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METHODS OF IMPROVING AN INTERFACE BETWEEN AN ELECTRODE AND AN ELECTROLYTE OF AN ELECTROCHEMICAL CELL, AND RELATED APPARATUSES, AND SYSTEMS

Abstract

A method of improving an interface between an electrode and an electrolyte of an electrochemical cell is disclosed. The method includes forming an electrolyte material on an electrode of an electrochemical cell. The electrolyte may include a perovskite material. The electrolyte material may be exposed to one or more of an acid solution, a plasma, thermal shock, and gamma radiation to increase the surface roughness of the electrolyte material. Additional methods, electrochemical cells, and systems are disclosed.

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Background/Summary

CROSS-REFERENCE TO RELATED APPLICATIONS [0001] This application is a national phase entry under 35 U.S.C. § 371 of International Patent Application PCT/US2023/060386, filed Jan. 10, 2023, designating the United States of America and published as International Patent Publication WO2023/133587 A2 on Jul. 13, 2023, which claims the benefit under Article 8 of the Patent Cooperation Treaty to U.S. Patent Application Ser. No. 63/298,084, filed Jan. 10, 2022.

TECHNICAL FIELD

[0003] Embodiments of the disclosure generally relate to electrochemical cells (e.g., fuel cells, electrolysis cells). In particular, embodiments of the disclosure relate to surface treatments for electrolytes in the electrochemical cells (e.g., fuel cells, electrolysis cells).

BACKGROUND

[0004] Protonic ceramic electrochemical cells (e.g., proton ceramic fuel cells (PCFCs), proton ceramic electrolysis cells (PCECs)) are promising for fuel and hydrogen (H.sub.2) production and power generation at intermediate temperatures (e.g., temperatures within a range of from about 400° C. to about 600° C.). Conventional PCFCs/PCECs include a perovskite-structure oxide electrolyte (e.g., a yttrium- and ytterbium-doped barium-zirconate-cerate (BZCYYb), a yttrium-doped barium-zirconate (BZY)). The conventional perovskite-structure oxide electrolytes exhibit a high bulk proton conductivity that enables lower temperature operations than those for oxygen ion conductor-based solid oxide fuel/electrolysis cells (SOFCs/SOECs) due to smaller activation energy. However, there remain electrolyte-related challenges that limit PCFC/PCEC applications.

[0005] Even though the electrolyte exhibits high proton conductivity (e.g., greater than 10 mS cm.sup.-1 at 500° C.), an observed ohmic loss in the electrochemical cells is much larger than a theoretical value estimated from bulk proton conductivity alone, and observed bulk proton conductivity in the electrochemical cell is much lower than a theoretical value bulk proton conductivity. In addition, an interface between an oxygen electrode and the electrolyte of the electrochemical cells is mechanically weak, which causes delamination and other forms of degradation, resulting in poor electrochemical performance of the electrochemical cell, especially under high-current-density PCEC operation conditions.

BRIEF SUMMARY

[0006] This summary does not identify key features or essential features of the claimed subject matter, nor does it limit the scope of the claimed subject matter.

[0007] Accordingly, in some embodiments, a method of improving an interface between an electrode and an electrolyte of an electrochemical cell is disclosed. The method comprises forming an electrolyte material on an electrode of an electrochemical cell. The electrolyte comprises a perovskite material. The electrolyte material is exposed to one or more of an acid solution, a plasma, thermal shock, and gamma radiation to increase a surface roughness of the electrolyte material.

[0008] Accordingly, in some embodiments, a method of forming an electrochemical cell is

disclosed. The method comprises forming an electrolyte material on a first electrode of an electrochemical cell, the electrolyte material comprising a perovskite material. The electrolyte material is exposed to at least one acid solution to increase a surface roughness of the electrolyte material. After exposing the electrolyte material to the at least one acid solution, a second electrode is formed on the electrolyte material.

[0009] Accordingly, in some embodiments, an electrochemical cell is disclosed. The electrochemical cell comprises a first electrode, a second electrode, and a proton-conducting membrane between the first electrode and the second electrode. The proton-conducting membrane comprises a perovskite material. An interface between the proton-conducting membrane and the second electrode exhibits a peeling strength within a range of from about 17 N to about 40 N.

[0010] Accordingly, in some embodiments, a system for H₂ gas production and electricity generation is disclosed. The system comprises at least one steam source, at least one electrochemical apparatus in fluid communication with the at least one steam source, and a power source electrically connected to the at least one electrochemical apparatus. The at least one electrochemical apparatus comprises one or more electrochemical cells.

Description

BRIEF DESCRIPTION OF THE DRAWINGS

[0011] The illustrations presented in this disclosure are not meant to be actual views of any particular electrochemical cell or component thereof, but are merely idealized representations employed to describe illustrative embodiments. Thus, the drawings are not necessarily to scale.

[0012] While this disclosure concludes with claims particularly pointing out and distinctly claiming specific embodiments, various features and advantages of embodiments within the scope of this disclosure may be more readily ascertained from the following description when read in conjunction with the accompanying drawings, in which:

[0013] FIG. 1 is a simplified cross-sectional view of an electrochemical cell, according to embodiments of the disclosure;

[0014] FIG. 2 is a simplified cross-sectional view that illustrates a method of forming the electrochemical cell **100** of FIG. 1, according to embodiments of the disclosure;

[0015] FIG. 3 is a simplified schematic view of a system for H₂ gas production and electricity generation including the electrochemical cell of FIG. 1, according to embodiments of the disclosure;

[0016] FIG. 4 includes atomic force microscopy (AFM) scans of untreated and treated electrolyte surfaces, as described in Example 2;

[0017] FIG. 5A is a graphical representation of results associated with electron impedance spectroscopy (EIS) measurements, as described in Example 4;

[0018] FIG. 5B is a graphical representation of an Arrhenius plot of ohmic resistance associated with the EIS measurements of FIG. 5A, as described in Example 4;

[0019] FIG. 5C is a graphical representation of an Arrhenius plot of polarization resistance associated with the EIS measurements of FIG. 5A, as described in Example 4;

[0020] FIG. 5D is a graphical representation of calculated activation energies associated with the EIS measurements of FIG. 5A, as described in Example 4;

[0021] FIG. 5E is a graphical representation of a correlation between enhanced kinetics associated with the EIS measurements of FIG. 5A, as described in Example 4;

[0022] FIG. 5F is a graphical representation of an Arrhenius plot correlating a relative “resistance” and the inverse of a reduced temperature associated with the EIS measurements of FIG. 5A, as described in Example 4;

[0023] FIG. 6 is a graphical representation showing current density results (e.g., polarization

curves) of electrochemical cells operated as electrolysis cells, as described in Example 5;

[0024] FIG. 7A is a graphical representation showing Faradic efficiency of electrochemical cells operated as electrolysis cells, as described in Example 5;

[0025] FIG. 7B is a graphical representation showing hydrogen production rate of electrochemical cells operated as electrolysis cells, as described in Example 5;

[0026] FIG. 8 is a graphical representation showing current density results (e.g., polarization curves) and power density results of electrochemical cells operated as fuel cells, as described in Example 5;

[0027] FIG. 9 is a graphical representation showing degradation results of electrochemical cells continuously operated as electrolysis cells for 200 hours, as described in Example 6;

[0028] FIG. 10A is a graphical representation showing voltage and power density curves for an untreated electrochemical cell and a treated electrochemical cell including an LSCF oxygen electrode, as described in Example 8;

[0029] FIG. 10B is a graphical representation showing voltage and power density curves for an untreated electrochemical cell and a treated electrochemical cell including a PBSCF oxygen electrode, as described in Example 8;

[0030] FIG. 10C is a graphical representation showing voltage and power density curves for an untreated electrochemical cell and a treated electrochemical cell including a PNC55 oxygen electrode, as described in Example 8;

[0031] FIG. 10D is a graphical representation showing voltage and power density curves for an untreated electrochemical cell and a treated electrochemical cell including a PNC73 oxygen electrode, as described in Example 8;

[0032] FIG. 10E is a graphical representation showing voltage and power density curves for an untreated electrochemical cell and a treated electrochemical cell including a 3D PNC73 mesh oxygen electrode, as described in Example 8;

[0033] FIG. 11A is a graphical representation comparing peak power density of untreated electrochemical cells and treated electrochemical cells, as described in Example 8;

[0034] FIG. 11B is a graphical representation comparing current density of untreated electrochemical cells and treated electrochemical cells, as described in Example 8;

[0035] FIG. 12A is a graphical representation of gas chromatography results for three reference gases (pure H₂, pure air, and an H₂+air mixture), as described in Example 9;

[0036] FIG. 12B is a graphical representation of gas chromatography results for an untreated electrochemical cell, as described in Example 9; and

[0037] FIG. 12C is a graphical representation of gas chromatography results for a treated electrochemical cell, as described in Example 9.

DETAILED DESCRIPTION

[0038] In the following detailed description, reference is made to the accompanying drawings, which form a part hereof, and in which are shown, by way of illustration, specific examples of embodiments in which the present disclosure may be practiced. These embodiments are described in sufficient detail to enable a person of ordinary skill in the art to practice the present disclosure. However, other embodiments enabled herein may be utilized, and structural, material, and process changes may be made without departing from the scope of the disclosure.

[0039] The illustrations presented herein are not meant to be actual views of any particular method, system, device, or structure, but are merely idealized representations that are employed to describe the embodiments of the present disclosure. In some instances, similar structures or components in the various drawings may retain the same or similar numbering for the convenience of the reader; however, the similarity in numbering does not necessarily mean that the structures or components are identical in size, composition, configuration, or any other property.

[0040] It will be readily understood that the components of the embodiments as generally described herein and illustrated in the drawings could be arranged and designed in a wide variety of different

configurations. Thus, the following description of various embodiments is not intended to limit the scope of the present disclosure, but is merely representative of various embodiments. While the various aspects of the embodiments may be presented in the drawings, the drawings are not necessarily drawn to scale unless specifically indicated.

[0041] The following description may include examples to help enable one of ordinary skill in the art to practice the disclosed embodiments. The use of the terms “exemplary,” “by example,” and “for example,” means that the related description is explanatory, and though the scope of the disclosure is intended to encompass the examples and legal equivalents, the use of such terms is not intended to limit the scope of an embodiment or this disclosure to the specified components, steps, features, functions, or the like.

[0042] As used herein, the term “negative electrode” means and includes an electrode having a relatively lower electrode potential in an electrochemical cell (i.e., lower than the electrode potential in a positive electrode therein). Conversely, as used herein, the term “positive electrode” means and includes an electrode having a relatively higher electrode potential in an electrochemical cell (i.e., higher than the electrode potential in a negative electrode therein).

[0043] As used herein, the term “electrolyte” means and includes an ionic conductor, which can be in a solid state, a liquid state, or a gas state (e.g., plasma).

[0044] As used herein, the term “compatible” means that a material does not undesirably react, decompose, or absorb another material, and also that the material does not undesirably impair the chemical and/or mechanical properties of the another material.

