2018 U.S. DOE HYDROGEN and FUEL CELLS PROGRAM and VEHICLE TECHNOLOGIES OFFICE ANNUAL MERIT REVIEW and PEER EVALUATION MEETING, June 13-15, 2018

Project ID: FC175

Polymer-based fuel cells that operate from 80–220 °C

PI: Yu Seung Kim

Los Alamos National Laboratory

June 15, 2018

This presentation does not contain any proprietary, confidential, or other wise restricted information



Overview

Timeline

- Project start date: 5/1/2017
- Project end date: 10/30/2018
- Percent complete: 70%

Budget

- Total project funding: \$300K
 DOE share: 100%
- Funding received in \$270K
 FY18:
- Total DOE Funds Spent*: \$200K

*As of 4/17/2018

Barriers

- B. Cost
- C. Electrode performance
- A. Durability

Project lead

- Los Alamos National Laboratory
 - Yu Seung Kim (PI), EunJoo Park, Albert Lee, Dongguo Li, Gerie Purdy

Collaborators (No cost)

Sandia National Laboratory
 - Cy Fujimoto



Universität Stuttgart

- Rensselaer Polytechnic Institute
 Chul Sung Bae, Junyoung Han
 Rensselaer
- National Institute of Advanced Industrial Science and Technology
 - Yoong-Kee Choe
- University of Stuttgart
 - Vladimir Atanasov
- Nanosonic, Inc.
 William Harrison
- Toyota Motor North America, Inc.
 Hongfei Jia
 TOYOTA



Relevance

Objective

 Development of a feasible ion-pair coordinated polymers to demonstrate a fuel cell that is operational from 80–220°C without humidification.

Cost Reduction of a Fuel Cell Car

- Current fuel cell car retail price: \$60,000.
- Projected fuel cell system cost: \$45/kW for 500,000 vehicles
- Target cost: \$40/kW (2020)
 \$30/kW (ultimate)
- Further cost reduction strategy is necessary







High temperature and low RH fuel cell operation could enable fixed cost savings of \$7.5/kW_{net} by eliminating or reducing the size of BOP components such as humidifier and radiator.

N. Dale, Nissan Motors

Approach: Ion-Pair Coordinated Fuel Cells







• Los Alamos

* Nature Energy, 1, 16120 (2016)

Approach: Project Phases and Milestones

DFT Calcul	lation	√	Determine the interaction energy between quaternary ammonium and phosphate	100 %	
Small Molecule Study		е	 Determine the best quaternary ammonium and phosphate phase 	100 %	
	MEA verificati	on	✓ Demonstrate the membrane <i>in-situ</i> and <i>ex</i> - performance (conductivity, water tolerance)	- <i>situ</i> e)	60 % See below

Milestone (4/30/18)	Units	Baseline* (2016)	Target	Current Status (as of April 15 2018)
ASR (160 °C)	Ω cm ²	0.24	< 0.10	0.09
Peak Power Density (H ₂ /O ₂)	mW cm ⁻²	800	1000	1134
Water Tolerance (Go-No-Go, April 30, 2018)	kPa	21.3	38.5	42.6
Durability during 80-180 °C AST	V loss at 160 °C	30%	<10%	Not evaluated yet



* Nature Energy, 1, 16120 (2016)

Accomplishment: DFT Calculation



- Ion pair coordinated system has higher interaction energy with PA than acid-base coordinated system at a given number of PA.
- Better water retention can be achieved with stronger interaction energy.





Accomplishment: Interaction Measurement (³¹P NMR)



- ³¹P NMR shows a single peak, indicating exchanging electrons in the phosphate phase.
- Ion pair coordinated system has higher interaction energy consistent with DFT study.
- Interaction energy HMG > TBA > TMA >> guanidine > benzimidazole > pyridine



Accomplishment: Proton Conductivity



- Ion pair coordinated system has stronger interaction at a given conductivity.
- Hexamethyl guanidinium (HMG⁺) exhibits the strongest interaction at a given conductivity.



