Advanced Fuel Cell Membranes Based on Heteropolyacids

John A. Turner and F. J. John Pern

Hydrogen and Electricity, Systems and Infrastructure Group National Renewable Energy Laboratory Golden, CO 80401-3393

Andrew M. Herring* and Steven F. Dec**

* Department of Chemical Engineering and ** Department of Chemistry and Geochemistry Colorado School of Mines Golden, CO 80401-1887

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Overview

Timeline

- Project start date: FY 2005
- Project end date: tbd
- Percent complete: tbd

Budget

- Total project funding
 - DOE share: \$300K
- Funding received in FY05:
 - \$150K (0.3 FTE)
- Funding for FY06:
 - \$150K (0.3 FTE)

Targets

- Low humidity operation (25% RH).
- High conductivity ~0.1 S/cm
- Cost \$40/m²

Barriers

- Barriers addressed
 - A. Durability.
 - B. Cost.
 - D. Thermal, Air and Water Management.

Partner/Subcontract

- Colorado School of Mines
 - Prof. Andrew M. Herring
 - Dr. Steven F. Dec



Objectives

 Develop the methodology for the fabrication of 3D crosslinked, hydrocarbon-based membranes using immobilized heteropolyacids (HPAs) as the proton conducting moiety.

- Conductivity ~0.1 S/cm at 120°C and <1.5 kPa H_2O

- Develop immobilization technology based on covalent attachment of HPAs to oxide nano-particles.
- Acquire an improved understanding of HPAs and their salts made by custom synthesis.
 - HPAs make up a class of inorganic proton conductors that exhibit high proton conductivity at low humidity (below 25% RH) and at elevated temperatures (well above 100°C).
- Conduct relevant characterizations of the membranes to better understand their structural, chemical, and thermal properties/stability and proton conductivity.

HPAs: High H⁺ Conductivity, High Thermal Stability; Vast Structural Diversity; Known Redox Catalysts



Lacunary (allows easy attachment points) $[SiW_{11}O_{39}]^{8-}$ $H_8SiW_{11}O_{39}-26H_2O_{5}$ (W11-STA)

Strategies for Immobilizing HPAs

A. Binding Approaches:

- Covalent bonding to oxide nano-particles insitu, which can bond covalently 1. to, or embed physically in, a polymeric matrix
- 2. 3.
- Direct embedding in a polymeric matrix Covalent bonding directly to a polymeric matrix (CSM/3M collaboration, poster #FCP-6)

B. Modification of Lacunary HPAs:

By bonding with functional silanes that can then be cross-linked or polymerized

C. Fabrication Approaches:

- Sol gel method 1.
- 2 Immobilized via silulation onto supporting particles
- Simple blending 3.

D. Polymeric Matrix:

- Organic 1.
- 2. Inorganic
- 3. Organic-inorganic hybrid



Ref. 14: "Heteropoly and Isopoly Oxometalates," by M. T. Pope, Springer-Verlag, New York, 1983, Chap. 7, Fig. 7.8, p. 126.



Key Concept and Components in Composite Membrane Fabrication

3-D Cross-linked Composite Matrix



Procedure for Fabricating 3D Cross-Linked HPA/SiO₂/Functional Silane Sol Gel Composite & PEM Membrane with PMG







Formation of SiO₂ Nano-Particles in Composite Matrix upon Thermal Treatments (TEM Analysis)



Flexibility of PEM Membranes Fabricated with High HPA Loading



PEM-#9B,C,D Films: W12-STA/(PMG + Cross-Linker) = 174 Wt%



Immobilizing the HPA Binding HPA with Z-6030 Silane in Sol Gel Composite → W12-STA Retained



Chemical Stability of Membrane and Composite PEM



PEM Mechanical Strength and Flexibility Reduced by Increasing HPA Loading

FTIR-ATR Spectra of Cured Control Blanks and PEM-#7



H⁺ Conductivity as a Function of Cell Temperature at 100% RH



Improving H⁺ Conductivity with Higher HPA Loading and Better Membrane Fabrication

Table 1. PEM Compositions vs Proton Condutivity Derived from I-V Curves of CV Scans