[0045] As used herein, the term “triple conducting perovskite” means and includes a perovskite formulated to conduct hydrogen ions ($H_{sup.}^{+}$) (e.g., protons), oxygen ions ($O_{sub.}^{2-}$), and electrons ($e_{sup.}^{-}$). A triple conducting perovskite exhibits a cubic lattice structure, with the general formula $ABO_{3-\delta}$, where A consists of one or more lanthanide elements (e.g., lanthanum (La), cerium (Ce), praseodymium (Pr), neodymium (Nd), promethium (Pm), samarium (Sm), europium (Eu), gadolinium (Gd), terbium (Tb), dysprosium (Dy), holmium (Ho), erbium (Er), thulium (Tm), ytterbium (Yb), lutetium (Lu)), B consists of cobalt (Co) and one or more of nickel (Ni), manganese (Mn), and iron (Fe), and δ is the oxygen deficit.

[0046] As used herein, spatially relative terms, such as “adjacent,” “beneath,” “below,” “lower,” “bottom,” “above,” “upper,” “top,” “front,” “rear,” “left,” “right,” and the like, may be used for ease of description to describe one element's or feature's relationship to another element(s) or feature(s) as illustrated in the figures. Unless otherwise specified, the spatially relative terms are intended to encompass different orientations of the materials in addition to the orientation depicted in the figures. For example, if materials in the figures are inverted, elements described as “below” or “beneath” or “under” or “on bottom of” other elements or features would then be oriented “above” or “on top of” the other elements or features. Thus, the term “below” can encompass both an orientation of above and below, depending on the context in which the term is used, which will be evident to one of ordinary skill in the art. The materials may be otherwise oriented (e.g., rotated 90 degrees, inverted, flipped) and the spatially relative descriptors used herein interpreted accordingly.

[0047] As used herein, the singular forms “a,” “an,” and “the” are intended to include the plural forms as well, unless the context clearly indicates otherwise.

[0048] As used herein, “and/or” includes any and all combinations of one or more of the associated listed items.

[0049] As used herein, the term “configured” refers to a size, shape, material composition, material distribution, orientation, and arrangement of one or more of at least one structure and at least one apparatus facilitating operation of one or more of the structures and the apparatus in a pre-determined way.

[0050] As used herein, the term “substantially” in reference to a given parameter, property, or condition means and includes to a degree that one of ordinary skill in the art would understand that

the given parameter, property, or condition is met with a degree of variance, such as within acceptable manufacturing tolerances. By way of example, depending on the particular parameter, property, or condition that is substantially met, the parameter, property, or condition may be at least 90.0% met, at least 95.0% met, at least 99.0% met, at least 99.9% met, or even 100.0% met.

[0051] As used herein, “about” or “approximately” in reference to a numerical value for a particular parameter is inclusive of the numerical value and a degree of variance from the numerical value that one of ordinary skill in the art would understand is within acceptable tolerances for the particular parameter. For example, “about” or “approximately” in reference to a numerical value may include additional numerical values within a range of from 90.0 percent to 110.0 percent of the numerical value, such as within a range of from 95.0 percent to 105.0 percent of the numerical value, within a range of from 97.5 percent to 102.5 percent of the numerical value, within a range of from 99.0 percent to 101.0 percent of the numerical value, within a range of from 99.5 percent to 100.5 percent of the numerical value, or within a range of from 99.9 percent to 100.1 percent of the numerical value.

[0052] FIG. 1 illustrates a cross-sectional view of an electrochemical cell **100**, according to embodiments of this disclosure. As shown in FIG. 1, the electrochemical cell includes a hydrogen electrode **102** (e.g., a H₂ gas side electrode), an oxygen electrode **104** (e.g., a steam side electrode), and an electrolyte **106** (e.g., a proton-conducting electrolyte, a proton-conducting membrane) disposed between the hydrogen electrode **102** and the oxygen electrode **104**. The hydrogen electrode **102** and the electrolyte **106** may collectively form a hydrogen electrode-electrolyte bi-layer **108**. In some embodiments, the electrochemical cell **100** is a protonic ceramic electrochemical cell. The electrochemical cell **100** may operate as an electrolysis cell to produce H₂ gas from steam or may operate in reverse as a fuel cell to generate electricity from H₂ gas (e.g., at least a portion of the H₂ gas produced when the electrochemical cell **100** is operated as an electrolysis cell). The electrochemical cell **100** may operate at an operational temperature within a range of from about 400° C. to about 600° C. The electrochemical cell **100** may operate at current densities greater than or equal to about 0.1 amperes per square centimeter (A/cm²), such as greater than or equal to about 0.5 A/cm², greater than or equal to about 1.0 A/cm², greater than or equal to about 2.0 A/cm², greater than or equal to about 3.0 A/cm², or greater than equal to about 3.5 A/cm².

[0053] The electrolyte **106** may be a proton-conducting membrane. The electrolyte **106** may be formed of and include at least one electrolyte material exhibiting an ionic conductivity (e.g., H⁺ conductivity) greater than or equal to about 10⁻² S/cm, such as within a range of from about 10⁻² S/cm to about 1 S/cm, at one or more temperatures within a range of from about 150° C. to about 650° C., such as from about 200° C. to about 600° C. The electrolyte material may be formulated to remain substantially adhered (e.g., laminated) to the hydrogen electrode **102** and the oxygen electrode **104** at relatively high current densities, such as at current densities greater than or equal to about 0.1 A/cm² (e.g., greater than or equal to about 0.5 A/cm², greater than or equal to about 1.0 A/cm², greater than or equal to about 2.0 A/cm², greater than or equal to about 3.0 A/cm², greater than equal to about 3.5 A/cm²). In some embodiments, the electrolyte **106** is formed of and includes at least one perovskite material having an operational temperature (e.g., a temperature at which the conductivity of the perovskite material is greater than or equal to about 10⁻² S/cm, such as within a range of from about 10⁻² S/cm to about 1 S/cm) within a range of from about 350° C. to about 650° C. The electrolyte **106** may be formed of and include a perovskite material exhibiting a cubic lattice structure with a general formula ABO_{3-δ}, where A may comprise barium (Ba), B may comprise one or more of zirconium (Zr), cerium (Ce), yttrium (Y), and ytterbium (Yb), and δ is the oxygen deficit. By way of non-limiting example, the electrolyte **106** may be formed of and include one or more of a yttrium- and ytterbium-doped barium-zirconate-cerate (BZCYYb), such as BaZr_{0.8-y}Ce_yY_{0.2-x}Yb_xO_{3-δ}, where x and y are dopant levels and δ is the oxygen deficit (e.g.,

BaZr.sub.0.1Ce.sub.0.7Y.sub.0.1Yb.sub.0.1O.sub.3- δ (BZCYYb1711), BaZr.sub.0.4Ce.sub.0.4Y.sub.0.1Yb.sub.0.1O.sub.3- δ (BZCYYb4411), BaZr.sub.0.3Ce.sub.0.5Y.sub.0.1Yb.sub.0.1O.sub.3- δ (BZCYYb3511)), doped barium-zirconate (BaZrO.sub.3) (e.g., yttrium-doped BaZrO.sub.3 (BZY), such as BaZr.sub.0.8Y.sub.0.2O.sub.3- δ where δ is the oxygen deficit), barium-yttrium-stannate (Ba.sub.2(YSn)O.sub.5.5), and barium-calcium-niobate (Ba.sub.3(CaNb.sub.2)O.sub.9). In some embodiments, the electrolyte **106** is formed of and includes BZCYYb.

[0054] The hydrogen electrode **102** may be formed of and include a material compatible with the material of the electrolyte **106** and a material of the oxygen electrode **104** under the operating conditions (e.g., temperature, pressure, current density) of the electrochemical cell **100**. The material composition of the hydrogen electrode **102** may facilitate the production of H.sub.2 gas from steam when the electrochemical cell is operated as an electrolysis cell and may also permit electricity generation from H.sub.2 gas when the electrochemical cell **100** is operated as a fuel cell. The hydrogen electrode **102** may be formed of and include a cermet material including at least one metal and at least one perovskite. By way of non-limiting example, the hydrogen electrode **102** may be formed of and include one or more of a nickel/perovskite cermet (Ni-perovskite) material, such as Ni-BZCYYb (e.g., Ni-BZCYYb1711, Ni-BZCYYb4411, Ni-BZCYYb3511), Ni-BSNYYb, Ni-BaCeO.sub.3, Ni—BaZrO.sub.3, Ni-BZY, Ni—Ba.sub.2(YSn)O.sub.5.5, and Ni—Ba.sub.3(CaNb.sub.2)O.sub.9. The hydrogen electrode **102** may include the same perovskite material(s) as the electrolyte **106**. In some embodiments, the hydrogen electrode **102** is formed of and includes Ni-BZCYYb.

[0055] The oxygen electrode **104** may be formed of and include material compatible with the material of the electrolyte **106** and the material of the hydrogen electrode **102** under the operating conditions (e.g., temperature, pressure, current density) of the electrochemical cell **100**. The material composition of the oxygen electrode **104** may facilitate the production of H.sub.2 gas from steam when the electrochemical cell is operated as an electrolysis cell and may also permit electricity generation from H.sub.2 gas when the electrochemical cell **100** is operated as a fuel cell. The material of the oxygen electrode **104** may be a porous material. The oxygen electrode **104** may be formed of and include at least one perovskite material. By way of non-limiting example, the oxygen electrode **104** may be formed of and include one or more of a triple conducting perovskite material, such as Pr(Co.sub.1-x-y-z, Ni.sub.x, Mn.sub.y, Fe.sub.z)O.sub.3- δ , wherein $0 \leq x \leq 0.9$, $0 \leq y \leq 0.9$, $0 \leq z \leq 0.9$, and δ is an oxygen deficit (e.g., PrNi.sub.0.5Co.sub.0.5O.sub.3- δ (PNC55)), a double perovskite material, such as MBa.sub.1-xSr.sub.xCO.sub.2-yFe.sub.yO.sub.5+ δ , wherein x and y are dopant levels, δ is the oxygen deficit, and M is Pr, Nd, or Sm (e.g., PrBa.sub.0.5Sr.sub.0.5Co.sub.1.5Fe.sub.0.5O.sub.5+ δ (PBSCF), NdBa.sub.0.5Sr.sub.0.5Co.sub.1.5Fe.sub.0.5O.sub.5+ δ , SmBa.sub.0.5Sr.sub.0.5Co.sub.1.5Fe.sub.0.5O.sub.5+ δ), a single perovskite material, such as Sm.sub.1-xSr.sub.xCoO.sub.3- δ (SSC), BaZr.sub.1-x-y-zCo.sub.xFe.sub.yY.sub.zO.sub.3- δ , or SrSc.sub.xNd.sub.yCo.sub.1-x-yO.sub.3- δ , wherein x, y, and z are dopant levels and δ is the oxygen deficit; a Ruddleson-Popper-type perovskite material, such as M.sub.2NiO.sub.4- δ , wherein δ is the oxygen deficit and M is La, Pr, Gd, or Sm (e.g., La.sub.2NiO.sub.4- δ , Pr.sub.2NiO.sub.4- δ , Gd.sub.2NiO.sub.4- δ , Sm.sub.2NiO.sub.4- δ); and a single perovskite/perovskite composite material such as SSC-BZCYYb. In some embodiments, the oxygen electrode **104** is formed of and includes PNC55.