Accomplishment: Water Tolerance



Highlight: Met the water tolerance go-no-go decision criteria (max. conductivity at $P_{H2O} > 38.5$ kPa) with two newly designed ion-pair coordinated systems.





Membrane Synthesis

Membrane: Ammonium-phosphate ion-pair coordinated cross-linked poly(biphenylene) (PA-XL-BPN, Technical Back-up Slide #2)*



High IEC ion-pair coordinated crosslinked poly(biphenylene) is developed.
Highlight: Fuel cell peak power density (> 1 W cm⁻²) milestones.



 H_2/O_2 , 285 kPa abs backpressure; membrane: PA-XL-BPN; ionomer: PA-QAPS, Pt 0.6 mg/cm² for both electrodes



Accomplishment: Ionomer development





Ionomer: Phosphonated poly(pentafluorostyrene) (PPFS, Technical Slide 4)*



Developed phosphonated ionomer (non-acid leachable) for ion-par system.

Highlight: Obtained ASR of 0.09 Ω cm² at 160°C Exceed the ASR (< 0.1 Ω cm²) milestone



Measured in H_2/O_2 , 147 kPa abs backpressure; membrane: PA-DAPP; Pt-Ru 0.5 mg_{Pt}/cm² for anode and Pt 0.6 mg/cm² for cathode







Responses to Previous Year Reviewers' Comments

This project was not reviewed last year.



Project Coordination





Collaboration

• Material Exchange: Critical component to achieve the project objective

Sandia National Laboratory (Cy Fujimoto) - Federal Lab., within DOE program

- Hexamethyl ammonium functionalized poly(phenylene)s 6 pieces (5" × 5")
- Benzyl ammonium functionalized poly(phenylene)s 3 pieces (5" × 5")

Rensselaer Polytechnic Institute (Chulsung Bae) – University, within DOE program

- Crosslinked biphenylene membranes: 6 pieces (4 " × 4 ")
- Crosslinked terphenylene membranes: 6 pieces (4" × 4 ")

University of Stuttgart (Vladimir Atanassov) - University (foreign), outside DOE program

- Phosphonated polystyrenes with different IECs: 10 pieces (2" × 2")

Nanosonic Inc. (William Harrison) - Industry, inside DOE program (SBIR)

- Quaterary ammonium functionalized poly(arylene)s: 9 pieces (4" × 4")

Toyota Motor North America, Inc. (Hongfei Jia) – Industry, outside DOE program - Ion liquid electrolyte: few grams

• **DFT Modeling:** Useful information provided to design materials

National Institute of Advanced Industrial Science and Technology (Yoong-Kee Choe, LANL visit) – National Lab (Foreign), outside DOE program

- Ion-pair interaction calculation



Remaining Challenges and Barriers

Within the project (ending October 30/2018)

No substantial technical challenges remained.

After the project

- Electrode development
 - Ionomer design
 - Understanding catalyst-ionomer interfacial reaction
 - Low Pt loading and non-PGM catalysts
- o Membrane development
 - Increasing molecular weight & IEC
 - Mechanically stable thin film construction
 - Incorporating super-base cations into polymeric materials
 - Understanding proton conduction in the ion pair networks
- Tech validation
 - Correlation of fuel cell AST and field test performance
 - Fuel cell start-up stability
 - Techno-economic analysis



Current and Proposed Future Work - Technical

With this project

Membrane development

• Complete synthesis of guanidinium poly(phenylene)s (Technical Backup Slide #5).

Ionomer development

Synthesis of phosphonated ionomers.

Fuel cell testing

- Initial performance with Pt-based catalysts under H_2/O_2 and H_2/air conditions.
- In-situ water tolerance test (humidity-temperature cycling 80 160 °C).
- Currently no alternative development pathways is planned; however we may add some additional work based on AMR comments.

