



(51) International Patent Classification:

C25B 3/04 (2006.01) **H01M 4/90** (2006.01)
C25B 11/04 (2006.01) **H01M 4/86** (2006.01)
H01M 8/00 (2016.01) **G01N 27/30** (2006.01)

(21) International Application Number:

PCT/US2017/025624

(22) International Filing Date:

31 March 2017 (31.03.2017)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

15/090,477 4 April 2016 (04.04.2016) US
 15/158,227 18 May 2016 (18.05.2016) US

(71) Applicant: **DIOXIDE MATERIALS, INC.** [US/US];
 3998 Fau Boulevard, Suite 300, Boca Raton, Florida
 33431 (US).

(72) Inventors: **MASEL, Richard I.**; 6503 Las Flores, Boca
 Raton, Florida 33433 (US). **LIU, Zengcai**; 5771 Coach
 House Circle, Unit D, Boca Raton, Florida 33486 (US).
KUTZ, Robert; 6055 Town Colony Drive, Unit 1211,
 Boca Raton, Florida 33433 (US). **SAJJAD, Syed Dawar**;
 5573 Pacific Boulevard, Unit 3502, Boca Raton, Florida
 33433 (US).

(74) Agents: **CORRIDOR LAW GROUP, P.C.** et al.; 2135
 CityGate Lane, Suite 300, Naperville, Illinois 60563 (US).

(81) Designated States (unless otherwise indicated, for every
 kind of national protection available): AE, AG, AL, AM,
 AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY,
 BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM,
 DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT,
 HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KH, KN,
 KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA,
 MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG,
 NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS,
 RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY,
 TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN,
 ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every
 kind of regional protection available): ARIPO (BW, GH,
 GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ,
 TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU,
 TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE,
 DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU,
 LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK,
 SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
 GW, KM, ML, MR, NE, SN, TD, TG).

Published:

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the
 claims and to be republished in the event of receipt of
 amendments (Rule 48.2(h))

(54) Title: CATALYST LAYERS AND ELECTROLYZERS

(57) Abstract: A catalyst layer for an electrochemical device comprises a catalytically active element and an ion conducting polymer. The ion conducting polymer comprises positively charged cyclic amine groups. The ion conducting polymer comprises at least one of an imidazolium, a pyridinium, a pyrazolium, a pyrrolidinium, a pyrrolium, a pyrimidium, a piperidinium, an indolium, a triazinium, and polymers thereof. The catalytically active element comprises at least one of V, Cr, Mn, Fe, Co, Ni, Cu, Sn, Zr, Nb, Mo, Ru, Rh, Pd, Ag, Cd, Hf, Ta, W, Re, Ir, Pt, Au, Hg, Al, Si, In, Tl, Pb, Bi, Sb, Te, U, Sm, Tb, La, Ce and Nd. In an electrolyzer comprising the present catalyst layer, the feed to the electrolyzer comprises at least one of CO₂ and H₂O



WO 2017/176597 A1

CATALYST LAYERS AND ELECTROLYZERS

Field of the Invention

[0001] The field of the invention is electrochemistry. The devices, systems and compositions described involve the electrochemical conversion of carbon dioxide into useful products, the electrolysis of water, electric power generation using fuel cells and electrochemical water purification.

Background of the Invention

[0002] There is a desire to decrease carbon dioxide (CO₂) emissions from industrial facilities and power plants as a way of reducing global warming and protecting the environment. One solution, known as carbon sequestration, involves the capture and storage of CO₂. Often the CO₂ is simply buried. It would be valuable if instead of simply burying or storing the CO₂, it could be converted into another product and put to a beneficial use.

[0003] Over the years, a number of electrochemical processes have been suggested for the conversion of CO₂ into useful products. Some of these processes and their related catalysts are discussed in U.S. Patent Nos. 3,959,094; 4,240,882; 4,349,464; 4,523,981; 4,545,872; 4,595,465; 4,608,132; 4,608,133; 4,609,440; 4,609,441; 4,609,451; 4,620,906; 4,668,349; 4,673,473; 4,711,708; 4,756,807; 4,818,353; 5,064,733; 5,284,563; 5,382,332; 5,457,079; 5,709,789; 5,928,806; 5,952,540; 6,024,855; 6,660,680; 6,664,207; 6,987,134; 7,157,404; 7,378,561; 7,479,570; U.S. Patent App. Pub. No. 2008/0223727; Hori, Y., "Electrochemical CO₂ reduction on metal electrodes", *Modern Aspects of Electrochemistry* 42 (2008), pages 89-189; Gattrell, M. et al. "A review of the aqueous electrochemical reduction of CO₂ to hydrocarbons at copper", *Journal of Electroanalytical Chemistry* 594 (2006), pages 1-19; and DuBois, D., *Encyclopedia of Electrochemistry*, 7a, Springer (2006), pages 202-225.

[0004] Processes utilizing electrochemical cells for chemical conversions have been known for years. Generally, an electrochemical cell contains an anode, a cathode and an electrolyte. Catalysts can be placed on the anode, the cathode, and/or in the electrolyte to promote the desired chemical reactions. During operation, reactants or a solution containing reactants are fed into the cell. In an electrolytic cell, voltage is then applied between the anode and the

cathode to promote the desired electrochemical reaction. Note that the convention for designating the cathode and anode is that the cathode is the electrode at which chemical reduction occurs. This is different for electrolytic cells (also known as electrolyzers, devices in which electrical energy is supplied in order to force a chemical oxidation-reduction reaction) than for galvanic cells (such as fuel cells and batteries) in which a spontaneous electrochemical reaction produces electricity in an external circuit during normal operation. In either case, a chemical species acquires electrons from the cathode, thus becoming more negative and reducing its formal oxidation number.

[0005] When an electrochemical cell is used as a CO₂ conversion system, a reactant comprising CO₂, carbonate or bicarbonate is fed into the cell. A voltage is applied to the cell, and the CO₂ reacts to form new chemical compounds.

[0006] One of the issues at present is that to obtain high currents, one needs to run the electrochemical cells for conversion of CO₂ either at low voltage efficiency or with continuous additions of co-reactants. For example, Schmidt et al., Electrochemical Reduction of CO₂ (available at http://www.sccer-hae.ch/resources/Talks/08_Krause_Power_to_Value.pdf; last accessed on May 17, 2016) report that over 6 volts need to be applied to achieve a current of 600 mA/cm². This corresponds to an electrical efficiency of 21%. Verma, S. et al. Phys. Chem. Chem. Phys., 2016, 18: p. 7075-7084 find that they can achieve 400 mA/cm² at a cell potential of 3 V (42% electrical efficiency) by continuously supplying 3 M KOH to the cell. Unfortunately, if the KOH is recycled, the KOH will react with CO₂ to form KHCO₃. Verma et al. find that the current drops to 40 mA/cm² in KHCO₃.

[0007] An electrochemical cell for the conversion of CO₂ that achieves (a) reasonable energy efficiencies (at least 40%), (b) reasonable currents (above 150 mA/cm²), and (c) reasonable selectivities (above 50%), without needing to continuously introduce other reactants, would represent a significant advance in the electrochemical field.

Summary of the Invention

[0008] In one embodiment, an electrolyzer cathode catalyst layer allows high currents to be obtained at lower voltages. The catalyst layer contains one or more catalytically active chemical elements and an anion conducting polymer, wherein the ion conducting polymer is comprised of positively charged cyclic amine groups.

[0009] Preferably the cyclic amine is a one or more of imidazoliums, pyridiniums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums, triaziniums, and polymers thereof; more preferably the polymer is one or more of imidazoliums, pyridiniums and pyrazoliums.

[0010] Preferably said pyridiniums have no hydrogens directly bound to the nitrogen.

[0011] Preferably said imidazoliums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums, triaziniums, have at least one nitrogen that has no hydrogens directly bound to it.

[0012] Preferably such positive charged cyclic amine groups are aromatic.

[0013] Preferably the catalytic chemical element is chosen from the list: V, Cr, Mn, Fe, Co, Ni, Cu, Sn, Zr, Nb, Mo, Ru, Rh, Pd, Ag, Cd, Hf, Ta, W, Re, Ir, Pt, Au, Hg, Al, Si, In, Tl, Pb, Bi, Sb, Te, U, Sm, Tb, La, Ce, and Nd; more preferably the catalytically active element is chosen from the list Pt, Pd, Au, Ag, Cu, Ni, Fe, Sn, Bi, Co, In, Ru and Rh; most preferably the catalytically active element is chosen from the list Au, Ag, Cu, Sn, Sb, Bi, and In.