[0056] The hydrogen electrode **102**, the oxygen electrode **104**, and the electrolyte **106** may each individually exhibit any desired dimensions (e.g., length, width, thickness) and any desired shape, such as one of a cubic shape, a cuboidal shape, a tubular shape, a tubular spiral shape, a spherical shape, a semi-spherical shape, a cylindrical shape, a semi-cylindrical shape, a conical shape, a triangular prismatic shape, a truncated version of one or more of the foregoing, or an irregular shape. The dimensions and the shapes of the hydrogen electrode **102**, the oxygen electrode **104**,

and the electrolyte **106** may be selected relative to one another such that the electrolyte **106** at least substantially intervenes between opposing surfaces of the hydrogen electrode **102** and the oxygen electrode **104**. In some embodiments, the hydrogen electrode **102** and the oxygen electrode **104** each individually exhibit a thickness within a range of from about 10 micrometers (μm) to about 1000 μm , and the electrolyte **106** exhibits a thickness within a range of from about 5 μm to about 1000 μm .

[0057] FIG. 2 is a cross-sectional view that illustrates a method of forming the electrochemical cell **100** of FIG. 1, according to embodiments of the disclosure. As shown in FIG. 2, the electrolyte **106** is formed adjacent (e.g., directly adjacent) to the hydrogen electrode **102** to collectively form the hydrogen electrode-electrolyte bi-layer **108**. The electrolyte **106** is formed on the hydrogen electrode **102**.

[0058] The hydrogen electrode **102** and the electrolyte **106** may each individually be formed using conventional processes (e.g., rolling processing, milling processing, shaping processes, pressing processes, consolidation processes), which are not described in detail herein. The hydrogen electrode **102** and the electrolyte **106** are annealed (e.g., sintered, co-sintered) to densify the electrolyte **106** and form the hydrogen electrode-electrolyte bi-layer. The hydrogen electrode **102** and the electrolyte **106** may be annealed at a temperature greater than about 1300° C., such as within a range of from about 1300° C. to about 1700° C. or from about 1400° C. to about 1600° C., for a period of time greater than about 3 hours, such as within a range of from about 3 hours to about 7 hours, from about 4 hours to about 6 hours, or about 5 hours. An exposed surface **110** of the electrolyte **106** opposite the hydrogen electrode **102** may be relatively flat (e.g., relatively smooth) after the hydrogen electrode **102** and the electrolyte **106** are annealed. By way of non-limiting example, surface roughness of the surface **110** of the electrolyte **106** after annealing may be within a range of from about 0.1 μm to about 5 μm .

[0059] In some embodiments, the hydrogen electrode-electrolyte bi-layer **108** is formed by a tape-casting process. A green tape of the hydrogen electrode **102** may be prepared by depositing a powder slurry including the hydrogen electrode **102** material(s) onto a substrate having a release material. A green tape of the electrolyte **106** may be prepared by depositing a powder slurry including the electrolyte **106** material(s) onto an additional substrate having a release material. The powder slurry including the hydrogen electrode **102** material(s) and the powder slurry including the electrolyte **106** material(s) may each individually include one or more of a binder, a dispersant, a solvent, or a plasticizer. The powder slurries may each individually be dried on the respective substrates to form the green tape of the hydrogen electrode **102** and the green tape of the electrolyte **106**.

[0060] The green tape of the hydrogen electrode **102** and the green tape of the electrolyte **106** may each individually exhibit any desired dimensions (e.g., length, width, thickness) and any desired shape, such as one of a cubic shape, a cuboidal shape, a tubular shape, a tubular spiral shape, a spherical shape, a semi-spherical shape, a cylindrical shape, a semi-cylindrical shape, a conical shape, a triangular prismatic shape, a truncated version of one or more of the foregoing, or an irregular shape, in order to produce the desired dimensions of the hydrogen electrode **102** and the electrolyte **106**.

[0061] At least one piece (e.g., layer) of the green tape of the hydrogen electrode **102** and at least one piece (e.g., layer) of the green tape of the electrolyte **106** may be laminated by hot press at a temperature within a range of from about 50° C. to about 100° C., such as about 70° C., under a pressure within a range of from about 3 tons to about 5 tons, such as about 4 tons (e.g., from about 3000 kilograms (kg) to about 5000 kilograms, such as about 4000 kg) for a period of time within a range of from about 4 hours to about 6 hours, such as about 5 hours. In some embodiments, two or more pieces (e.g., layers) of the green tape of the hydrogen electrode **102** and one piece (e.g., layer) of the green tape of the electrolyte **106** are laminated. The laminated green tapes may be pre-annealed (e.g., pre-sintered) at a temperature within a range of from about 800° C. to about 1100°

C., such as from about 900° C. to about 1000° C., for a period of time within a range of from about 1 hour to about 5 hours, such as from about 2 hour to about 4 hours or about 3 hours, to remove any organic materials from the laminated green tapes. The laminated green tapes may be annealed (e.g., sintered, co-sintered) at a temperature greater than about 1300° C., such as within a range of from about 1300° C. to about 1700° C. or from about 1400° C. to about 1600° C., for a period of time greater than about 3 hours, such as within a range of from about 3 hours to about 7 hours, from about 4 hours to 6 hours, or about 5 hours. In some embodiments, the pre-annealed laminated green tapes are annealed at a temperature of about 1400° C. for about 5 hours.

[0062] After annealing the hydrogen electrode **102** and the electrolyte **106**, the electrolyte **106** is exposed to one or more surface treatment acts (e.g., surface treatments) to increase surface roughness of a surface **110** of the electrolyte **106** opposite the hydrogen electrode **102**. However, one or more of the surface treatment acts may be conducted without first annealing the hydrogen electrode **102** and the electrolyte **106**. The one or more surface treatments may remove a portion of the electrolyte **106** adjacent to (e.g., directly adjacent to) the surface **110**. The one or more surface treatments may include exposing the electrolyte **106** to one or more of an acid solution, a plasma, thermal shock, or gamma radiation. In some embodiments, the one or more surface treatments includes a series of (e.g., more than one) surface treatments. The series of surface treatments may include one or more repeated surface treatments. In other words, if multiple surface treatments are conducted, the multiple treatments may include multiple surface treatments of the same type or may include a combination of different types of surface treatments. By way of non-limiting example, the one or more surface treatments may include exposing the electrolyte **106** to one or more acid solutions, one or more plasmas, thermal shock at one or more temperatures and/or for one or more periods of time, or gamma radiation at one or more wavelengths and/or for one or more periods of time. The increased surface roughness of the electrolyte **106** may result in an increased effective surface area of the surface **110** of the electrolyte **106**.

[0063] The electrolyte **106** may be exposed to the one or more surface treatments for any period of time sufficient to obtain a suitable (e.g., desired) surface roughness of the surface **110** of the electrolyte **106** and maintain a desired thickness of the electrolyte **106**. As a non-limiting example, the electrolyte **106** may be exposed to each of the one or more surface treatments individually for a period of time within a range of from about 10 seconds to about 60 minutes, such as from about 10 seconds to about 2 minutes, from about 10 seconds to about 5 minutes, from about 10 seconds to about 10 minutes, from about 1 minute to about 10 minutes, 1 minute to about 30 minutes, from about 5 minutes to about 10 minutes, from about 5 minutes to about 15 minutes, from about 5 minutes to about 30 minutes, from about 10 minutes to about 20 minutes, from about 10 minutes to about 30 minutes, from about 10 minutes to about 50 minutes, from about 20 minutes to about 40 minutes, from about 20 minutes to about 60 minutes, or from about 30 minutes to about 60 minutes. In some embodiments, the electrolyte **106** is exposed to each of the one or more surface treatments for a period of time within a range of from about 5 minutes to about 15 minutes. In some embodiments, the electrolyte **106** is exposed to each of the one or more surface treatments for a period of time of about 10 minutes.

[0064] In some embodiments, at least one of the one or more surface treatments is an etch process, such as a wet etch process, in which the electrolyte **106** is exposed to an acid solution to remove a portion of the electrolyte **106** and increase surface roughness of the surface **110** of the electrolyte **106**. By way of non-limiting example, the acid solution may be a solution of nitric acid (HNO₃), hydrochloric acid (HCl), sulfuric acid (H₂SO₄), phosphoric acid (H₃PO₄), or a combination thereof. The acid solution may be an aqueous solution. A concentration of the acid in the acid solution may be sufficient to remove the portion of the electrolyte **106**, such within a range of from about 0.4% by weight (wt %) to about 80 wt %, such as from about 0.45 wt % to about 73 wt %, from about 0.4 wt % to about 10 wt %, from about 0.4 wt % to about 30 wt %, from about 0.5 wt % to about 80 wt %, from about 1 wt % to about 15 wt

%, from about 1 wt % to about 40 wt %, from about 1 wt % to about 60 wt %, from about 1 wt % to about 75 wt %, from about 10 wt % to about 40 wt %, from about 20 wt % to about 60 wt %, from about 30 wt % to about 70 wt %, from about 50 wt % to about 80 wt %, or from about 60 wt % to about 80 wt %. In some embodiments, a concentration of acid in the acid solution is within a range of from about 60 wt % to about 80 wt %. In some embodiments, a concentration of acid in the acid solution is about 70 wt %.

[0065] Exposing the electrolyte **106** to a plasma (e.g., a plasma treatment) may include exposing the electrolyte **106** to one or more of an oxygen plasma, an argon plasma, a hydrogen plasma, or a plasma of air. Exposing the electrolyte **106** to the plasma may remove a portion of the electrolyte **106** to increase surface roughness of the surface **110** of the electrolyte **106**.

[0066] Exposing the electrolyte **106** to thermal shock (e.g., a thermal shock treatment) may include one or more of increasing or decreasing a temperature of the electrolyte **106**. The thermal shock treatment may include increasing a temperature of the electrolyte **106** at a rate within a range of from about 10° C. sec.sup.-1 to about 1000° C. sec.sup.-1 to an elevated temperature within a range of from about 100° C. to about 2000° C. The thermal shock treatment may include decreasing a temperature of the electrolyte **106** at a rate within a range of from about -10° C. sec.sup.-1 to about -1000° C. sec.sup.-1 to a decreased temperature within a range of from about 100° C. to about 2000° C. Exposing the electrolyte **106** to the thermal shock treatment increases surface roughness of the surface **110** of the electrolyte **106**.

[0067] Exposing the electrolyte **106** to gamma radiation (e.g., a gamma radiation treatment) may include exposing the electrolyte **106** to gamma radiation having a wavelength less than about 100 picometers (pm), such as within a range of from about 10 pm to about 100 pm. Exposing the electrolyte **106** to the gamma radiation treatment increases surface roughness of the surface **110** of the electrolyte **106**.

[0068] After the electrolyte **106** is exposed to the one or more surface treatments, surface roughness of the surface **110** of the electrolyte may be determined by calculating an average distance between adjacent peaks and valleys of the surface **110**. The surface roughness may be within a range of from about 0.1 μm to about 5 μm, such as from about 0.1 μm to about 1 μm, from about 0.1 μm to about 2 μm, from about 0.1 μm to about 4 μm, from about 0.5 μm to about 1 μm, from about 0.5 μm to about 3 μm, from about 0.5 μm to about 5 μm, from about 1 μm to about 3 μm, from about 1 μm to about 5 μm, from about 2 μm to about 4 μm, from about 2 μm to about 5 μm, from about 3 μm to about 5 μm, or from about 4 μm to about 5 μm. In some embodiments, surface roughness of the surface **110** of the electrolyte **106** is within a range of from about 0.5 μm to about 1 μm after exposure to the one or more surface treatments.