Beyond this project

Investigation of interface between catalyst and ion-paired ionomer

- Hydrogen oxidation reaction
- Oxygen reduction reaction
- Funding (LANL-Toyota CRADA)



Current and Proposed Future Work - Technology Transfer Activities

1.2

SBIR Phase II

 Technology transfer effort to Nanosonic Inc. (SBIR Phase II) to produce scale-up synthesis of ionpair coordinated membranes.

Patent application

- Kwan-Soo Lee and Yu Seung Kim, "Proton exchange membrane fuel cells" US Patent Application No. 62/377,163.
- Sarah Park, Sandip Maurya, Yu Seung Kim, "Polymer electrolytes for alkaline membrane fuel cells" S133606 (March, 2, 2018)
- Albert Lee and Yu Seung Kim, "Polymer electrolytes for fuel cells" (Feb. 9, 2018).

80 C 1.0 160 C 200 C 0.8 Voltage / V 0.6 0.4 0.2 H_2/O_2 0.0 0.0 0.2 0.4 0.6 0.8 1.0 1.2 1.4 i / A cm⁻² 0.40 0.35







William Harrison

The performance of Nanosonic membrane tested by LANL

Summary

- **Objective:** Development of feasible ion-pair coordinated polymers to demonstrate a fuel cell that operates at 80–220°C without humidification.
- **Relevance:** Aiming to make ion-pair coordinated membranes having comparable proton conductivity and durability without humidification. Successful development of such ion-pair coordinated membranes enable to operate fuel cells at the wide-range of temperature without humidification.
- **Approach:** DFT modeling and small molecule study (³¹P NMR) determine the best candidate materials that have strong ionic interactions. The selected ion pairs are incorporated in membranes and demonstrate good water tolerance and low cell resistance in membrane electrode assemblies.
- Accomplish-
ments (FY 18)Completed the DFT modeling and small molecule study, demonstrating
guanidinium-phosphate is the best candidate having strongest
interaction. Achieved water tolerance at 80 °C, 80% RH (P_{H20} = 38.5
kPa), met the go-no go decision criteria. Demonstrated > 1 W/cm² peak
power density of ion-pair coordinated HT-PEMFC.
- **Collaborations:** Collaboration in the area of chemical synthesis, MEA integration, electrochemistry and fuel cell testing. Extensive communications with several industrial partners, including RPI, University of Stuttgart (Germany), Sandia National Laboratories, AIST (Japan) Nanosonic, Inc and Toyota Motors North America.



Technical Back-Up Slides



Determination of Phosphoric Acid Doping Level



Typical doping process

- → Completely submerge membrane in 85 wt% phosphoric acid bath under ambient conditions.
- → Hang dry membrane until excess phosphoric acid drips off.
- \rightarrow Dry at 80°C under air for 2 hr.
- \rightarrow 12 h was found to be appropriate to achieve non-excess doping levels.



Technical Back-up #1

Synthesis and Properties of Cross-linked BPN





Tensselaer J.Y. Han and C. Bae

Technical Back-up #2

Synthesis and Properties of PA-QAPS*





* Nature Energy, 1, 16120 (2016)

Technical Back-up #3

(%)

35

Per PA

5.2

Synthesis of Phosphonated Poly(pentafluoro styrene)*





TMPS/PPFS (mol %)	Phosphonation ¹⁹ F NMR (%)	Phosphonation IEC (%)	IEC-direct/total (mequiv. g⁻¹)
200	100	99	7.4/8.0
90	66	51	1.9/4.1
80	42	41	1.4/3.5
70	41	44	1.6/3.8
60	36	32	1.1/2.9
40	17	17	0.5/1.6



J. Power Sources, 343, 363-372 (2017)

Technical Back-up #4





Synthesis of Phenyl Guanidinium Funct. Poly(pheneylene)s



Poly(phenylene)-based anion exchange polymers and methods thereof" USP applications US IEC = 1.9 – 2.5 meq/g 2017/0190831 A1, K.S. Lee, Y.S. Kim, C. Fujimoto (2017)





C. Fujimoto

Technical Back-up #5