[0014] In a preferred embodiment, the weight of the ion conducting polymer in the catalyst layer is less than 64% of the weight of the catalytically active element. Most preferably the weight of the ion conducting polymer in the catalyst layer is between 1 and 10% of the weight of the catalytically active element.

[0015] In a preferred embodiment the catalyst layer also contains elemental carbon, most preferably carbon black, such as that available from Cabot Corporation, Boston, Massachusetts, USA, under the trade designation Vulcan XC-72R.

[0016] The catalyst layer can be part of an electrolyzer, fuel cell, battery or sensor.

[0017] In a preferred embodiment the catalyst layer is part of an electrolyzer. Electrolysis products may include CO , OH^- , HCO^- , H_2CO , $(\text{HCO}_2)^-$, HCOOH , H_2O_2 , CH_3OH , CH_4 , C_2H_4 , $\text{CH}_3\text{CH}_2\text{OH}$, CH_3COO^- , CH_3COOH , C_2H_6 , O_2 , H_2 , $(\text{COOH})_2$, and $(\text{COO}^-)_2$.

[0018] The feed to the electrolyzer may include at least one of CO_2 , HCO_3^- , CO_3^{2-} and H_2O .

[0019] The catalyst layer may be in electrical contact with the anode or cathode.

[0020] In one embodiment, the catalyst layer is part of a CO₂ electrolyzer. Preferably CO₂ is fed into the electrolyzer cathode and water is fed into the anode.

[0021] In a preferred embodiment, the water feed contains an electrolyte, preferably a carbonate, bicarbonate or hydroxide.

Brief Description of the Drawings

[0022] FIG. 1 is an exploded side view of a fuel cell hardware assembly including a membrane electrode assembly interposed between two fluid flow field plates having reactant flow channels formed in the major surfaces of the plates facing the electrodes.

[0023] FIG. 2 shows how the voltammograms of a 5 cm² cell change when 0% (200), 1% (201), 5% (202), and 10% (203) of PSMIMCl are added to the CO₂ electrolyzer cathode catalyst layer, where the percentage is calculated as the weight of the PSMIMCl divided by the weight of the silver.

[0024] FIG. 3 shows how the voltammograms of a 5 cm² CO₂ electrolyzer cell change when 1% (210), 4% (211), 8% (212), 16% (213) and 32% (214) of PSTMIMCl are added to the cathode catalyst layer.

[0025] FIG. 4 shows how the voltammograms of a 5 cm² CO₂ electrolyzer cell change when 5x% PSMP is added to the cathode catalyst layer.

[0026] FIG. 5 shows how the voltammograms of a 5 cm² CO₂ electrolyzer cell change when 5% of PSPY is added to the cathode catalyst layer.

[0027] FIG 6. Shows how the voltage (300) and selectivity (301) of a 5 cm² CO₂ electrolyzer varies with time when the cell is run at a constant current of 600 mA/cm².

Detailed Description of Illustrative Embodiments

[0028] It is understood that the process is not limited to the particular methodology, protocols and reagents described herein, as these can vary as persons familiar with the technology involved here will recognize. It is also to be understood that the terminology used herein is used for the purpose of describing particular embodiments only, and is not intended to limit the scope of the process. It also is to be noted that as used herein and in the appended claims, the singular forms “a,” “an,” and “the” include the plural reference unless the context clearly

dictates otherwise. Thus, for example, a reference to “a linker” is a reference to one or more linkers and equivalents thereof known to those skilled in the art. Similarly, the phrase “and/or” is used to indicate one or both stated cases can occur, for example, A and/or B includes (A and B) and (A or B).

[0029] Unless defined otherwise, technical and scientific terms used herein have the same meanings as commonly understood by one of ordinary skill in the art to which the process pertains. The embodiments of the process and the various features and advantageous details thereof are explained more fully with reference to the non-limiting embodiments and/or illustrated in the accompanying drawings and detailed in the following description. It should be noted that the features illustrated in the drawings are not necessarily drawn to scale, and features of one embodiment can be employed with other embodiments as the skilled artisan would recognize, even if not explicitly stated herein.

[0030] Any numerical value ranges recited herein include all values from the lower value to the upper value in increments of one unit, provided that there is a separation of at least two units between any lower value and any higher value. As an example, if it is stated that the concentration of a component or value of a process variable such as, for example, size, angle, pressure, time and the like, is, for example, from 1 to 98, specifically from 20 to 80, more specifically from 30 to 70, it is intended that values such as 15 to 85, 22 to 68, 43 to 51, 30 to 32, and the like, are expressly enumerated in this specification. For values which are less than one, one unit is considered to be 0.0001, 0.001, 0.01 or 0.1 as appropriate. These are only examples of what is specifically intended and all possible combinations of numerical values between the lowest value and the highest value are to be treated in a similar manner.

[0031] Moreover, provided immediately below is a “Definitions” section, where certain terms related to the articles and process are defined specifically. Particular methods, devices, and materials are described, although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of the process or articles.

Definitions

[0032] The term “electrochemical conversion of CO₂” as used here refers to any electrochemical process where carbon dioxide, carbonate, or bicarbonate is converted into another chemical substance in any step of the process.

[0033] The term polymer electrolyte membrane refers to both cation exchange membranes, which generally comprise polymers having multiple covalently attached negatively charged groups, and anion exchange membranes, which generally comprise polymers having multiple covalently attached positively charged groups. Typical cation exchange membranes include proton conducting membranes, such as the perfluorosulfonic acid polymer available under the trade designation NAFION from E. I. du Pont de Nemours and Company (DuPont) of Wilmington, DE.

[0034] The term anion exchange polymer refers to polymers having multiple covalently attached positively charged groups.

[0035] The term “anion exchange membrane electrolyzer” as used here refers to an electrolyzer with an anion-conducting polymer electrolyte membrane separating the anode from the cathode.

[0036] The term "Hydrogen Evolution Reaction" also called "HER" as used here refers to the electrochemical reaction $2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2$.

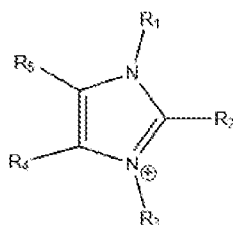
[0037] The term "MEA" as used here refers to a membrane electrode assembly, which typically comprises at least an ion-conducting membrane having an anode layer attached or in close proximity to one face of the membrane and a cathode layer attached or in close proximity to the other side of the membrane.

[0038] The Term “CV” refers to cyclic voltammetry or cyclic voltammogram.

[0039] The term "Millipore water" is water that is produced by a Millipore filtration system with a resistivity of at least 18.2 megohm-cm.

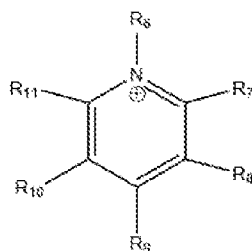
[0040] The term “GC” as used here refers to a gas chromatograph.

[0041] The term "imidazolium" as used here refers to a positively charged ligand containing an imidazole group. This includes a bare imidazole or a substituted imidazole. Ligands of the form:



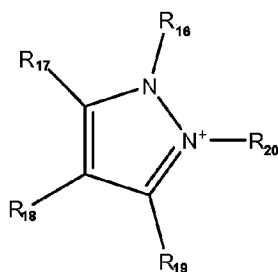
where R₁- R₅ are each independently selected from hydrogen, halogens, linear alkyls, branched alkyls, cyclic alkyls, heteroalkyls, aryls, cyclic aryls, heteroaryls, alkylaryls, heteroalkylaryls, and polymers thereof, such as the vinyl benzyl copolymers described herein, are specifically included.

[0042] The term "pyridinium" as used here refers to a positively charged ligand containing a pyridinium group. This includes a protonated bare pyridine or a substituted pyridine or pyridinium. Ligands of the form



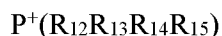
where R₆-R₁₁ are each independently selected from hydrogen, halogens, linear alkyls, branched alkyls, cyclic alkyls, heteroalkyls, aryls, cyclic aryls, heteroaryls, alkylaryls, heteroalkylaryls, and polymers thereof, such as the vinyl benzyl copolymers described herein, are specifically included.