[0069] With reference to FIG. 1, the oxygen electrode **104** is formed adjacent to (e.g., directly adjacent to) the surface **110** of the electrolyte **106** opposite the hydrogen electrode **102**. An electrolyte/oxygen electrode/inlet gas of the oxygen electrode triple-phase boundary may be formed at the interface when the oxygen electrode **104** is formed directly adjacent to the electrolyte **106**. The oxygen electrode **104** may be formed using conventional processes (e.g., rolling processing, milling processing, shaping processes, pressing processes, consolidation processes, screen-printing processes, painting processes), which are not described in detail herein. The oxygen electrode **104**, the electrolyte **106**, and the hydrogen electrode **102** are annealed at a temperature within a range of from about 800° C. to about 1200° C., such as about 1000° C., to bond the oxygen electrode **104** to the electrolyte **106** along an oxygen electrode-electrolyte interface **112** disposed between the oxygen electrode **104** to the electrolyte **106** and form the electrochemical cell **100**.

[0070] The oxygen electrode-electrolyte interface **112** may exhibit a peeling strength corresponding (e.g., correlating) to the surface roughness of the surface **110** of the electrolyte **106**. By way of a non-limiting example, the oxygen electrode-electrolyte interface **112** may exhibit a peeling strength within a range of from about 17 Newtons (N) to about 40 N, such as from about 17 N to about 35

N, from about 17 N to about 25 N, from about 20 N to about 35 N, from about 20 N to about 30 N, from about 20 N to about 40 N, or from about 25 N to about 40 N. In some embodiments, the oxygen electrode-electrolyte interface **112** exhibits a peeling strength of at least about 23 N. In some embodiments, the oxygen electrode-electrolyte interface **112** exhibits a peeling strength of about 23.5 N.

[0071] Exposing the surface **110** of the electrolyte **106** to the one or more surface treatments may increase the interfacial bond strength of the oxygen electrode **104** and the electrolyte **106** along the oxygen electrode-electrolyte interface **112**. Removing a portion of the electrolyte **106** adjacent to the surface **110** by exposing the surface **110** to the one or more surface treatments enables deficiency of an A-site cation at the surface **110** of the electrolyte to be increased, improving A-site cation inter-diffusion (e.g., oxide-oxide wetting) between the material of the oxygen electrode **104** and the material of the electrolyte **106** and further increasing the bond strength between the oxygen electrode **104** and the electrolyte **106**. In addition, increasing surface roughness of the surface **110** of the electrolyte **106** by exposing the surface **110** to the one or more surface treatments enables an effective surface area of the surface **110** to be increased. By increasing the effective surface area of the surface **110** and improving interfacial strength and bond strength along the oxygen electrode-electrolyte interface **112**, a contact area (e.g., number of contacts) and a number of reaction sites near the oxygen electrode-electrolyte interface **112** may be increased, enabling the electrochemical cell **100** to approach or achieve intrinsic bulk proton conductivity, improving electrochemical performance and long-term stability of the electrochemical cell **100**. In comparison, conventional electrochemical cells formed without exposing the electrolyte **106** to the one or more surface treatments exhibit high ohmic loss as a result of extended proton conduction through the oxygen electrode due to poor bonding, limited contact, and a decreased number of reaction sites near the interface between the electrolyte and the oxygen electrode.

[0072] Electrochemical cells (e.g., the electrochemical cell **100**) in accordance with embodiments of this disclosure may be used in embodiments of H.sub.2 gas production and electricity generation systems of the disclosure. For example, FIG. 3 schematically illustrates a system **300** for producing H.sub.2 gas and generating electricity, according to embodiments of the disclosure. As shown in FIG. 3, the system **300** includes at least one steam source **302**, and at least one electrochemical apparatus **304** in fluid communication with the steam source **302**. The electrochemical apparatus **304** includes a housing structure **306**, and one or more embodiments of the electrochemical cell **100** previously described with reference to FIG. 1 contained within the housing structure **306**. The electrochemical cell **100** is electrically connected (e.g., coupled) to a power source **308**, and includes the oxygen electrode **104** (e.g., steam side electrode), the hydrogen electrode **102** (e.g., H.sub.2 gas side electrode), and the electrolyte **106** between the oxygen electrode **104** and the hydrogen electrode **102**. As shown in FIG. 3, the system **300** may optionally include one or more of at least one H.sub.2 gas source **310** in fluid communication with the electrochemical apparatus **304**, at least one O.sub.2 gas source **332** in fluid communication with the electrochemical apparatus **304**, and at least one heating apparatus **312** operatively associated with the electrochemical apparatus **304**.

[0073] The steam source **302** comprises at least one apparatus configured and operated to produce a steam stream **314** including steam (e.g., gaseous H.sub.2O). The steam stream **314** may be directed into the electrochemical apparatus **304** from the steam source **302** to interact with the oxygen electrode **104** of the electrochemical cell **100** when the electrochemical cell **100** is operated as an electrolysis cell, as described in further detail below. The steam source **302** may also receive an H.sub.2O stream **316** containing one or more phases of H.sub.2O (e.g., steam) exiting the electrochemical apparatus **304** when the electrochemical cell **100** is operated as a fuel cell, as also described in further detail below. By way of non-limiting example, the steam source **302** may comprise a boiler apparatus configured and operated to heat liquid H.sub.2O to a temperature greater than or equal to about 100° C. In some embodiments, the steam source **302** is configured

and operated to convert the liquid H.sub.2O to steam having a temperature within a range of an operating temperature of the electrochemical cell **100** of the electrochemical apparatus **304**, such as a temperature within a range of from about 400° C. to about 600° C. In some embodiments, the steam source **302** is configured and operated to convert the liquid H.sub.2O into steam having a temperature below the operating temperature of the electrochemical cell **100**. In such embodiments, the heating apparatus **312** may be employed to further heat the steam stream **314** to the operational temperature of the electrochemical cell **100**, as described in further detail below.

[0074] The electrochemical apparatus **304**, including the housing structure **306** and the electrochemical cell **100** thereof, is configured and operated to facilitate the production of H.sub.2 gas from steam (e.g., steam of the steam stream **314**) when the electrochemical cell **100** is operated as an electrolysis cell, and to facilitate the electricity generation from H.sub.2 gas (e.g., the H.sub.2 gas produced when the electrochemical cell **100** is operated as an electrolysis cell) when the electrochemical cell **100** is operated as a fuel cell. The housing structure **306** may exhibit any shape (e.g., a tubular shape, a quadrilateral shape, a spherical shape, a semi-spherical shape, a cylindrical shape, a semi-cylindrical shape, truncated versions thereof, or an irregular shape) and size able to contain (e.g., hold) the electrochemical cell **100** therein. In addition, the housing structure **306** is configured, such that when the electrochemical cell **100** is operated as an electrolysis cell, the housing structure **306** may receive and direct the steam stream **314** to the oxygen electrode **104** of the electrochemical cell **100**, may direct O.sub.2 gas produced at the oxygen electrode **104** of the electrochemical cell **100** away from the electrochemical apparatus **304** as an O.sub.2 gas stream **318**, and may optionally direct H.sub.2 gas produced at the hydrogen electrode **102** of the electrochemical cell **100** away from the electrochemical apparatus **304** as an H.sub.2 gas stream **322**. The housing structure **306** may also be configured, such that when the electrochemical cell **100** is operated as a fuel cell, the housing structure **306** may receive and direct a H.sub.2 gas containing stream **324** to the hydrogen electrode **102** of the electrochemical cell **100**, may receive and direct an O.sub.2 gas containing stream **320** to the oxygen electrode **104** of the electrochemical cell **100**, and may direct H.sub.2O produced at the oxygen electrode **104** of the electrochemical cell **100** away from the electrochemical apparatus **304** as an H.sub.2O stream **316**. The housing structure **306** may be formed of and include any material (e.g., glass, metal, alloy, polymer, ceramic, composite, combination thereof, etc.) compatible with the operating conditions (e.g., temperatures, pressures, etc.) of the electrochemical apparatus **304**.

[0075] The housing structure **306** of the electrochemical apparatus **304** may at least partially define at least one internal chamber **326** at least partially surrounding the electrochemical cell **100**. The electrochemical cell **100** may serve as a boundary between a first region **328** (e.g., a steam region) of the internal chamber **326** configured and positioned to temporarily contain steam, and a second region **330** (e.g., an H.sub.2 gas region) of the internal chamber **326** configured and positioned to temporarily contain H.sub.2 gas. H.sub.2O (e.g., steam) may be substantially limited to the first region **328** of the internal chamber **326** by the configurations and positions of the housing structure **306** and the electrochemical cell **100**. Keeping the second region **330** of the internal chamber **326** substantially free of the H.sub.2O circumvents additional processing of produced H.sub.2 gas (e.g., to separate the produced H.sub.2 gas from steam) that may otherwise be necessary if the H.sub.2O (e.g., steam) was provided within the second region **330** of the internal chamber **326**. In addition, protecting the hydrogen electrode **102** of the electrochemical cell **100** from exposure to H.sub.2O may enhance the operational life (e.g., durability) of the electrochemical cell **100** as compared to conventional electrochemical cells by preventing undesirable oxidation of the hydrogen electrode **102** that may otherwise occur in the presence of H.sub.2O.

[0076] Although the electrochemical apparatus **304** is depicted in FIG. 3 as including a single (i.e., only one) electrochemical cell **100**, the electrochemical apparatus **304** may include any number of electrochemical cells **100**. Put another way, the electrochemical apparatus **304** may include a single (e.g., only one) electrochemical cell **100**, or may include multiple (e.g., more than one)

electrochemical cells **100** that are operatively coupled to one another. If the electrochemical apparatus **304** includes multiple electrochemical cells **100**, each of the electrochemical cells **100** may be substantially the same (e.g., exhibit substantially the same components, component sizes, component shapes, component material compositions, component material distributions, component positions, component orientations, etc.) and may be operated under substantially the same conditions (e.g., substantially the same temperatures, pressures, flow rates, etc.), or at least one of the electrochemical cells **100** may be different (e.g., exhibit one or more of different components, different component sizes, different component shapes, different component material compositions, different component material distributions, different component positions, different component orientations, etc.) than at least one other of the electrochemical cells **100** and/or may be operated under different conditions (e.g., different temperatures, different pressures, different flow rates, etc.) than at least one other of the electrochemical cells **100**. By way of non-limiting example, one of the electrochemical cells **100** may be configured for and operated under a different temperature (e.g., different operating temperature resulting from a different material composition of one of more components thereof) than at least one other of the electrochemical cells **100**. In some embodiments, two or more electrochemical cells **100** are provided in parallel with one another within the housing structure **306** of the electrochemical apparatus **304**.