		Components		Weight Ratio	Best Proton Conductivity (mS/cm)		
PEM ID	HPA	Host Polymer	X-Linker	HPA/(PMG + X-Linker)	80°C/100%RH	100°C/46%RH	120°C/23%RH ¹
1	HSiW12Ox	BSPPO	No	0.56	0.15		
2	HSiW12Ox	PMG	Yes	0.81	6.9		
3	HSiW11Ox	PMG	Yes	1.09	6.4, 10.46	2.41	0.85
4	KSiW10Ox	PMG	Yes	1.05	7.56, 13.3	1.61	0.25
5	HSiW12Ox	PMG	Yes	1.50	8.8		
6	HSiW12Ox	PMG	Yes	1.54	15.57		
7	HSiW12Ox	PMG	Yes	1.74	14.55	2.1	
8	HSiW12Ox	PMG	Yes	1.74	19.17	3.81	
9B 9C	HSiW12Ox HSiW12Ox	PMG PMG	Yes Yes	1.74 1 74	22.28 21 15		
9D	HSiW12Ox	PMG	Yes	1.74	25.45	[28.25 at 70°C/	100%RH]
Nafion 112	SO3H				149.9	98.99	49.25

¹ Values of the proton conductivity at 120°C/23%RH are with large uncertainty because of rapidly lost linearity on I-V curves

High H⁺ Diffusion Coefficients for Composite Membrane





Summary of Accomplishments

PEM Fabrication and Performance

 We have shown the ability to retain HPAs into a polymercomposite matrix of our design.

Properties of HPA-based composite PEMs:

- high chemical stability (Fenton's reagent test)
- good thermal stability (with highly reactive W12-STA)
- good mechanical flexibility
- effective binding of silicotungstic acids (Wn-STA) with select functional silanes (n = 10, 11, 12)
- high Wn-STA loading [HPA/(PMG + X-Linker) > 150 wt%]
- moderate proton conductivity (25 mS/cm at 80°C/100%RH)

Clear progress towards meeting the DOE targets





Achieving Fundamental Goals Future Work

- To continue to improve/modify/optimize the current PEM composite formulation, fabrication, and processing conditions
 - to enhance PEM's thermal stability in the 90-120°C range
 - to improve mechanical strength and flexibility
 - to reduce membrane thickness and improve film uniformity
- To continue to develop immobilization strategies for various HPAs, custom-synthesized at CSM, that show high proton diffusion coefficients and thermal stability.
- To understand the binding mechanism of HPA with functional silanes and SiO₂ nano-particles in the polymer matrix.
- To understand the proton conduction mechanism in the 3D cross-linked composite membranes in order to further improve proton conductivity at low humidity and elevated temperatures.

2005 Reviewers' Comments

- "One of the few new, alternative ideas for membranes in the whole DOE program"
- Issues:
 - ...needs to present conductivity values for membranes with "fixed" HPAs...
 - Done
 - HPA approach is sound as a demonstration but water solubility must be addressed...
 - Excellent progress has been made in this regard
 - Nafion doped in HPAs has been shown to be feasible...the PI is in need of new insight.
 - Not part of our project, those figures were for introduction to HPAs only
 - Our project is focused on developing a composite hydrocarbon membrane using HPAs as the proton conducting moiety that will meet the DOE targets for operation at low RH and higher temperatures
- Future:
 - Need durability studies in actual operating fuel cell conditions and ...thermal and RH cycling...gas crossover measurements
 - PEMs of 3D cross-linked PMG matrix were not available yet at the time
 - These subjects will be investigated for HPA-based PEMs this summer



Presentations and Publications

- F. J. Pern, J. A. Turner, Fanqin Meng, and A. M. Herring; "Sol-Gel SiO₂-Polymer Hybrid Heteropoly Acid-Based Proton Exchange Membranes," MRS 2005 Fall Meeting, Energy and The Environment Symposium, Session A: The Hydrogen Cycle—Generation, Storage, and Fuel Cells. In press.
- F. J. Pern, J. A. Turner, and A. M. Herring; "Hybrid Proton Exchange Membranes Based on Heteropoly-Acid and Sulfonic-Acid Proton Conductors," ECS 2006, Abstract (accepted for oral presentation)
- J. L. Horan , J. Turner , A. M. Herring, and S. Dec; "Structure and Dynamics of Non-Commercial Heteropoly Acids for Fuel Cell Applications," ECS 2006, Abstract
- N. V. Aieta, M. Kuo, F. Meng, J. Turner, and A. M. Herring; "The Use of Heteropolyacids as Additives for Low Humidity Operation of Nafion Membranes for PEM Fuel Cell Applications," ECS 2006, Abstract
- A. M. Herring, R. J. Stanis, J. Ferrell III, M. Kuo, J. Turner, and M. Samaroo; "The Use of Heteropoly Acids as Electrocatalysts for the Oxygen Reduction Reaction in PEM Fuel Cells," ECS 2006, Abstract
- R. J. Stanis, A. M. Herring, M. Kuo, and J. Turner; "Increased CO Tolerance of Pt Electrodes by Addition of Adsorbed Heteropoly Acids and Salts in PEM Fuel Cell Anode Catalysts," ECS 2006, Abstract