[0043] The term "pyrazoliums" as used here refers to a positively charged ligand containing a pyrazolium group. This includes a bare pyrazolium or a substituted pyrazolium. Ligands of the form



where R₁₆-R₂₀ are each independently selected from hydrogen, halogens, linear alkyls, branched alkyls, cyclic alkyls, heteroalkyls, aryls, cyclic aryls, heteroaryl, alkylaryls, heteroalkylaryls, and polymers thereof, such as the vinyl benzyl copolymers described herein, are specifically included.

[0044] The term "phosphonium" as used here refers to a positively charged ligand containing phosphorous. This includes substituted phosphorous. Ligands of the form:

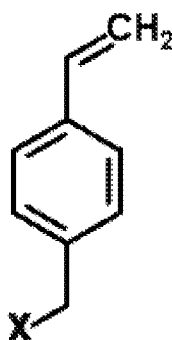


where R₁₂-R₁₅ are each independently selected from hydrogen, halogens, linear alkyls, branched alkyls, cyclic alkyls, heteroalkyls, aryls, cyclic aryls, heteroaryl, alkylaryls, heteroalkylaryls, and polymers thereof, such as the vinyl benzyl copolymers described herein, are specifically included.

[0045] The term "positively charged cyclic amine" as used here refers to a positively charged ligand containing a cyclic amine. This specifically includes imidazoliums, pyridiniums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums, triaziniums, and polymers thereof, such as the vinyl benzyl copolymers described herein.

[0046] The term "electrochemical device" as used here refers to a device capable of either generating electrical energy from chemical reactions or facilitating chemical reactions through the introduction of electrical energy. Batteries, fuel cells, electrolyzers, and electrochemical reactors are specifically included.

[0047] The term "vinyl benzyl derivatives" as used here refers to a chemical of the form.



or polymers thereof where X is hydrogen, halogens, linear alkyls, branched alkyls, cyclic alkyls, heteroalkyls, aryls, cyclic aryls, heteroaryls, alkylaryls, heteroalkylaryls, imidazoliums, pyridiniums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums, or triaziniums. Polymers thereof, such as the vinyl benzyl copolymers described herein, are specifically included.

Specific Description

[0048] One embodiment of the advance is an electrolyzer cathode catalyst layer that improves the output of a CO₂ electrolyzer.

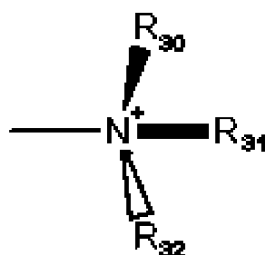
[0049] FIG. 1 illustrates a fuel cell hardware assembly 30, which includes a membrane electrode assembly 32 interposed between rigid flow field plates 34 and 36, the flow fields typically being formed of graphite or a graphite composite material. Membrane electrode assembly 32 consists of a polymer electrolyte (ion exchange) membrane 42 interposed between two electrodes, namely, anode 44 and cathode 46. Anode 44 and cathode 46 are typically formed of porous electrically conductive sheet material, preferably carbon fiber paper, and have planar major surfaces. Electrodes 44 and 46 have a thin layer of catalyst material disposed on their major surfaces at the interface with membrane 42 to render them electrochemically active.

[0050] A distinguishing feature here is the addition of anion conducting polymers to the catalyst layer to improve the cell output.

[0051] There are many previous descriptions of catalyst layers for electrochemical cells. Patents include U.S. Patent Nos. 5,234,777; 5,869,416; 6,156,449; 6,696,382; 6,800,391; 6,844,286; 7,364,813; 7,754,369; 7,754,369; 7,855,160; 7,906,452; 8,198,206; 8,481,231; 8,940,460; 9,127,182; and 9,160,008; U.S. Pat. App. Pub. Nos. 2002/0034674;

2002/0098405; 2004/0023104; 2004/0107869; 2005/0151121; 2006/0110631; 2008/0248944; 2010/0196785; 2010/0285951; 2011/0003071; 2011/0166009; 2011/0262828; 2012/0094210; 2012/0148936; 2012/0171583; 2012/0196741; 2012/0258381; 2013/0260278; 2014/0162170; 2014/0220474; and 2014/0228200; and International Publication Nos. WO2015/092371 and WO2015/124250. However, in most cases acidic polymers such as perfluorosulfonic acids, including those available from DuPont under the trade designation Nafion, are used. Acidic polymers substantially decrease the selectivity of CO₂ electrolyzers, so they are typically not useful in practical CO₂ electrolyzers.

[0052] There are a few reports of anionic fuel cells with anion conducting polymers having multiple covalently attached positively charged groups in their catalyst layer. See for example U.S. Patent Nos. 3,403,054; 7,785,750; and 8,257,872; U.S. Pat. App. Pub. No. 20150171453; International Publication Nos. WO/2013/0137011 and WO/2012/078513; and Matsuoka et al., Journal of Power Sources, Volume 150, 4 October 2005, pages 27-31. However, these authors use polymers containing amines of the form:



where the bond on the left is an attachment to the polymer and the R groups are either, hydrogens, methyls or ethyls, not positively charged cyclic amines such as imidazolium or pyrazolium. Si et al. (J. Mater. Chem. A, 2014, 2: p. 4413-4421.), Yan et al (J. Power Sources, 2014, 250: p. 90-97.), and Schauer et al. (the Schauer paper) Journal of Applied Polymer Science, 2015, 132: 42581, disclose imidazole functionalized poly(arylene ether sulfone) and poly(ether ketone) polymers, but Schauer et al. Figure 2 shows that these polymers are unstable under alkaline conditions so they are unsuitable for use in alkaline electrolyzers.

[0053] U.S. Patent No. 6,841,285, notes that, “Thus imidazole and pyrazole may act as both hydrogen donors and acceptors in proton conduction processes. While these compounds may show increased conductivity within membrane systems, it is unlikely that they are suitable for

use within the fuel cell environment. For example, a recent study by C. Yang et al., Journal of Power Sources 103:1, 2001, reports that imidazole impregnated membranes poisoned the catalysts.”

[0054] Note that imidazole and pyrazole would be protonated under the acidic conditions in U.S. Patent No. 6,841,285 thus forming imidazolium and pyrazolium ions, so that the implication of the ‘285 patent is that imidazoliums and pyrazoliums should not be used in a catalyst layer.

[0055] The present work shows that in contrast to the implication in the ‘285 patent, polymers containing imidazoliums, pyrazoliums and pyridiniums enhance the performance of a CO₂ electrolyzer.

[0056] The catalyst layer can also include at least one Catalytically Active Element. “Catalytically Active Element” as used here refers to a chemical element that can serve as a catalyst for the electrochemical conversion of CO₂ or another species of interest in a desired reaction. In particular, the device can include one or more of the following Catalytically Active Elements: V, Cr, Mn, Fe, Co, Ni, Cu, Sn, Zr, Nb, Mo, Ru, Rh, Pd, Ag, Cd, Hf, Ta, W, Re, Ir, Pt, Au, Hg, Al, Si, In, Tl, Pb, Bi, Sb, Te, U, Sm, Tb, La, Ce, and Nd. Research has established that Pt, Pd, Au, Ag, Cu, Ni, Fe, Sn, Bi, Co, In, Ru and Rh perform well, with Au, Ag, Cu, Sn, Sb, Bi, and In perform especially well.

[0057] Embodiments of the present invention also include the addition of electrically conductive species to the catalyst layer, with carbon being a conductive preferred component.

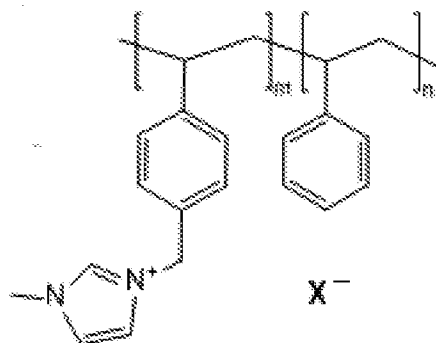
[0058] Without further elaboration, it is believed that persons familiar with the technology involved here using the preceding description can utilize the invention to the fullest extent. The following examples are illustrative only, and are not meant to be an exhaustive list of all possible embodiments, applications or modifications of the invention.

Specific Example 1

[0059] The objective of this example is to show that in contrast to the indication in US 6,841,285, the addition of polymers containing imidazoliums to the catalyst layer enhances the performance of a CO₂ electrolyzer.