[0077] Although the system **300** is depicted as including a single (i.e., only one) electrochemical apparatus **304** in FIG. **3**, the system **300** may include any number of electrochemical apparatuses **304**. Put another way, the system **300** may include a single (e.g., only one) electrochemical apparatus **304**, or may include multiple (e.g., more than one) electrochemical apparatuses **304** that are operatively coupled to one another. If the system **300** includes multiple electrochemical apparatuses **304**, each of the electrochemical apparatuses **304** may be substantially the same (e.g., exhibit substantially the same components, component sizes, component shapes, component material compositions, component material distributions, component positions, component orientations, etc.) and may be operated under substantially the same conditions (e.g., substantially the same temperatures, pressures, flow rates, etc.), or at least one of the electrochemical apparatus **304** may be different (e.g., exhibit one or more of different components, different component sizes, different component shapes, different component material compositions, different component material distributions, different component positions, different component orientations, etc.) than at least one other of the electrochemical apparatuses **304** and/or may be operated under different conditions (e.g., different temperatures, different pressures, different flow rates, etc.) than at least one other of the electrochemical apparatuses **304**. By way of non-limiting example, one of the electrochemical apparatuses **304** may be configured for and operated under a different temperature (e.g., a different operating temperature resulting from a different material composition of one of more components of one or more electrochemical cell(s) **100** thereof) than at least one other of the electrochemical apparatuses **304**. In some embodiments, two or more electrochemical apparatuses **304** are provided in parallel with one another. In some embodiments, two or more electrochemical apparatuses **304** are provided in series with one another.

[0078] The power source **308** may comprise one or more of a device, structure, and apparatus able to apply a potential difference (e.g., voltage) between the oxygen electrode **104** of the electrochemical cell **100** and the hydrogen electrode **102** of the electrochemical cell **100** to facilitate desired operation (e.g., operation as an electrolysis cell, operation as a fuel cell) of the electrochemical cell **100**. During operation of the electrochemical cell **100** as an electrolysis cell, the potential difference applied between the oxygen electrode **104** and the hydrogen electrode **102** permits the oxygen electrode **104** to serve as a positive electrode (e.g., anode) and the hydrogen electrode **102** to serve as a negative electrode (e.g., cathode) to facilitate water splitting reaction (WSR) and the production of H₂ gas from steam, as described in further detail below. During operation of the electrochemical cell **100** as a fuel cell, the potential difference applied between the oxygen electrode **104** and the hydrogen electrode **102** permits the hydrogen electrode **102** to serve

as the positive electrode (e.g., anode) and the oxygen electrode **104** to serve as the negative electrode (e.g., cathode) to facilitate oxygen reduction reaction (ORR) and the electricity generation using H.sub.2 gas as a fuel, as also described in further detail below. The power source **308** may, for example, comprise one or more of a device, structure, or apparatus configured and operated to exploit one or more of solar energy, wind (e.g., wind turbine) energy, hydropower energy, geothermal energy, nuclear energy, combustion-based energy, and waste heat (e.g., heat generated from one or more of an engine, a chemical process, and a phase change process) to apply a potential difference between the oxygen electrode **104** and the hydrogen electrode **102** of the electrochemical cell **100**.

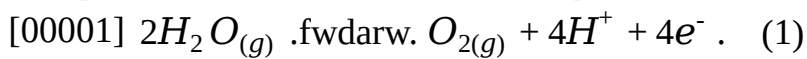
[0079] The heating apparatus **312**, if present, may comprise at least one apparatus (e.g., one or more of a combustion heater, an electrical resistance heater, an inductive heater, and an electromagnetic heater) configured and operated to heat one or more of at least a portion of the electrochemical apparatus **304** and one or more of the streams (e.g., one or more of the steam stream **314**, the H.sub.2 gas containing stream **324**, and the O.sub.2 gas containing stream **320**) directed into the electrochemical apparatus **304** during desired operation (e.g., operation as an electrolysis cell, operation as a fuel cell) of the electrochemical cell **100** to an operating temperature of the electrochemical apparatus **304**. The operating temperature of the electrochemical apparatus **304** may at least partially depend on the material compositions of the oxygen electrode **104**, the electrolyte **106**, and the hydrogen electrode **102** thereof. In some embodiments, the heating apparatus **312** heats one or more of at least a portion of the electrochemical apparatus **304** and one or more of the streams directed into the electrochemical apparatus **304** to a temperature within a range of from about 400° C. to about 600° C. In additional embodiments, such as in embodiments where the temperature of the streams is already within the operating temperature range of the electrochemical cell **100** of the electrochemical apparatus **304**, the heating apparatus **312** may be omitted (e.g., absent) from the system **300**.

[0080] The H.sub.2 gas source **310**, if present, may comprise one or more of a device, structure, and apparatus configured and operated to produce an H.sub.2 gas containing stream **324** including H.sub.2 gas. The H.sub.2 gas containing stream **324** may be directed into the electrochemical apparatus **304** from the H.sub.2 gas source **310** to interact with the hydrogen electrode **102** of the electrochemical cell **100** when the electrochemical cell **100** is operated in as a fuel cell, as described in further detail below. The H.sub.2 gas source **310** may also receive and temporarily store (e.g., contain) one or more portions of the H.sub.2 gas stream **322** including H.sub.2 gas exiting the electrochemical apparatus **304** when the electrochemical cell **100** is operated as an electrolysis cell, as also described in detail below. The H.sub.2 gas exiting the electrochemical apparatus **304** in the H.sub.2 gas stream **322** during operation of the electrochemical cell **100** as an electrolysis cell may be employed as at least a portion of the H.sub.2 gas of the H.sub.2 gas containing stream **324** being directed into the electrochemical apparatus **304** when the electrochemical cell **100** is operated as a fuel cell. In additional embodiments, such as in embodiments wherein the electrochemical cell **100** is rapidly switched (e.g., in under five (5) minutes, such as in under two (2) minutes, or in under one (1) minute) between operation as an electrolysis cell and operation as a fuel cell, the H.sub.2 gas source **310** may be omitted. In such embodiments, the at least a portion (e.g., substantially all) of the H.sub.2 gas produced during operation of the electrochemical cell **100** as an electrolysis cell may be employed as fuel during operation of the electrochemical cell **100** as a fuel cell before the produced H.sub.2 gas exits the second region **330** of the internal chamber **326** of the housing structure **306**.

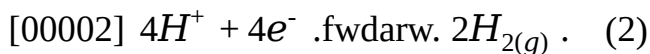
[0081] The O.sub.2 gas source **332**, if present, may comprise one or more of a device, structure, and apparatus configured and operated to produce an O.sub.2 gas containing stream **320** including O.sub.2 gas. The O.sub.2 gas containing stream **320** may be directed into the electrochemical apparatus **304** from the O.sub.2 gas source **332** to interact with the oxygen electrode **104** of the electrochemical cell **100** when the electrochemical cell **100** is operated as a fuel cell, as described

in further detail below. The O.sub.2 gas source **332** may also receive and temporarily store (e.g., contain) one or more portions of the O.sub.2 gas stream **318** including O.sub.2 gas exiting the electrochemical apparatus **304** when the electrochemical cell **100** is operated as an electrolysis cell. The O.sub.2 gas exiting the electrochemical apparatus **304** in the O.sub.2 gas stream **318** during operation of the electrochemical cell **100** as an electrolysis cell may be employed as at least a portion of the O.sub.2 gas of the O.sub.2 gas containing stream **320** being directed into the electrochemical apparatus **304** when the electrochemical cell **100** is operated as a fuel cell.

[0082] When the electrochemical cell **100** of the electrochemical apparatus **304** (and, hence, the electrochemical apparatus **304** itself) is operated as an electrolysis cell, the system **300** directs the steam stream **314** from the steam source **302** and into the electrochemical apparatus **304** to interact with the oxygen electrode **104** (e.g., steam side electrode) of the electrochemical cell **100** contained therein. A potential difference (e.g., voltage) is applied between the oxygen electrode **104** (serving as an anode) and the hydrogen electrode **102** (serving as a cathode) by the power source **308** so that as steam interacts with the oxygen electrode **104**, H atoms of the steam release their electrons (e.sup.-) to generate oxygen gas (O.sub.2(g)), hydrogen ions (H.sup.+) (i.e., protons), and electrons (e.sup.-) according to the following equation:



[0083] The generated H.sup.+ permeate (e.g., diffuse) across the electrolyte **106** to the hydrogen electrode **102**, and the generated e.sup.- are directed to the power source **308** through external circuitry (not shown). The produced O.sub.2 gas may exit the electrochemical apparatus **304** as an O.sub.2 gas stream **318**. At the hydrogen electrode **102**, the generated H.sup.+ exiting the electrolyte **106** react with e received from the power source **308** to form H atoms, which then combine to form H.sub.2 gas (H.sub.2(g)), according to the following equation:

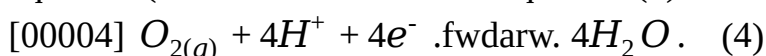


[0084] The produced H.sub.2 gas may exit the electrochemical apparatus **304** as the H.sub.2 gas stream **322**.

[0085] When the electrochemical cell **100** of the electrochemical apparatus **304** (and, hence, the electrochemical apparatus **304** itself) is operated as a fuel cell, the system **300** employs H.sub.2 gas previously produced by the electrochemical cell **100** when operated as an electrolysis cell and/or directed into electrochemical apparatus **304** (e.g., into the second region **330** thereof) from the H.sub.2 gas source **310** as a gaseous H.sub.2 stream **316** to interact with the hydrogen electrode **102** (e.g., H.sub.2 gas side electrode) of the electrochemical cell **100**. A potential difference (e.g., voltage) is applied between the hydrogen electrode **102** (serving as an anode) and the oxygen electrode **104** (serving as a cathode) by the power source **308** so that as H.sub.2 gas interacts with the hydrogen electrode **102**, H atoms of the H.sub.2 gas release their electrons (e.sup.-) to generate hydrogen ions (H.sup.+) (i.e., protons) and electrons (e.sup.-) according to the following equation (the reverse reaction of Equation (2) above):



[0086] The generated H.sup.+ permeate (e.g., diffuse) across the electrolyte **106** to the oxygen electrode **104**, and the generated e are directed to the power source **308** through external circuitry. At the oxygen electrode **104**, the generated H.sup.+ exiting the electrolyte **106** react with e.sup.- received from the power source **308** and O.sub.2 gas previously produced by the electrochemical cell **100** when operated in as an electrolysis cell and/or directed into electrochemical apparatus **304** (e.g., into the first region **328** thereof) from the O.sub.2 gas source **332** as an O.sub.2 gas containing stream **320** to generate electricity and produce H.sub.2O, according to the following equation (the reverse reaction of Equation (1) above):



[0087] The produced H.sub.2O may exit the electrochemical apparatus **304** as the H.sub.2O stream

316 and may be directed into the steam source **302**, and/or may be employed to produce additional H.sub.2 gas when the electrochemical cell **100** of the electrochemical apparatus **304** (and, hence, the electrochemical apparatus **304** itself) is operated as an electrolysis cell.

[0088] Switching between operation of the electrochemical cell **100** as an electrolysis cell and operation of the electrochemical cell **100** as a fuel cell may be rapid (e.g., electrolysis and fuel cell operation modes may alternate between one another using relatively short time periods for each operation, such as time periods less than or equal to about five (5) minutes, less than or equal to about two (2) minutes, or less than or equal to about one (1) minute), or may be delayed (e.g., the electrolysis and fuel cell operation modes may not alternate between one another using relatively short time periods). In some embodiments, such as embodiments wherein the electrochemical cell **100** is rapidly switched (e.g., cyclically alternated) between operation as an electrolysis cell and operation as a fuel cell, at least a portion (e.g., substantially all) of the H.sub.2 gas produced during operation of the electrochemical cell **100** as an electrolysis cell is consumed as fuel during operation of the electrochemical cell **100** as a fuel cell before the produced H.sub.2 gas can exit the electrochemical apparatus **304** as the H.sub.2 gas stream **322**. In additional embodiments, such as embodiments wherein the electrochemical cell **100** is not rapidly switched between operation as an electrolysis cell and operation as a fuel cell, at least a portion (e.g., substantially all) of the H.sub.2 gas produced during operation of the electrochemical cell **100** as an electrolysis cell may exit the electrochemical apparatus **304** as the H.sub.2 gas stream **322** and may be stored (e.g., at the H.sub.2 gas source **310**, if any) for subsequent use (e.g., for subsequent use as fuel during relatively delayed operation of the electrochemical cell **100** as a fuel cell), as desired.

