide Fuel Cells”, Journal of Power Sources 2008, 185, 13, incorporated by reference herein), thin film SDC electrolyte and Ni-SDC anode tested at 500°C, with humidified hydrogen as fuel is shown in Table 1 (FIG. 9), along with other anode-supported cells using the conventional cobaltite-electrolyte composite cathode (see, e.g., Xia, C.; Chen, F.; Liu, M. “Reduced-Temperature SOFC systems using Ceria-Based Electrolytes Fabricated by Screen Printing”, Electrochemical and Solid-State Letters 2001, 4, A52, Zhang, X.; Robertson, M.; Yick, S.; De Petit, C.; Styles, E.; Qu, W.; Xie, Y. S.; Hui, R.; Roller, J.; Kesler, O.; Marie, R.; Ghosh, D. “SmO2CeO2FeO3 Nano-catalyst aggregates”, Journal of Physical Chemistry 2009, 113, 17262, incorporated by reference herein). Using the self-rising approach, hierarchically porous LSCF (La0.8Sr0.2CoO3-δ) has been obtained, as shown in FIG. 8(a), FIG. 8(b) revealing a three dimensional honeycomb like morphology. Large fractions of the sample have highly ordered porous structures in three dimensions over a range of hundreds of micrometers. X-ray diffraction (XRD) results indicate that the LSCF is highly crystalline and the energy dispersive X-ray (EDX) analysis confirms its composition. One significant feature of this self-rising approach is that the high temperature calcination does not destroy the hierarchically porous structure. This is quite different from previous reports in which, even after calcinations at 500°C, macroporous metal oxide started to collapse when using a hard template. The larger and thicker channel walls in LSCF most likely lead to enhanced thermal stability of the macropores. The average macropore diameter of LSCF (FIG. 8c) is about 5 µm. TEM image analysis shows that mesopores with average pore size of a few nanometers are formed on the walls of the macropores.

In the interests of brevity and conciseness, any ranges of values set forth in this specification are to be construed as written description support for claims reciting any sub-ranges having endpoints which are whole number values within the specified range in question. By way of a hypothetical illustrative example, a disclosure in this specification of a range of 1-5 shall be considered to support claims to any of the following sub-ranges: 1-4; 1-3; 1-2; 2-5; 2-4; 2-3; 3-5; 3-4; and 4-5.

These and other modifications and variations to the present disclosure can be practiced by those of ordinary skill in the art, without departing from the spirit and scope of the present disclosure, which is more particularly set forth in the appended claims. In addition, it should be understood that aspects of the various embodiments can be interchanged both in whole or in part. Furthermore, those of ordinary skill in the art will appreciate that the foregoing description is by way of example only, and is not intended to limit the disclosure.
What is claimed is:

1. A solid oxide fuel cell comprising:
   A cathode comprising a porous cathode support, the porous cathode support comprising samaria doped ceria (SDC) with an impregnated cobaltite active material deposited thereon, the cobaltite active material comprising $\text{La}_0.6 \text{Sr}_0.4 \text{CoO}_3$, the porous cathode support comprising macropores and mesopores, each macropore having a pore size of greater than 50 nm and 10 micrometers or less, and each mesopore having a pore size from 2 to 50 nm;
   an electrolyte comprising samaria doped ceria (SDC); and
   an anode comprising an anode support, the anode support comprising nickel-samaria doped ceria (Ni-SDC) with samaria doped ceria (SDC) nanoparticles deposited thereon, the SDC nanoparticles catalyzing oxidation of a hydrocarbon fuel, wherein the cathode support, electrolyte, and anode support are joined together and form an integral component and wherein the solid oxide fuel cell operates a temperature of between 400° C. and 600° C.

2. The solid oxide fuel cell of claim 1, wherein the solid oxide fuel cell operates at a temperature of 550° C. or less.

3. The solid oxide fuel cell of claim 1, wherein the solid oxide fuel cell operates at a temperature of 500° C. or less.

4. The solid oxide fuel cell of claim 1, wherein the solid oxide fuel cell operates at a temperature of 450° C. or less.

5. A method for forming a solid oxide fuel cell comprising:
   forming a cathode, the cathode comprising a porous cathode support comprising samaria doped ceria (SDC);
   depositing a cobaltite active material on the cathode support such that the cobaltite active material is impregnated on the cathode support, the cobaltite active material comprising $\text{La}_0.6 \text{Sr}_0.4 \text{CoO}_3$, the porous cathode support comprising macropores and mesopores, each macropore having a pore size of greater than 50 nm and 10 micrometers or less, and each mesopore having a pore size from 2 to 50 nm;
   forming an anode comprising an anode support, the anode support comprising nickel-samaria doped ceria (Ni-SDC);
   depositing samaria doped ceria (SDC) nanoparticles on the anode support, the SDC nanoparticles catalyzing oxidation of a hydrocarbon fuel;
   co-sintering the cathode support and the anode support with an electrolyte therebetween to form an integral component, wherein the electrolyte comprises samaria doped ceria (SDC), wherein the solid oxide fuel cell is capable of operating at a temperature of between 400° C. and 600° C.