[0060] A copolymer, which is designated here as PSMIM (Cl), was prepared following the

synthetic route in patent application 14/704,935. “PSMIM” refers to a co-polymer of polystyrene and poly 1-(p-vinylbenzyl)-3-methyl-imidazolium:



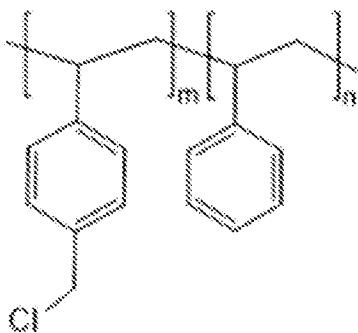
where X^- is an anion, $m > 0$ and $n > 0$.

[0061] The inhibitor-free styrene was prepared by passing styrene (Sigma-Aldrich, Saint Louis, MO) through the tert-butylcatechol (TBC) inhibitor remover (Sigma-Aldrich 311340). In general, 40 ml of remover is sufficient to yield 50 ml of clear, inhibitor free styrene. Inhibitor TBC in 4-vinylbenzyl chloride (4-VBC) was removed by the same inhibitor remover in a similar fashion.

[0062] Poly(4-vinylbenzyl chloride-co-styrene) was then synthesized by heating a solution of inhibitor-free styrene (Sigma-Aldrich) (36.139 g, 350 mmol) and 4-vinylbenzyl chloride (Sigma-Aldrich) (29.7272 g, 190 mmol) in chlorobenzene (Sigma-Aldrich) (45 ml) at 60-65°C in an oil bath for approximately 20 hours under argon gas with AIBN (α,α' -Azoisobutyronitrile, Sigma-Aldrich) (0.5927 g, 0.90 wt% based on the total monomers' weight) as initiator. The copolymer was precipitated in CH_3OH (methanol) and dried under vacuum.

[0063] “Polystyrene methylimidazolium chloride” (PSMIM) was synthesized by adding 1-methylimidazole (Sigma-Aldrich) (2.8650 g, 0.0349 mmol), which is an alkylimidazole, to the solution of the poly(4-VBC-co-St) (5.0034 g, 19.4 mmol) in anhydrous *N,N*-Dimethylformamide (DMF) (Sigma-Aldrich) (30 mL). The mixture was then stirred at around 30°C for around 50 hours to form a PSMIM solution.

[0064] “4-VBC-co-St” or “poly(4-vinylbenzyl chloride co-styrene)” as used here refers to a co-polymer of styrene and 4-vinylbenzyl chloride:



[0065] PSMIM-DVB was synthesized starting with poly(4-vinylbenzyl chloride co-styrene.) 1-methylimidazole (Sigma-Aldrich) (3.912 g, 47.7 mmol) was added in a 250 ml 3-neck round bottom flask to the solution of the poly(4-VBC-co-St) (15.358 g, 59.8mmol) in anhydrous N,N-Dimethylformamide (DMF) (Sigma-Aldrich) (105 mL). 0.22 ml of a divinylbenzene (DVB) in DMF solution (DVB concentration = 0.0083 g/ml) was carefully added through a pipette to the mixture with continual magnetic stirring. After this, 0.22 ml of AIBN-DMF solution (AIBN concentration = 0.0083 g/ml) was added to the mixture in a similar fashion. The reaction was then kept under nitrogen atmosphere at 50°C for about 60 hours. PSMIM-DVB was obtained as a white powder after purification by precipitation into diethyl ether.

[0066] Membranes were prepared by casting the PSMIM-DVB solution prepared above directly onto a flat glass surface. The thickness of the solution on the glass was controlled by a film applicator (MTI Corporation, Richmond, CA) with an adjustable doctor blade. The membranes were then dried in a vacuum oven in the following step wise fashion. They were first kept at 80°C for 120 minutes, then at 100°C for 60 minutes, at 120°C for 30 minutes and finally at 150°C for 60 minutes. Chloride ions in the membranes were removed by soaking the membranes in 1 M KOH solution for 24 hours or longer.

[0067] The cathode layer in Specific Example 1 was prepared as follows. Silver ink was made by mixing 100 mg of silver nanoparticles (20-40 nm, 45509, Alfa Aesar, Ward Hill, MA), 5 mg porous carbon (Vulcan XC-72R, Fuel Cell Earth, Woburn, MA) and different amounts of PSMIM-Cl in 3 ml of ethanol (459844, Sigma-Aldrich, St. Louis, MO). The mixture was then sonicated for 10 minutes. The silver ink was painted onto a gas diffusion layer (Sigracet 35 BC GDL, Ion Power Inc., New Castle, DE) covering an area of 6 cm x 6 cm. The electrode was immersed in 1 M KOH for at least 1 hour so that PSMIM-Cl converted by ion exchange to PSMIM-OH. Then the electrode was cut into 2.5 cm x 2.5 cm

sections for cell testing.

[0068] The anode in Specific Example 1 was prepared as follows: 100 mg of IrO₂ (43396, Alfa Aesar, Ward Hill, MA) was dispersed in the mixture of 1 ml of deionized water, 2 ml of isopropanol (3032-16, Macron Fine Chemicals, Avantor Performance Materials, Center Valley, PA) and 0.1 ml of 5wt.% poly-tetrafluoroethylene (PTFE) dispersion (665800, Sigma-Aldrich, St. Louis, MO). The mixture was sonicated for 10 min using a water bath sonicator. The ink was painted onto 6 cm X 6 cm of carbon fiber paper (Toray Paper 120, Fuel Cell Earth, Woburn, MA). The actual IrO₂ loading was about 2 mg/cm². The electrode was cut into 3 cm X 3 cm sections for cell testing.

[0069] The membrane was sandwiched between the anode and the cathode with the metal-containing layers on the anode and cathode facing the membrane, and the whole assembly was mounted in a Fuel Cell Technologies 5 cm² fuel cell hardware assembly with serpentine flow fields.

[0070] CO₂ humidified at 25°C was fed into the cathode flow field at a rate of 20 sccm, and 10 mM KHCO₃ was fed into the anode flow field. The cyclic voltammograms were collected by scanning the cell potential from 1.2 to 3.0 V. All of the scans were made at room temperature and atmospheric pressure.

[0071] FIG. 2 shows the results. Plot 200 is a base case with no PSMIM in the cathode catalyst layer ink. Notice that the current increases when PSMIM is added to the catalyst layer in a later sample, such that the PSMIM weight is 1% of the weight of the silver (plot 201). Further increases in the current are seen as the PSMIM concentration is increased so that the PSMIM weight is 5% of the weight of the silver (plot 202). Then there is a small decrease when the weight of the PSMIM is increased to 10% of the weight of the silver (plot 203). A decrease in the selectivity of the reaction was observed.

[0072] A run in which the PSMIM weight was 20% of the weight of the silver was also performed. The cell showed a small current, but analysis of the exit stream did not show significant CO₂ conversion.

[0073] These results demonstrate that the addition of an ionomer containing an imidazolium enhances the performance of a CO₂ electrolyzer, in contrast to the findings in U.S. Patent Nos. 6,841,285.

Specific Example 2

[0074] The objective of this example is to show that in contrast to the findings in U.S. Patent Nos. 6,841,285, the addition of polymers containing tetra-methyl-imidazolium to the catalyst layer enhances the performance of a CO₂ electrolyzer.

[0075] Preparation of PSTMIM: poly(4-vinylbenzyl chloride-co-styrene) was prepared as in Specific Example 1. Tetra-methyl-imidazolium (TCI) (5.934 g) was added to the solution of the poly(4-VBC-co-St) (10 g) in anhydrous N,N-Dimethylformamide (DMF) (Sigma-Aldrich) (85 mL). The mixture was stirred at 30-35°C for around 60 hours. PSTMIM was obtained as white solid particles after purification by precipitation into diethyl ether. PSTMIM refers to a material that contains a co-polymer of styrene and 1-(p-vinylbenzyl)-tetra-methyl-imidazolium.

[0076] The cathode in Specific Example 2 was prepared as follows. Silver ink was made by mixing 100 mg of silver nanoparticles (20-40 nm, 45509, Alfa Aesar, Ward Hill, MA), 5 mg porous carbon (Vulcan XC-72R, Fuel Cell Earth, Woburn, MA) and different amounts of PSTMIM-Cl in 3 ml of ethanol (459844, Sigma-Aldrich, St. Louis, MO). The mixture was then sonicated for 10 minutes. The silver ink was painted onto a gas diffusion layer (Sigracet 35 BC GDL, Ion Power Inc., New Castle, DE) covering an area of 6 cm x 6 cm. The electrode was immersed in 1 M KOH for at least 1 hour so that PSTMIM-Cl converted to PSTMIM-OH. Then the electrode was cut into 2.5 cm X 2.5 cm for cell testing.