[0089] Still referring to FIG. 3, streams exiting the electrochemical apparatus **304** during the different modes of operation (e.g., operation as an electrolysis cell, operation of a fuel cell) of the electrochemical cell **100** may individually be utilized or disposed of as desired. In some embodiments, one or more of the H.sub.2 gas stream **322** and the O.sub.2 gas stream **318** produced during operation of the electrochemical cell **100** as an electrolysis cell are respectively delivered into one or more storage vessels of the H.sub.2 gas source **310** and the O.sub.2 gas source **332** for subsequent use (e.g., to respectively form the H.sub.2 gas containing stream **324** and the O.sub.2 gas containing stream **320** employed during operation of the electrochemical cell **100** as a fuel cell), as desired. In additional embodiments, the H.sub.2O stream **316** produced during operation of the electrochemical cell **100** as a fuel cell is delivered into one or more storage vessels of the steam source **302** for subsequent use (e.g., to form the steam stream **314** employed during operation of the electrochemical cell **100** as an electrolysis cell), as desired. In further embodiments, at least a portion of one or more of the streams (e.g., the H.sub.2 gas stream **322** and the O.sub.2 gas stream **318**) may be utilized (e.g., combusted) to heat one or more components (e.g., the heating apparatus **312** (if present); the electrochemical apparatus **304**; etc.) and/or other streams (e.g., the steam stream **314**) of the system **300**. By way of non-limiting example, if the heating apparatus **312** (if present) is a combustion-based apparatus, at least a portion of one or more of the H.sub.2 gas stream **322** and the O.sub.2 gas stream **318** may be directed into the heating apparatus **312** and undergo a combustion reaction to efficiently heat the steam stream **314** entering the electrochemical apparatus **304** and/or at least a portion of the electrochemical apparatus **304** during operation of the electrochemical cell **100** as an electrolysis cell. Utilizing the hydrocarbon H.sub.2 gas stream **322** and/or the O.sub.2 gas stream **318** as described above may reduce the electrical power requirements of the system **300** by enabling the utilization of direct thermal energy.

[0090] Thermal energy input into (e.g., through the heating apparatus **312** (if present)) and/or generated by the electrochemical apparatus **304** may also be used to heat one or more other components and/or streams of the system **300**. As a non-limiting example, during operation of the electrochemical cell **100** as an electrolysis cell, one or more of the H.sub.2 gas stream **322** and the O.sub.2 gas stream **318** exiting the electrochemical apparatus **304** may be directed into a heat exchanger configured and operated to facilitate heat exchange between the H.sub.2 gas stream **322**

and/or the O.sub.2 gas stream **318** of the system **300** and one or more other relatively cooler streams (e.g., in some embodiments, the steam stream **314**) of the system **300** to transfer heat from the H.sub.2 gas stream **322** and/or the O.sub.2 gas stream **318** to the relatively cooler stream(s) to facilitate the recovery of the thermal energy input into and generated within the electrochemical apparatus **304**. The recovered thermal energy may increase process efficiency and/or reduce operational costs without having to react (e.g., combust) the H.sub.2 gas stream **322** and/or the O.sub.2 gas stream **318**. As another non-limiting example, during operation of the electrochemical cell **100** as a fuel cell, the H.sub.2O stream **316** exiting the electrochemical apparatus **304** may be directed into a heat exchanger configured and operated to facilitate heat exchange between the H.sub.2O stream **316** of the system **300** and one or more other relatively cooler streams (e.g., in some embodiments, one or more of the O.sub.2 gas containing stream **320** and the H.sub.2 gas containing stream **324**) of the system **300** to transfer heat from the H.sub.2O stream **316** to the relatively cooler stream(s) to facilitate the recovery of the thermal energy input into and generated within the electrochemical apparatus **304**. The recovered thermal energy may increase process efficiency and/or reduce operational costs without having to react (e.g., combust) one or more of the streams employed in the system **300**.

[0091] The electrochemical cells (e.g., the electrochemical cell **100**), systems (e.g., the system **300**), and methods of the disclosure enable a bulk proton conductivity to approach or achieve a theoretical bulk proton conductivity (e.g., intrinsic bulk proton conductivity), improving electrochemical performance at intermediate temperatures, such as temperatures within a range of from about 400° to about 600° C. The electrochemical cells, systems, and methods of the disclosure may reduce one or more of the time (e.g., processing acts), costs (e.g., material costs), and energy (e.g., thermal energy, electrical energy, etc.) utilized to produce H.sub.2 gas and/or generate electricity relative to conventional electrochemical cells, systems, and methods. The electrochemical cells, systems, and methods of the disclosure may be more efficient, durable, and reliable than conventional electrochemical cells, conventional systems, and conventional methods of H.sub.2 gas production and electricity. The methods of the disclosure may be applied in various technologies and industries including, but not limited to, power industry applications, petrochemical industry applications, and other energy-intense industrial applications, including, but not limited to ammonia industry applications, Li-ion batteries, fuel cells, electrolyzers, supercapacitors, biomass conversion, and fuel-efficient vehicles.

[0092] The following examples serve to explain embodiments of the disclosure in more detail. These examples are not to be construed as being exhaustive, exclusive, or otherwise limiting as to the scope of the disclosure.

EXAMPLES

Example 1

Materials Synthesis

[0093] BZCYYb was synthesized by a solid-state reaction method. Stoichiometric amounts of BaCO.sub.3 (99.8% purity), ZrO.sub.2 (99% purity), CeO.sub.2 (99.9% purity), Y.sub.2O.sub.3 (99.9% purity), and Yb.sub.2O.sub.3 (99.9% purity) were mixed by ball milling in ethanol for 12 hours, followed by drying, grinding, and heat treatment at 1100° C. for 8 hours to obtain phase-pure BZCYYb powders.

[0094] PNC55 was synthesized by the Pechini method. Stoichiometric Pr(NO.sub.3).sub.3.Math.6H.sub.2O (99.9% purity), Ni(NO.sub.3).sub.2.Math.6H.sub.2O (99% purity), and Co(NO.sub.3).sub.2.Math.6H.sub.2O (99% purity) were dissolved in deionized water to prepare an aqueous solution containing 0.05 mol L.sup.-1 Pr.sup.3+, 0.025 mol L.sup.-1 Ni.sup.2+, and 0.025 mol L.sup.-1 Co.sup.2+. Next, 0.2 mol L.sup.-1 glycol and 0.1 mol L.sup.-1 citric acid were added to the aqueous solution. The prepared solution was heated to 80° C. on a hot plate with continuous stirring until converted to a gel. The obtained gel was heated to 350° C. and followed with an auto-ignition process to produce a black foamy intermediate product. Final

PNC55 powders were obtained by annealing the intermediate product at 1100° C. for 4 hours. La.sub.0.6Sr.sub.0.4Co.sub.0.2Fe.sub.0.8O.sub.3-δ (LSCF), PrBa.sub.0.5Sr.sub.0.5Co.sub.1.5Fe.sub.0.5O.sub.5+δ (PBSCF), and PrNi.sub.0.7Co.sub.0.3O.sub.3-δ (PNC73) were similarly synthesized by the same Pechini method following their stoichiometries.

[0095] A three-dimensional (3D) PNC73 mesh was fabricated by a template-derived method. Stoichiometric Pr(NO.sub.3).sub.3.Math.6H.sub.2O (99.9% purity), Ni(NO.sub.3).sub.2.Math.6H.sub.2O (99% purity), and Co(NO.sub.3).sub.2.Math.6H.sub.2O (99%, purity) were dissolved in deionized water to prepare a nitrate precursor solution containing 0.05 mol L.sup.-1 Pr.sup.3+, 0.035 mol L.sup.-1 Ni.sup.2+, and 0.015 mol L.sup.-1 Co.sup.2+. A piece of fabric textile (Telio, Montreal, CA) was immersed into the precursor solution for 24 hours and then heat-treated at 750° C. for 2 hours to form a 3D PNC73 mesh.

Example 2

Cell Fabrication

[0096] Hydrogen electrode-supported cells were fabricated by a tape-casting process. To prepare green tapes of the hydrogen electrode, NiO and BZCYYb powders as described in Example 1 were mixed with 6:4 weight ratio by ball milling in ethanol and toluene for 24 hours. Next, a binder of polyvinyl butyral (PVB), a plasticizer of butyl benzyl phthalate (BBP), and a dispersant of fish oil were added to form a powder slurry. The powder slurry was mixed by ball milling for an additional 24 hours to yield the desired slip rheology. Tape casting was performed using a laboratory tape casting machine. The thickness of hydrogen-electrode green tapes was controlled to be about 1 mm after drying at 37.8° C. for 4 hours.

[0097] Green tapes of electrolyte were prepared similarly without adding NiO and by controlling the thickness to about 0.12 mm (for a 22 μm thick BZCYYb electrolyte after sintering) or about 0.08 mm (for a 16 μm thick BZCYYb electrolyte after sintering) after drying. Three pieces of hydrogen-electrode green tapes and one piece of electrolyte green tape were laminated by a hot press at 70° C. under 4 tons (e.g., 4000 kg) for 5 hours. Laminated green tapes were punched and pre-sintered at 920° C. for 3 hours to remove the organics. The co-sintering of hydrogen electrode-electrolyte bi-layer was conducted at T.sub.1=1400° C. for 5 hours (heating rate: 1° C. min.sup.-1 to 1000° C. and 2° C. min.sup.-1 to 1400° C.). The co-sintered hydrogen electrode-electrolyte bi-layer was then allowed to cool.

[0098] A surface treatment including concentrated nitric acid was applied to the surface of the electrolyte of the co-sintered hydrogen electrode-electrolyte bi-layer for a period of time within a range of from 1 minutes to 15 minutes (e.g., 1 minute, 2 minutes, 5 minutes, 10 minutes, and 15 minutes) and then washed with deionized water. For comparison purposes, a control co-sintered hydrogen electrode-electrolyte bi-layer was not exposed to the surface treatment.

[0099] Surface roughness of the electrolyte of the co-sintered hydrogen electrode-electrolyte bi-layer was inspected using a Dimension FASTSCAN® atomic force microscope (AFM) from Bruker using a tapping mode. Referring to FIG. 4, the AFM scan of an untreated electrolyte surface is shown at a), the AFM scan of an electrolyte surface treated for 1 minute is shown at b), the AFM scan of an electrolyte surface treated for 2 minutes is shown at c), the AFM scan of an electrolyte surface treated for 5 minutes is shown at d), the AFM scan of an electrolyte surface treated for 10 minutes is shown at e), and the AFM scan of an electrolyte surface treated for 15 minutes is shown at f). Table 1 includes the surface roughness measured by conventional techniques for the electrochemical cells (e.g., untreated, treated for 1 minute, treated for 2 minutes, treated for 5 minutes, treated for 10 minutes, treated for 15 minutes).

