

High Performance non-PGM Transition Metal Oxide ORR Catalysts of PEMFCs



P.I.: Timothy C. Davenport



ElectroCat Consortia Project

Project ID: FC306

DE-EE0008420

2020 Hydrogen and Fuel Cells Annual Merit Review

This presentation does not contain any proprietary, confidential, or otherwise restricted information.

This presentation does not contain technical data subject to the EAR or the ITAR.

Overview

Timeline

Project Start: March 2019

Project BP1: March 2019 – August 2020

Project End: August 2021*

* Project continuation and direction determined annually by DOE

Budget

Total Project Budget: \$1,250K

- Federal Share \$1,000K

- Cost Share (20%) \$250K

Total DOE Funds Spent*: **\$473K**

* as of 5/27/2020

Barriers Addressed

- Low performance for PGM-free oxygen reduction reaction catalysts for PEMFCs

Target I.D. #	Characteristic	Units	2020 Targets
FC-4	Loss in initial catalytic activity	% mass loss	< 40
FC-5	Loss in performance at 0.8 A cm ⁻²	mV	< 30
FC-8	PGM-free catalyst activity	A cm ⁻² at 900 mV _{iR-free}	> 0.044

Funded Partners

Massachusetts Institute of Technology

Relevance

Objective: Develop *acid-stable non-PGM metal oxides* and *optimize oxide catalytic activity for ORR reactivity*.

	Barrier	Approach	2020 Impact
A	Durability	Focusing on improving the acid stability of novel materials will lead to higher durability materials for PEMFCs	<ul style="list-style-type: none">• Acid-stability descriptors developed based on manganese oxides• Family of acid-stable antimony-based oxides developed
B	Cost	Development of non-PGM electrocatalysts is a key approach to reducing PEMFC cost	<ul style="list-style-type: none">• Project focuses on Mn-based oxide materials that do not contain PGMs
C	Performance	Optimize the catalytic activity by the doping of acid-stable oxides with multiple metal cations to fill multiple roles including stability, catalytic activity, and electronic conductivity	<ul style="list-style-type: none">• Evaluated electrocatalytic activity of Mn-based oxides

Approach

Develop acid-stable non-PGM metal oxides

- Identify descriptors for acid-stability to guide prediction of acid stable oxides
- DFT calculation of phase stability of doped metal oxides of interest as potential acid stable oxides

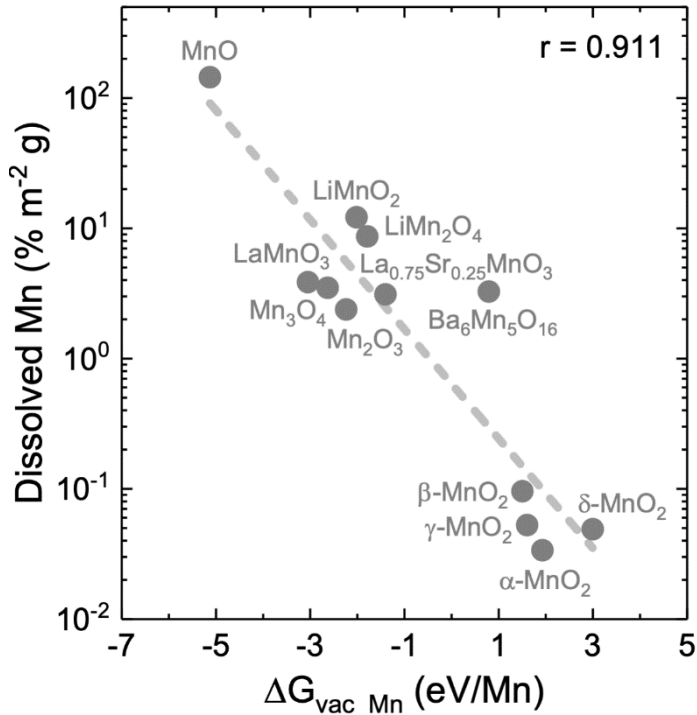
Optimize oxide catalytic activity for ORR