[0077] The anode in specific example 2 was the same as in Specific Example 1 and the cell was tested as in Specific Example 1.

[0078] FIG. 3 shows the voltammograms measured at varying PSTMIM content. Notice that the current is higher than the base case (plot 200) in FIG. 2 when PSTMIM is added to the catalyst later so that the PSTMIM weight is 1% of the weight of the silver (plot 210). Further increases in the current are seen as the PSTMIM concentration is increased so that the PSTMIM weight is 4% of the weight of the silver (plot 211). Then there is a small decrease when the weight of the PSTMIM is increased to 8% of the weight of the silver (plot 212). The performance continues to decrease when the weight of the PSTMIM is 16% of the weight of the silver (plot 213), but the cell current is still higher than the base case in FIG. 2 (plot 200). The performance is lower than the base case in FIG. 2 (plot 200) when the weight of the PSTMIM is 32% of the weight of the silver (plot 214), but the cell current is still

significant.

[0079] A run in which the PSTMIM weight was 64% of the weight of the silver was also performed. The cell showed a small current, but analysis of the exit stream did not show significant CO₂ conversion. These results demonstrate that the addition of an ionomer containing tetra-methyl-imidazolium enhances the performance of a CO₂ electrolyzer, in contrast to the findings in the '285 patent.

Specific Example 3

[0080] The objective of this example is to show that in contrast to the findings in US 6,841,285, the addition of polymers containing pyridiniums to the catalyst layer changes the performance of a CO₂ electrolyzer but does not poison it.

[0081] Preparation of PSMP: poly(4-vinylbenzyl chloride co-styrene) was prepared as in Specific Example 1. Pyridine (Sigma-Aldrich) (0.318 g, 4.68 mmol) was added to the solution of the poly(4-VBC-co-St) (1 g, 3.89 mmol) in anhydrous N,N-Dimethylformamide (DMF) (Sigma-Aldrich) (8 mL). The mixture was stirred at room temperature for 60 hours, and PSMP was obtained as a white solid after purification by precipitation into diethyl ether. PSMP refers to a material that contains a co-polymer of styrene and 1-(p-vinylbenzyl)-pyridinium.

[0082] The cathode in Specific Example 3 was prepared as follows. Silver ink was made by mixing 100 mg of silver nanoparticles (20-40 nm, 45509, Alfa Aesar, Ward Hill, MA), 5 mg porous carbon (Vulcan XC-72R, Fuel Cell Earth, Woburn, MA) and different amounts of PSMP-Cl in 3 ml of ethanol (459844, Sigma-Aldrich, St. Louis, MO). The mixture was then sonicated for 10 minutes. The silver ink was painted onto a gas diffusion layer (Sigracet 35 BC GDL, Ion Power Inc., New Castle, DE) covering an area of 6 cm x 6 cm. The electrode was immersed in 1 M KOH for at least 1 hour so that PSMP-Cl was converted to PSMP-OH. Then the electrode was cut into 2.5 cm X 2.5 cm sections for cell testing.

[0083] The anode in Specific Example 3 was the same as in Specific Example 1 and the cell was tested as in Specific Example 1.

[0084] FIG. 4 shows the voltammograms measured when the weight of the PSMP was 5% of the weight of the silver. Notice that the current is considerably above the base case (200) in FIG 2. These results demonstrate that the addition of an ionomer containing a pyridinium

enhances the performance of a CO₂ electrolyzer, in contrast to the findings in the '285 patent.

Specific Example 4

[0085] The objective of this example is to show that in contrast to the findings in the '285 patent, the addition of polymers containing pyrazoliums to the catalyst layer enhances the performance of a CO₂ electrolyzer.

[0086] Preparation of PSPZ: poly(4-vinylbenzyl chloride co-styrene) was prepared as in Specific Example 1. Pyrazole (Sigma-Aldrich) (0.593 g, 4.67 mmol) was added to the solution of the poly(4-VBC-co-St) (1 g, 3.89 mmol) in anhydrous N,N-Dimethylformamide (DMF) (Sigma-Aldrich) (8 mL). The mixture was stirred at room temperature for 60 hours and this PSPY-DMF was accordingly further diluted for use as the ionomer. PSPY refers to a material that contains a co-polymer of styrene and 1-(p-vinylbenzyl)-pyrazolium ionomers.

[0087] The anode in Specific Example 4 was the same as in Specific Example 1 and the cell was tested as in Specific Example 1.

[0088] FIG. 5 shows a voltammogram measured when the weight of the PSPY was 5% of the weight of the silver. Notice that the current is considerably above the base case (200) in FIG 2. These results demonstrate that the addition of an ionomer containing a pyrazolium enhances the performance of a CO₂ electrolyzer, in contrast to the findings in the '285 patent.

Specific Example 5

[0089] The objective of this example is to show that the addition of a PSMIM to the cathode of a CO₂ electrolyzer also improves the steady state performance of the electrolyzer.

[0090] The anode and cathode for this test were synthesized as in Specific Example 1. The weight of the PSMIM in the cathode layer was 2% of the weight of the silver.

[0091] A PSTMIM-DVB membrane was used in this experiment. The preparation of the PSTMIM-DVB membrane is as follows: Poly(4-vinylbenzyl chloride co-styrene) was formed as in specific example 1. Tetramethylimidazole (TMIM) (TCI) (4.05 g, 32.6 mmol) was added in a 250 ml 3-neck round bottom flask to the solution of the poly(4-VBC-co-St) (10 g, 38.9 mol) in anhydrous N,N-Dimethylformamide (DMF) (Sigma-Aldrich) (73 mL). After the TMIM was thoroughly dissolved within this reaction mixture, 1 mL of a DVB-DMF solution (DVB concentration = 0.052 g/ml) was carefully added through a pipette to the mixture with

continual magnetic stirring. After this, 1 ml of AIBN-DMF solution (AIBN concentration = 0.00135 g/ml) was added to the mixture in a similar fashion. The reaction was then kept under nitrogen atmosphere at 50°C for about 60 hours. PSTMIM was obtained as a white powder after purification by precipitation into diethyl ether. A PSTMIM-DVB membrane was then formed as in Specific Example 1.

[0092] The cell was assembled and tested as in Specific Example 1.

[0093] FIG. 6 shows how the voltage and selectivity varied with time when the cell was held at a fixed current of 3 amps (600 mA/cm²). Notice that the cell is producing 600 mA/cm² at 3.3 volts. GC analysis showed that the cell was converting CO₂ to CO with over 95% selectivity to CO. By comparison, the Schmidt paper (referred to above, available at http://www.sccer-hae.ch/resources/Talks/08_Krause_Power_to_Value.pdf; last accessed on May 17, 2016) requires over 6 volts to reach the same current. Consequently, our cell would only need about half as much energy as Schmidt et al.'s to convert CO₂ at a rate corresponding to 600 mA/cm². The results are stable. The loss of performance reported in the Schauer paper is not seen.

[0094] More generally, notice that Specific Examples 1, 2, 3, 4 and 5 show that adding three different ion conducting polymer containing positively charged cyclic amine groups to the catalyst layer in a CO₂ electrolyzer enhance the performance of a CO₂ electrolyzer, in contrast to the findings in the '285 patent. This observation is believed to be general. That is, adding ion conducting polymer containing positively charged cyclic amine groups can enhance the performance of electrolyzers and other devices.

[0095] While particular elements, embodiments and applications of the present invention have been shown and described, it will be understood that the invention is not limited thereto since modifications can be made by those skilled in the art without departing from the scope of the present disclosure, particularly in light of the foregoing teachings.

[0096] The examples given above are merely illustrative and are not meant to be an exhaustive list of all possible embodiments, applications or modifications of the present electrochemical device. Thus, various modifications and variations of the described articles, methods and systems of the invention will be apparent to those skilled in the art without departing from the scope and spirit of the invention. Although the invention has been described in connection with specific embodiments, it should be understood that the

invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the described modes for carrying out the invention which are obvious to those skilled in the chemical arts or in the relevant fields are intended to be within the scope of the appended claims.