TABLE-US-00001 TABLE 1 Surface Roughness of the Electrolyte Treatment Time (min)

Untreated	1	2	5	10	15	Surface Roughness	0.28	0.37	0.59	0.66	0.77	0.70 (μm)
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[0100] Without being bound by any theory, it is believed that the concentrated nitric acid reacts with an upper portion of the electrolyte layer to form nitrate. After the treatment, the electrolyte

surface is cleaned by deionized water to remove the residue including nitrate and extra nitric acid. The etching of the electrolyte surface by the nitric acid removes the well-annealed surface and increases the surface roughness of the electrolyte.

[0101] To fabricate full electrochemical cells, slurries of the oxygen electrode materials described in Example 1 were prepared by mixing the described oxygen electrode powders with ethanol and a texanol-based binder by ball milling. The slurries of the oxygen electrode materials were individually brush-painted on a surface of an electrolyte of a respective co-sintered hydrogen electrode-electrolyte bi-layer. For the 3D PNC73 mesh electrode, the heat-treated mesh was bonded to a surface of an electrolyte of a respective co-sintered hydrogen electrode-electrolyte layer using a thin layer of brush-painted PNC73 slurry. The surface area of the oxygen electrodes was controlled to be about 0.178 cm². The brush-painted cells were sintered at T_{sub.2}=1000° C. for 4 hours (heating rate: 3° C. min^{sup.-1}) and then allowed to cool.

Example 3

Cell Testing Assembly

[0102] Electrochemical cells including NiO and BZCYYb hydrogen electrodes and PNC55 oxygen electrodes as described in Examples 1 and 2 were sealed in a reactor using Aremco Ceramabond™ 552 adhesive with the oxygen electrode side of the electrochemical cells exposed. Silver mesh was used as a current collector with attached silver wires as leads. After assembly, the electrochemical cells were heated to 600° C. at a heating rate of 1° C. min^{sup.-1}. When a temperature of 600° C. was reached, H_{sub.2} gas was fed into the hydrogen electrodes at a flow rate of 20 mL min^{sup.-1} to reduce the NiO of the hydrogen electrodes to metallic Ni. After reducing the NiO to metallic Ni, a bubbler was connected to hydrolyze H_{sub.2} with 3 wt % steam.

Example 4

Electrochemical Impedance Spectroscopy (EIS) Measurements

[0103] EIS measurements were conducted under open-circuit voltage (OCV) at temperatures between 400° C. and 650° C. for electrochemical cells including Ni-BZCYYb hydrogen electrodes and PNC55 oxygen electrodes described in Examples 1-3 operating as fuel cells with feedstocks of H_{sub.2} and 3 wt % H_{sub.2}O for the hydrogen electrodes and O_{sub.2} for the oxygen electrodes. FIG. 5A illustrates a Cole-Cole plot obtained from the EIS measurements, where the x-axis is Z' and the y-axis is -Z'', where Z' and Z'' are the real and imaginary parts of the complex impedance, respectively. As shown by the Cole-Cole plot of FIG. 5A, the surface treatment described in Example 2 applied for a time up to 10 minutes lowered both ohmic resistance R_{sub.o} (shown by the left intercept of the EIS curve with the x-axis) and polarization resistance R_{sub.p} (shown by the right intercept of the EIS curve with the x-axis minus R_{sub.o}). Both R_{sub.o} and R_{sub.p} reached their corresponding minimum value in the electrochemical cell treated for 10 minutes, which correlates with the largest surface roughness of the electrolyte and highest interfacial strength between the electrolyte and the oxygen electrode. The decreased R_{sub.p} indicated an increase in activity of a triple-phase boundary between the electrolyte, oxygen electrode, and the O_{sub.2} gas feedstock of the oxygen electrode. R_{sub.o} and R_{sub.p} are assumed to follow an Arrhenius-type temperature dependence

$$[00005] R_o = A_o \exp\left(-\frac{E_{a,o}}{k_B T}\right) \quad (1) \quad R_p = A_p \exp\left(-\frac{E_{a,p}}{k_B T}\right) \quad (2)$$

[0104] Where A is the respective pre-exponent term, E_{sub.a} is the activation energy, the subscripts o and p denote that for R_{sub.o} and R_{sub.p}, respectively, k_{sub.B} is the Boltzmann constant, and T is the absolute temperature. The plot of log R_{sub.o} vs. 1/T illustrated in FIG. 5B and the plot of log R_{sub.p} vs. 1/T illustrated in FIG. 5C show the curves of the electrochemical cells treated with nitric acid for different times were parallel to each other, indicating that the treatment does not affect the activation energy of R_{sub.o} and R_{sub.p}. An activation energy of E_{sub.a,o}=0.358±0.001 eV for R_{sub.o} and E_{sub.a,p}=0.903±0.003 eV for R_{sub.p} were deduced for the analyzed electrochemical cells, as shown in the plot of calculated activation energy shown at FIG. 5D.

Instead, the treatment time modified $R_{\text{sub.o}}$ and $R_{\text{sub.p}}$ by changing the pre-exponential factor $A_{\text{sub.o}}$ and $A_{\text{sub.p}}$. This was distinct from the conventional practice of enhancing kinetics by lowering the activation energy.

[0105] The enhancement factor for $1/A_{\text{sub.o}}$ and $1/A_{\text{sub.p}}$ in the treated electrochemical cells over the pre-exponent term $1/A_{\text{sub.o}}^{\text{sup.0}}$ and $1/A_{\text{sub.p}}^{\text{sup.0}}$ in the untreated electrochemical cell was calculated and plotted in an $A_{\text{sub.o}}/A_{\text{sub.o}}^{\text{sup.0}}$ vs. $A_{\text{sub.p}}^{\text{sup.0}}/A_{\text{sub.p}}$ plot shown in FIG. 5E. A dimensionless reduced temperature $k_{\text{sub.BT}}/E_{\text{sub.a}}$ and a dimensionless relative “resistance” R were combined into a single curve in the Arrhenius plot shown in FIG. 5F. The EIS measurements suggest a single mechanism was responsible for the simultaneously lowered $R_{\text{sub.o}}$ and $R_{\text{sub.p}}$, which were lowered by lowering their pre-exponential factor without changing the mechanisms of electrode reactions and proton conduction (inferred by unchanged activation energy). Accordingly, the increased effective contact area between the oxygen electrode and electrolyte and resulting improved interfacial bonding were determined to be responsible for the lowered $R_{\text{sub.o}}$ and $R_{\text{sub.p}}$.

Example 5

Electrochemical PCFC and PCEC Testing

[0106] Electrochemical tests of the electrochemical cells including Ni-BZCYYb hydrogen electrodes and PNC55 oxygen electrodes described in Examples 1-3 operating as fuel cells (e.g., protonic ceramic fuel cells (PCFCs)) and operating as electrolysis cells (e.g., protonic ceramic electrolysis cells (PCECs)) were conducted. Current density-voltage curves and current density-power density curves of the electrochemical cells operating as fuel cells were measured using a Solartron **1400** and a Solartron **1470** electrochemical working station after a stable OCV was observed at the set temperature. When operated as fuel cells, the feedstocks were $\text{H}_{\text{sub.2}}$ and 3 wt % steam for the hydrogen electrode and pure $\text{O}_{\text{sub.2}}$ for the oxygen electrode. Electrochemical data for the electrochemical cells operating as fuel cells was collected for a voltage range from OCV to 0.2 V at the set temperature. Continuous operation as a fuel cell was conducted under a constant applied voltage of 0.75 V at 600° C. for up to 200 hours. After tests for operation as a fuel cell were completed, the electrochemical cells were switched to operate as electrolysis cells. When operated as electrolysis cells, the feedstocks were pure $\text{H}_{\text{sub.2}}$ for the hydrogen electrode and $\text{O}_{\text{sub.2}}$ and 30 wt % $\text{H}_{\text{sub.2}}\text{O}$ for the oxygen electrode. Electrochemical tests were conducted after a stable OCV was observed at the set temperature, for a voltage range from 1.5 V to OCV. Continuous operation as an electrolysis cell was conducted under a constant applied voltage of 1.4 V at 600° C. for up to 200 hours.

[0107] FIG. 6 is a current density vs. voltage plot (e.g., polarization curve) of the electrochemical cells operated as electrolysis cells under a constant applied voltage of 1.4 V at 600° C. When a voltage larger than OCV (e.g., 1.04 V at 600° C.) was applied, the absolute value of the current density j (defined as being negative for an electrolysis cell and positive for a fuel cell) was used to evaluate the electrochemical cell performance. As shown in FIG. 6, under the same applied voltage, a larger $|j|$ was achieved in the treated electrochemical cells than in the untreated electrochemical cell. A 2.8-fold increment over the untreated electrochemical cell was observed in the $|j|$ of the electrochemical cell treated for 10 minutes under 1.4 V at 600° C. ($|j|=3.07 \text{ A cm}^{\text{sup.-2}}$) during operation as an electrolysis cell.

[0108] FIG. 7A is a current density vs. Faradic efficiency plot for the untreated electrochemical cell and the electrochemical cell treated with nitric acid for 10 minutes operated as electrolysis cells. FIG. 7B is a voltage vs. hydrogen production rate plot for the untreated electrochemical cell and the electrochemical cell treated with nitric acid for 10 minutes operated as electrolysis cells. As shown in FIGS. 7A and 7B, the electrochemical cell treated for 10 minutes demonstrated a higher Faradic efficiency and a higher hydrogen production rate than the untreated electrochemical cell.

[0109] FIG. 8 is a current density vs. voltage plot (e.g., polarization curve) and a current density vs. power density plot (e.g., power density curve) of the electrochemical cells operated as fuel cells

under a constant applied voltage of 0.75 V at 600° C. As shown in FIG. 8, the current density was higher in the treated electrochemical cells than in the untreated electrochemical cell at the same voltage. A 2.5-fold increment over the untreated electrochemical cell was observed in the peak power density $P_{\text{sub.max}}$ of the electrochemical cell treated for 10 minutes ($P_{\text{sub.max}}=1.18 \text{ W cm.sup.-2}$) during operation as a fuel cell.

Example 6

Degradation Testing

[0110] Accelerated degradation tests were conducted under high-current-density operation of the electrochemical cells including Ni-BZCYYb hydrogen electrodes and PNC55 oxygen electrodes described in Examples 1-3 as electrolysis cells (e.g., PCECs). High-current-density PCEC operations are known to cause severe stability issues due to the harsh thermomechanical and electrochemical conditions. FIG. 9 is a testing time vs. current density plot (e.g., current density curve) of the untreated electrochemical cell and the electrochemical cell treated with nitric acid for 10 minutes operated as PCECs under a constant applied voltage of 1.4 V at 600° C. The electrochemical cell treated for 10 minutes exhibited a high initial j of about $-3.24 \text{ A cm.sup.-2}$ and was stable for over 200 hours of continuous operation. As shown in FIG. 9, $|j|$ of the electrochemical cell treated for 10 minutes exhibited a 0.94% decay between 0 hours and 100 hours of operation and a 0.05% decay between 100 hours and 200 hours of operation. In comparison, the untreated electrochemical cell exhibited a smaller initial j of about $-1.35 \text{ A cm.sup.-2}$. The untreated electrochemical cell additionally exhibited fast degradation, with a 10.2% decay in $|j|$ between 0 hours and 100 hours of operation and a 5.9% decay between 100 hours and 200 hours of operation. Post-testing analysis of the untreated electrochemical cell revealed severe delamination of the oxygen electrode after electrochemical cycling and a change in the fracture mode of the BZCYYb electrolyte layer from the intragranular-cracking behavior observed in an untested electrochemical cell. In comparison, post-testing analysis of the electrochemical cell treated with nitric acid for 10 minutes revealed no observable oxygen electrode delamination and the BZCYYb electrolyte layer retained the intragranular cracking behavior after 200 hours of operation.