- Electrochemical characterization to determine intrinsic activity of novel materials
- Optimize material synthesis for high surface area
- Optimize catalyst layer composition with high-throughput ink formulations

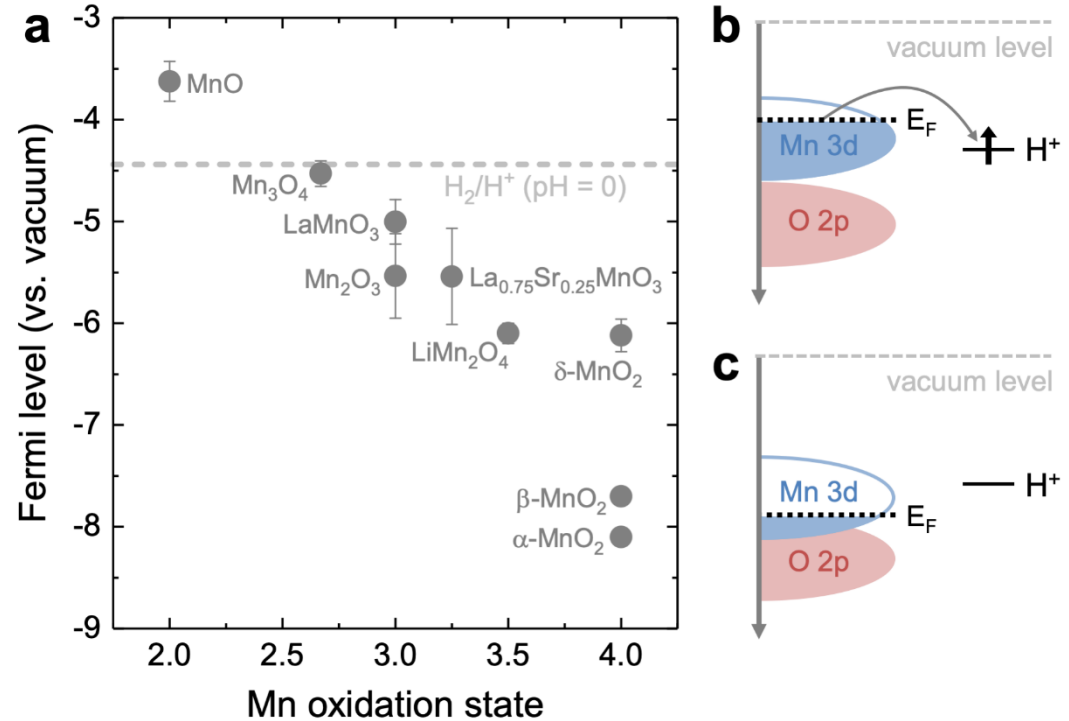
Project Milestones	Original Planned	Revised Planned	Percent Complete	Notes
Subcontracts completed	5/31/2019		100%	Completed
Evaluation of Acid Stability of A_xMnO_2	5/31/2019		100%	No doped MnO_x have been found to be acid stable under ORR conditions
Evaluation of Intrinsic ORR Activity of Acid-Stable A_xMnO_2	8/31/2019		100%	Evaluated activity of most acid-stable MnO_x systems – proceeding to 2 nd generation oxides
Demonstrate intrinsic ORR activity $\geq 0.5 \mu A\text{-cm}^{-2}_{\text{oxide}}$ at 0.9 V (iR-free) with acid stable oxide	8/31/2019	8/31/20	50%	Activity observed, but not with demonstrated acid stability
Optimize Catalyst Layer Composition with best catalyst developed	11/30/2019	8/31/20	0%	
Demonstrate intrinsic ORR activity $\geq 4.4 \mu A\text{-cm}^{-2}_{\text{oxide}}$ at 0.9 V (iR-free) with an acid stable oxide	11/30/2019	8/31/20	0	
Demonstrate MEA with performance of $0.025 A\text{-cm}^{-2}$ at 0.9 V (iR-free) under 1 atm O_2 and 80 °C	2/29/2020	8/31/20	0%	Go/No-Go