CLAIMS

What is claimed is:

1. An electrolyzer comprising a catalyst layer, the catalyst layer comprising a catalytically active element and an ion conducting polymer, wherein said ion conducting polymer comprises positively charged cyclic amine groups, and further wherein a reactant stream comprising carbon dioxide is in contact with said catalyst layer.
2. The electrolyzer of claim 1, wherein said ion conducting polymer comprises one or more of imidazoliums, pyridiniums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums or triaziniums.
3. The electrolyzer of claim 2, wherein said ion conducting polymer comprises one or more of imidazoliums, pyridiniums or pyrazoliums.
4. The electrolyzer of claim 2, wherein said ion conducting polymer also comprises at least one of styrene, vinyl benzyl derivatives or polymers thereof.
5. The catalyst layer of claim 1, wherein said catalytically active element comprises one or more of V, Cr, Mn, Fe, Co, Ni, Cu, Sn, Zr, Nb, Mo, Ru, Rh, Pd, Ag, Cd, Hf, Ta, W, Re, Ir, Pt, Au, Hg, Al, Si, In, Tl, Pb, Bi, Sb, Te, U, Sm, Tb, La, Ce or Nd.
6. The electrolyzer of claim 5, wherein said catalytically active element comprises one or more of Au, Ag, Cu, Sn, Sb, Bi or In.
7. The electrolyzer of claim 2, said electrolyzer comprising an anode, a cathode, a polymeric ion conducting membrane interposed therebetween, the reactant stream comprising carbon dioxide in contact with said cathode, and a source of electrical power for directing electrical current from said cathode to said anode.
8. The electrolyzer of claim 7, wherein said ion conducting membrane comprises a polymer comprising one or more of imidazoliums, pyridiniums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums or triaziniums.
9. The electrolyzer of claim 8, wherein said ion conducting membrane comprises an imidazolium.

10. The electrolyzer of claim 9, wherein said imidazolium is an alkyl-imidazolium.
11. The electrolyzer of claim 10, wherein said alkyl-imidazolium is one of 1-methyl-imidazolium or tetra-methyl-imidazolium.
12. The electrolyzer of claim 11, wherein said alkyl-imidazolium is tetra-methyl-imidazolium.
13. A catalyst layer for an electrochemical device comprising a catalytically active element and an ion conducting polymer wherein said ion conducting polymer comprises:
 - (a) positively charged cyclic amine groups; and
 - (b) polymers or copolymers of styrene
14. The catalyst layer of claim 13, where said ion conducting polymer comprises one or more of imidazoliums, pyridiniums, pyrazoliums, pyrrolidiniums, pyrroliums, pyrimidiums, piperidiniums, indoliums, or triaziniums.
15. The catalyst layer of claim 14, wherein said ion conducting polymer comprises one or more of imidazoliums, pyridiniums, or pyrazoliums.
16. The catalyst layer of claim 14, wherein said catalytically active element comprises one or more of V, Cr, Mn, Fe, Co, Ni, Cu, Sn, Zr, Nb, Mo, Ru, Rh, Pd, Ag, Cd, Hf, Ta, W, Re, Ir, Pt, Au, Hg, Al, Si, In, Tl, Pb, Bi, Sb, Te, U, Sm, Tb, La, Ce or Nd.
17. The catalyst layer of claim 16, wherein said catalytically active element comprises one or more of Pt, Pd, Au, Ag, Cu, Ni, Fe, Sn, Bi, Co, In, Ru or Rh.
18. The catalyst layer of claim 17, wherein said catalytically active element comprises one or more of Au, Ag, Cu, Sn, Sb, Bi or In.
19. The catalyst layer of claim 18, wherein the weight of the ion conducting polymer in the catalyst layer is less than 64% of the weight of the catalytically active element in the catalyst layer.
20. The catalyst layer of claim 19, wherein the weight of the ion conducting polymer is at least 1% and less than or equal to 10% of the weight of the catalytically active element in the catalyst layer.

21. The catalyst layer of claim 13, wherein said catalyst layer also comprises elemental carbon.
22. The catalyst layer of claim 21, wherein the carbon is a carbon black
23. The catalyst layer of claim 13, wherein the electrochemical device is an electrolyzer.
24. The electrolyzer of claim 23, wherein operation of the electrolyzer produces electrolysis products comprising one or more of CO , OH^- , HCO^- , H_2CO , $(\text{HCO}_2)^-$, HCOOH , H_2O_2 , CH_3OH , CH_4 , C_2H_4 , $\text{CH}_3\text{CH}_2\text{OH}$, CH_3COO^- , CH_3COOH , C_2H_6 , O_2 , H_2 , $(\text{COOH})_2$ or $(\text{COO}^-)_2$.
25. An electrolyzer comprising the catalyst layer of claim 13, the electrolyzer having a reactant stream input, wherein the reactant stream to the electrolyzer comprises at least one of CO_2 or H_2O .
26. The electrolyzer of claim 25, the electrolyzer having an anode and a cathode, wherein said catalyst layer is in electrical contact with at least one of the anode or the cathode.
27. The electrolyzer of claim 26, wherein the CO_2 is fed into the cathode.
28. The electrolyzer of claim 26, wherein said catalyst layer is in electrical contact with the cathode.
29. The electrolyzer of claim 26, wherein water is fed into the anode.
30. The electrolyzer of claim 29, wherein the water feed also includes an electrolyte.
31. The electrolyzer of claim 30, wherein the electrolyte is at least one of a carbonate, a bicarbonate or a hydroxide.
32. An electrochemical device comprising the catalyst layer of claim 13, wherein the electrochemical device is a sensor.
33. An electrochemical device comprising the catalyst layer of claim 13, wherein the device is a fuel cell.