Example 7

Peeling Strength

[0111] A peeling strength of the oxygen electrode-electrolyte interface of the electrochemical cells including Ni-BZCYYb hydrogen electrodes and PNC55 oxygen electrodes described in Examples 1-3 was measured. To measure the peeling strength of the oxygen electrode-electrolyte interface, double-sided tape was attached to both sides of electrochemical cells with a rectangular shape having a length of 1.18 inch and a width of 1 inch. The hydrogen electrode side of the electrochemical cells was attached to a testing bed with the oxygen electrode facing up and tape was attached to the double-sided tape on the oxygen electrode side. The oxygen electrode was then peeled off from the electrolyte while the force was recorded and converted to peeling strength. Table 2 includes the peeling strength observed for each of the electrochemical cells (e.g., untreated, treated for 1 minute, treated for 2 minutes, treated for 5 minutes, treated for 10 minutes, treated for 15 minutes).

TABLE-US-00002 TABLE 2 Peeling Strength of the Oxygen Electrode-Electrolyte Interface
Treatment Time (min) Untreated 1 2 5 10 15 Peeling Strength 18.6 18.8 19.9 22 23.5 20.2 (N)

Example 8

Oxygen Electrode Composition Testing

[0112] The electrochemical cells having a BZCYYb electrolyte exhibiting a thickness of about 16 μm and oxygen electrodes formed of one of LSCF, PBSCF, PNC55, PNC73, or 3D PNC73 mesh described in Examples 1-3 were operated as fuel cells at 600° C. using $\text{CH}_{\text{sub.4}}$ and $\text{C}_{\text{sub.2H}_{\text{sub.6}}}$ with 3 wt % steam at a flow rate of 20 mL min.sup.-1 as the feedstock for the hydrogen electrodes and pure $\text{O}_{\text{sub.2}}$ at a flow rate of 60 mL min.sup.-1 as the feedstock for the oxygen electrodes. FIGS. 10A-10E are current density vs. voltage and power density plots for each

electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte to a solution comprising nitric acid, hydrochloric acid, sulfuric acid, phosphoric acid, or a combination thereof.

[0119] Embodiment 4: The method of any of Embodiments 1 through 3 wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte to an acid solution comprising nitric acid.

[0120] Embodiment 5: The method of any of Embodiments 1 through 4, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte to an acid solution comprising a concentration of acid within a range of from about 60 wt % to about 80 wt %.

[0121] Embodiment 6: The method of any of Embodiments 1 through 5, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation for a period of time within a range of from about 5 minutes to about 15 minutes.

[0122] Embodiment 7: The method of any of Embodiments 1 through 6, wherein forming an electrolyte material on an electrode of an electrochemical cell comprises forming an electrolyte material on an electrode comprising a nickel/perovskite cermet.

[0123] Embodiment 8: The method of any of Embodiments 1 through 7, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation to increase a surface roughness of the electrolyte material comprises increasing the surface roughness of the electrolyte to a roughness within a range of from about 0.5 μm to about 1 μm .

[0124] Embodiment 9: The method of any of Embodiments 1 through 8, further comprising heating the electrolyte material on the electrode to a temperature greater than about 1300° C. for a period of time greater than about 3 hours.

[0125] Embodiment 10: A method of forming an electrochemical cell, the method comprising: forming an electrolyte material on a first electrode of an electrochemical cell, the electrolyte material comprising a perovskite material; exposing the electrolyte material to at least one acid solution to increase surface roughness of the electrolyte material; and after exposing the electrolyte material to the at least one acid solution, forming a second electrode on the electrolyte material.

[0126] Embodiment 11: The method of Embodiment 10, wherein exposing the electrolyte material to at least one acid solution comprises exposing the electrolyte material to a nitric acid solution comprising a concentration of nitric acid of about 70 wt %.

[0127] Embodiment 12: The method of Embodiment 10 or Embodiment 11, wherein forming a second electrode on the electrolyte material comprises forming a second electrode comprising at least one perovskite material on the electrolyte material.

[0128] Embodiment 13: The method of any of Embodiments 10 through 12, wherein forming an electrolyte material on a first electrode comprises forming an electrolyte material exhibiting an ionic conductivity greater than or equal to about 10^{-2} S/cm at one or more temperatures within a range of from about 150° C. to about 650° C. on the first electrode.

[0129] Embodiment 14: The method of any of Embodiments 10 through 13, wherein exposing the electrolyte material to at least one acid solution comprises removing a portion of the electrolyte material adjacent an exposed surface of the electrolyte material opposite the first electrode.

[0130] Embodiment 15: The method of any of Embodiments 10 through 14, further comprising heating the first electrode, the electrolyte, and the second electrode to diffusion bond the second electrode to the electrolyte material.

[0131] Embodiment 16: An electrochemical cell comprising: a first electrode; a second electrode; and a proton-conducting membrane between the first electrode and the second electrode, the proton-conducting membrane comprising a perovskite material, wherein an interface between the

proton-conducting membrane and the second electrode exhibits a peeling strength within a range of from about 17 N to about 40 N.

[0132] Embodiment 17: The electrochemical cell of Embodiment 16, wherein the perovskite material exhibits an ionic conductivity greater than or equal to about 10^{-2} S/cm at one or more temperatures within a range of from about 150° C. to about 650° C.

[0133] Embodiment 18: The electrochemical cell of Embodiment 16 or Embodiment 17, wherein: the first electrode comprises nickel and a yttrium- and ytterbium-doped barium-zirconate-cerate (BZCYYb); and the electrolyte material comprises BZCYYb.

[0134] Embodiment 19: The electrochemical cell of any of Embodiments 16 through 18, wherein the second electrode comprises $\text{PrNi}_{0.5}\text{Co}_{0.5}\text{O}_{3-\delta}$ (PNC55).

[0135] Embodiment 20: A system for H_2 gas production and electricity generation, comprising: at least one steam source, at least one electrochemical apparatus in fluid communication with the at least one steam source, and a power source electrically connected to the at least one electrochemical apparatus, the at least one electrochemical apparatus comprising one or more electrochemical cells of any of Embodiments 16 through 19.

[0136] While the disclosure is susceptible to various modifications and alternative forms, specific embodiments have been shown by way of example in the drawings and have been described in detail herein. However, the disclosure is not intended to be limited to the particular forms disclosed. Rather, the disclosure encompasses all modifications, equivalents, and alternatives falling within the scope of the disclosure as defined by the following appended claims and their legal equivalents.

Claims

1. A method of improving an interface between an electrode and an electrolyte of an electrochemical cell, the method comprising: forming an electrolyte material on an electrode of an electrochemical cell, the electrolyte material comprising a perovskite material; and exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation to increase a surface roughness of the electrolyte material.
2. The method of claim 1, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte material to an aqueous acid solution.
3. The method of claim 1, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte to a solution comprising nitric acid, hydrochloric acid, sulfuric acid, phosphoric acid, or a combination thereof.
4. The method of claim 1, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte to an acid solution comprising nitric acid.
5. The method of claim 1, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte to an acid solution comprising a concentration of acid within a range of from about 60 wt % to about 80 wt %.
6. The method of claim 1, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation comprises exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation for a period of time within a range of from about 5 minutes to about 15 minutes.
7. The method of claim 1, wherein forming an electrolyte material on an electrode of an electrochemical cell comprises forming an electrolyte material on an electrode comprising a nickel/perovskite cermet.

8. The method of claim 1, wherein exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation to increase a surface roughness of the electrolyte material comprises increasing the surface roughness of the electrolyte to a roughness within a range of from about 0.5 μm to about 1 μm .
 9. The method of claim 1, further comprising, before exposing the electrolyte material to one or more of an acid solution, a plasma, thermal shock, and gamma radiation, heating the electrolyte material on the electrode to a temperature greater than about 1300° C. for a period of time greater than about 3 hours.
 10. A method of forming an electrochemical cell, the method comprising: forming an electrolyte material on a first electrode of an electrochemical cell, the electrolyte material comprising a perovskite material; exposing the electrolyte material to at least one acid solution to increase surface roughness of the electrolyte material, and after exposing the electrolyte material to the at least one acid solution, forming a second electrode on the electrolyte material.
 11. The method of claim 10, wherein exposing the electrolyte material to at least one acid solution comprises exposing the electrolyte material to a nitric acid solution comprising a concentration of nitric acid of about 70 wt %.
 12. The method of claim 10, wherein forming a second electrode on the electrolyte material comprises forming a second electrode comprising at least one perovskite material on the electrolyte material.
 13. The method of claim 10, wherein forming an electrolyte material on a first electrode comprises forming an electrolyte material exhibiting an ionic conductivity greater than or equal to about 10.sup.-2 S/cm at one or more temperatures within a range of from about 150° C. to about 650° C. on the first electrode.
 14. The method of claim 10, wherein exposing the electrolyte material to at least one acid solution comprises removing a portion of the electrolyte material adjacent to an exposed surface of the electrolyte material opposite the first electrode.
 15. The method of claim 10, further comprising heating the first electrode, the electrolyte material, and the second electrode to diffusion bond the second electrode to the electrolyte material.
 16. An electrochemical cell comprising: a first electrode; a second electrode; and a proton-conducting membrane between the first electrode and the second electrode, the proton-conducting membrane comprising a perovskite material, wherein an interface between the proton-conducting membrane and the second electrode exhibits a peeling strength within a range of from about 17 N to about 40 N.
 17. The electrochemical cell of claim 16, wherein the perovskite material exhibits an ionic conductivity greater than or equal to about 10.sup.-2 S/cm at one or more temperatures within a range of from about 150° C. to about 650° C.
 18. The electrochemical cell of claim 16, wherein: the first electrode comprises nickel and a yttrium- and ytterbium-doped barium-zirconate-cerate (BZCYYb); and the proton-conducting membrane comprises BZCYYb.
 19. The electrochemical cell of claim 16, wherein the second electrode comprises PrNi.sub.0.5Co.sub.0.5O.sub.3- δ (PNC55).
 20. A system for H.sub.2 gas production and electricity generation, comprising: at least one steam source, at least one electrochemical apparatus in fluid communication with the at least one steam source, and a power source electrically connected to the at least one electrochemical apparatus, the at least one electrochemical apparatus comprising one or more electrochemical cells comprising: a first electrode; a second electrode; and a proton-conducting membrane between the first electrode and the second electrode, the proton-conducting membrane comprising a perovskite material, wherein an interface between the proton-conducting membrane and the second electrode exhibits a peeling strength within a range of from about 17 N to about 40 N.
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