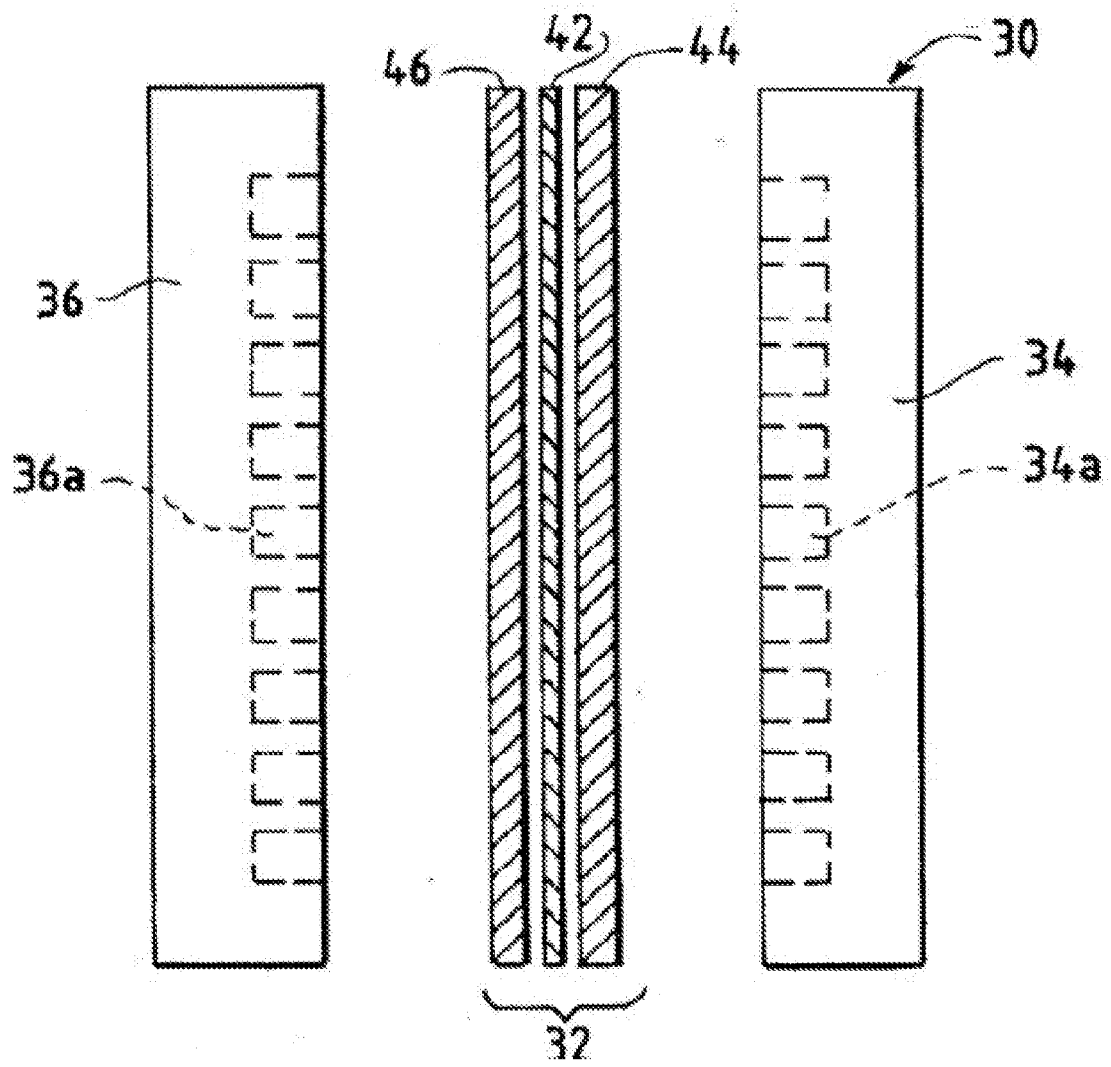
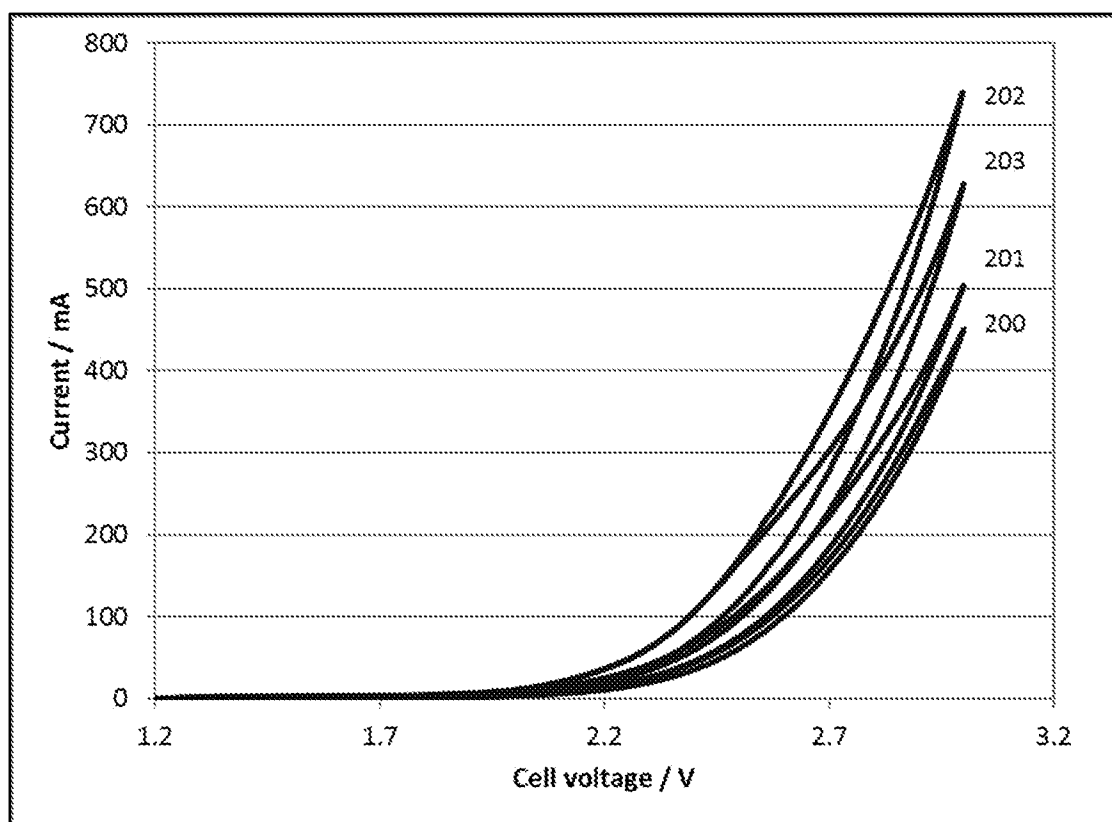
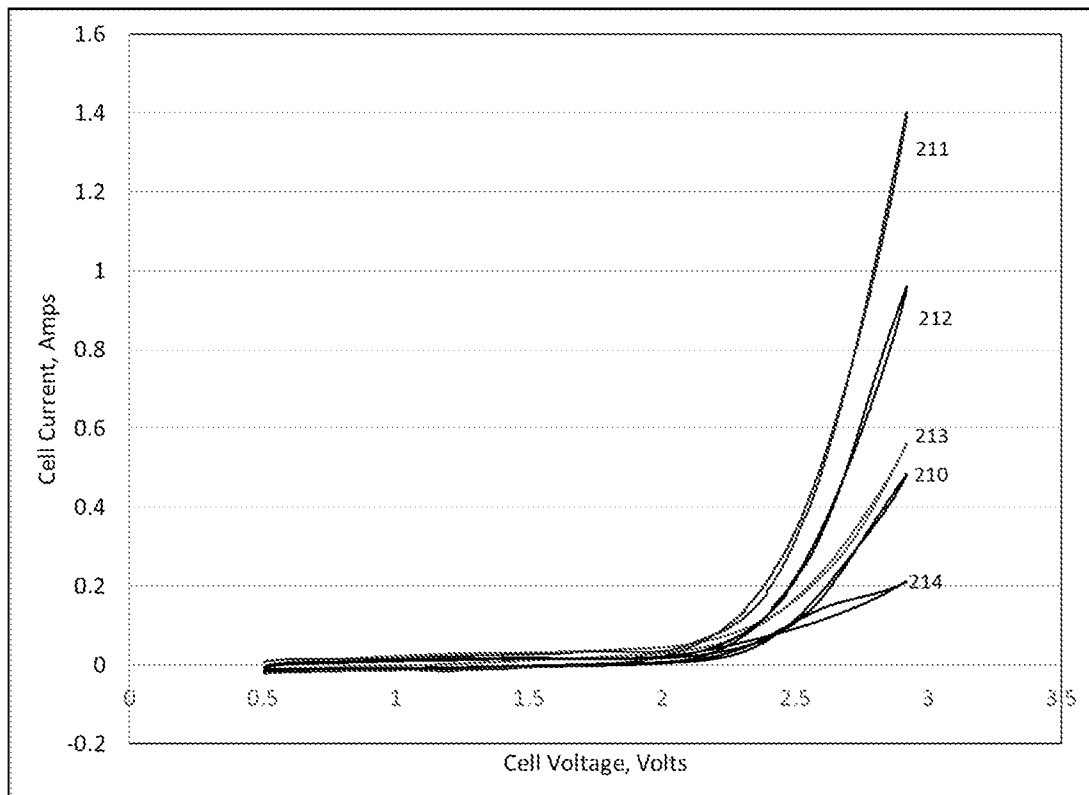
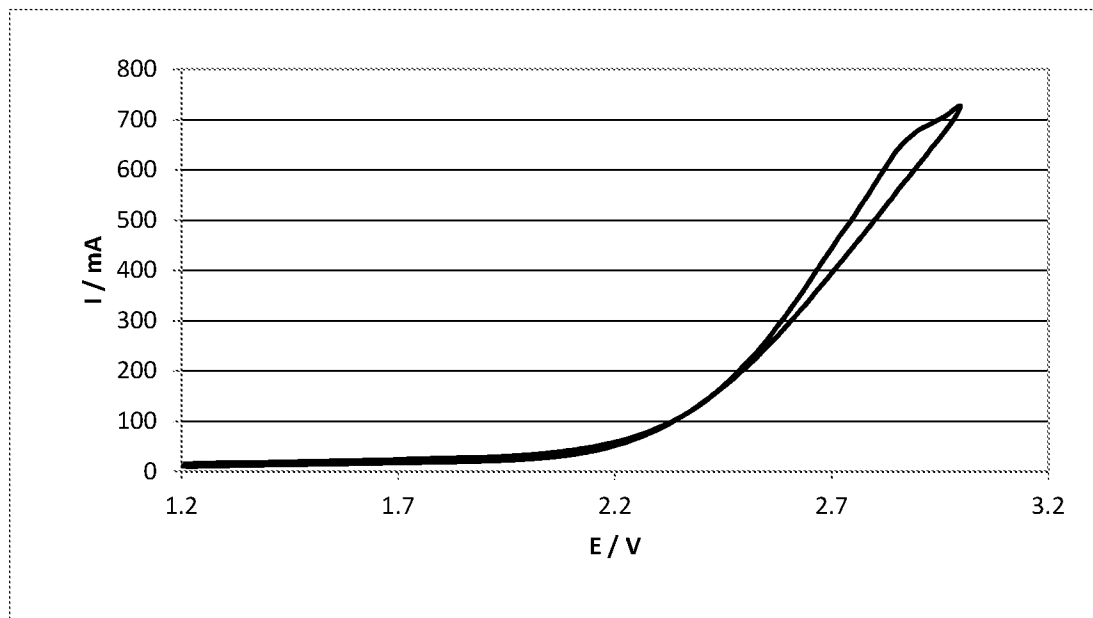


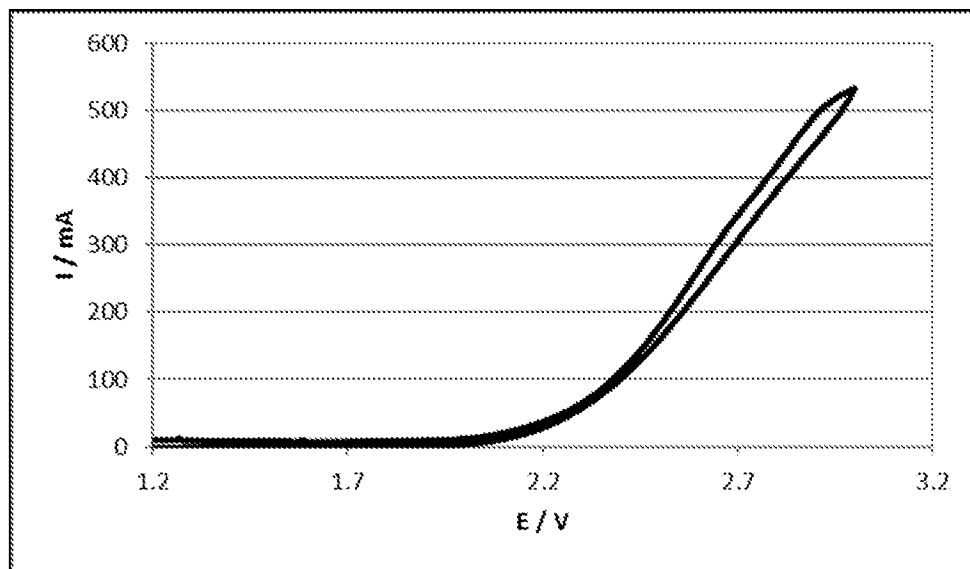
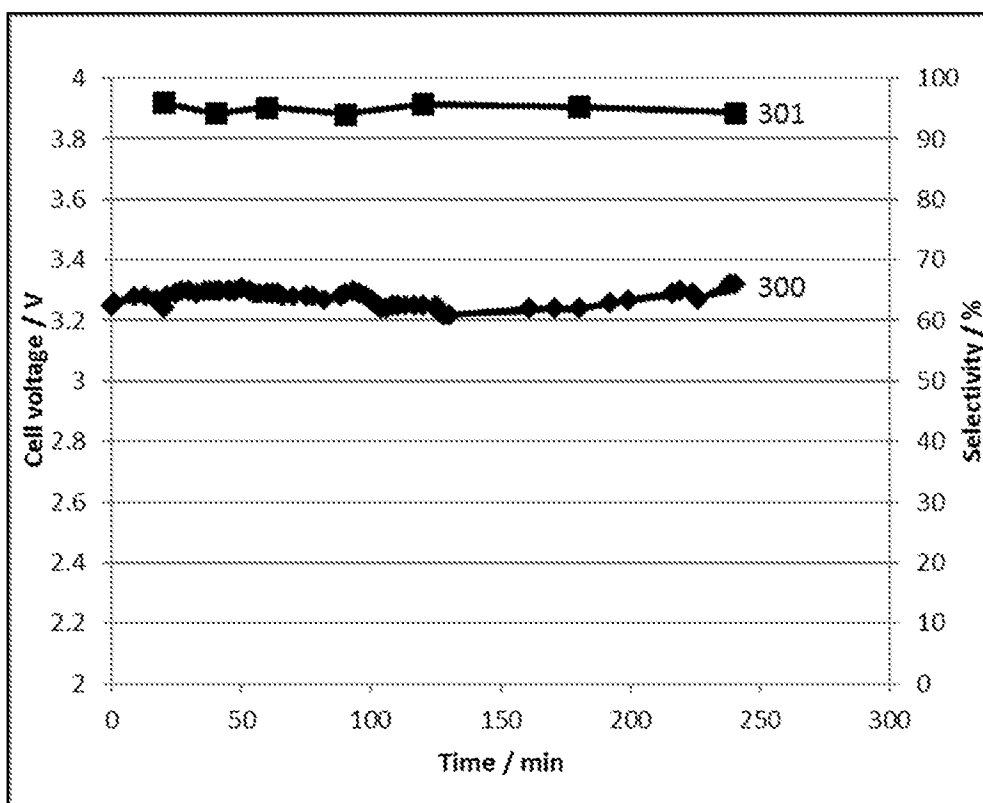
FIG. 1

**FIG. 2**

3/4

**FIG. 3****FIG. 4**

4/4

**FIG. 5****FIG. 6**

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2017/025624

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-12

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2017/025624

A. CLASSIFICATION OF SUBJECT MATTER		
INV.	C25B3/04 G01N27/30	C25B11/04 H01M8/00 H01M4/90 H01M4/86
ADD.		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols) C25B		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	XIAOMING YAN ET AL: "Imidazolium-functionalized poly(ether ether ketone) as membrane and electrode ionomer for low-temperature alkaline membrane direct methanol fuel cell", JOURNAL OF POWER SOURCES, vol. 250, 8 November 2013 (2013-11-08), pages 90-97, XP055319613, CH ISSN: 0378-7753, DOI: 10.1016/j.jpowsour.2013.10.140 cited in the application abstract, figure 1, sections 2.11 and 3.7 -----	1-12
X	US 2012/171583 A1 (BOCARSLY ANDREW B [US] ET AL) 5 July 2012 (2012-07-05) cited in the application paragraphs [0016], [0023]; claims 11-18 ----- -/-	1-3,5,7, 8,12
☒ Further documents are listed in the continuation of Box C. ☒ See patent family annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search 13 June 2017		Date of mailing of the international search report 16/08/2017
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Ritter, Thomas

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2017/025624

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2016/039999 A1 (3M INNOVATIVE PROPERTIES CO [US]) 17 March 2016 (2016-03-17) figure 1; example 1 -----	1-10,12
X	US 2015/345034 A1 (SUNDARA RAMAPRABHU [IN] ET AL) 3 December 2015 (2015-12-03) paragraphs [0004], [0019], [0025], [0035] - [0038]; examples 1, 3 -----	1-3,5-12
A	US 2011/237830 A1 (MASEL RICHARD ISAAC [US]) 29 September 2011 (2011-09-29) the whole document -----	1-12

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2017/025624

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2012171583 A1	05-07-2012	US 2012171583 A1	05-07-2012
		WO 2013102086 A1	04-07-2013

WO 2016039999 A1	17-03-2016	AU 2015315620 A1	23-03-2017
		CA 2960595 A1	17-03-2016
		CN 106715760 A	24-05-2017
		EP 3191621 A1	19-07-2017
		KR 20170056588 A	23-05-2017
		US 2017183789 A1	29-06-2017
		WO 2016039999 A1	17-03-2016

US 2015345034 A1	03-12-2015	NONE	

US 2011237830 A1	29-09-2011	AU 2011230545 A1	11-10-2012
		CA 2794105 A1	29-09-2011
		CN 102892929 A	23-01-2013
		EP 2553147 A1	06-02-2013
		JP 2013525088 A	20-06-2013
		JP 2016041427 A	31-03-2016
		KR 20130060185 A	07-06-2013
		US 2011237830 A1	29-09-2011
		US 2015122640 A1	07-05-2015
		WO 2011120021 A1	29-09-2011

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1-12

An electrolyzer comprising a catalyst layer, the catalyst layer comprising a catalytically active element and an ion conducting polymer, wherein said ion conducting polymer comprises positively charged cyclic amine groups, and further wherein a reactant stream comprising carbon dioxide is in contact with said catalyst layer.

2. claims: 13-33

A catalyst layer for an electrochemical device comprising a catalytically active element and an ion conducting polymer wherein said ion conducting polymer comprises: (a) positively charged cyclic amine groups; and (b) polymers or copolymers of styrene.