Progress: Acid Stability Descriptors

Mn vacancies are energetically preferred over O vacancies



Less acidic surface favors the attack of proton, where protonation weakens surface Mn-O bonds



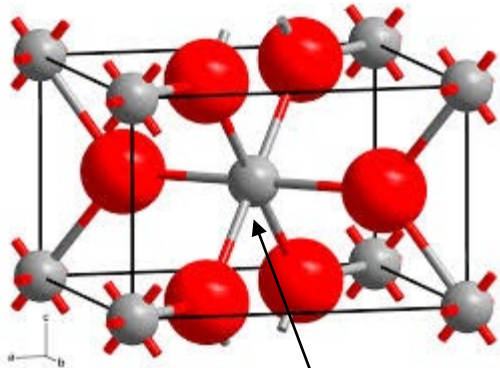
Acid stability descriptor developed for manganese oxides, indicating high stability of high oxidation state manganese oxides.

Identified rutile phase as a target for Mn-based acid stable oxides ($\beta\text{-MnO}_2$).

Progress: Acid Stable Rutile Structures

Overview:

Sb_2O_4 is a mixed valent (Sb^{III} , Sb^{V}) oxide
 Sb^{III} and Sb^{V} are randomly disordered on the single crystallographic metal ion site (4+ site charge)



$\frac{1}{2} Sb^{3+}$, $\frac{1}{2} Sb^{5+}$

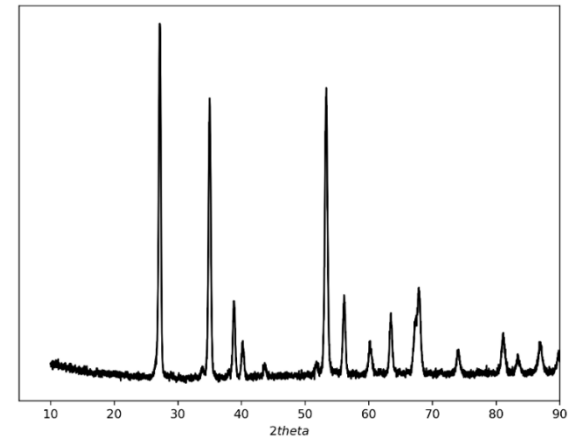
1 H hydrogen 1.00784, 1.00819	2 He helium 4.0026											13 B boron 10.811	14 C carbon 12.011	15 N nitrogen 14.0064, 14.0031	16 O oxygen 15.999, 16.003	17 F fluorine 18.998	18 Ne neon 20.180	
3 Li lithium 6.941	4 Be beryllium 9.0122											13 Al aluminum 26.982	14 Si silicon 28.086, 28.085	15 P phosphorus 30.974	16 S sulfur 32.06, 32.059	17 Cl chlorine 35.45, 35.453	18 Ar argon 39.95	
11 Na sodium 22.990	12 Mg magnesium 24.304, 24.305	3	4	5	6	7	8	9	10	11	12	13 Ga gallium 69.723	14 Ge germanium 72.630(8)	15 As arsenic 74.922	16 Se selenium 78.9718(8)	17 Br bromine 79.901, 79.907	18 Kr krypton 83.798(2)	
19 K potassium 39.098	20 Ca calcium 40.078(4)	21 Sc scandium 44.956	22 Ti titanium 47.867	23 V vanadium 50.942	24 Cr chromium 51.996	25 Mn manganese 54.938	26 Fe iron 55.845(2)	27 Co cobalt 58.933	28 Ni nickel 58.693	29 Cu copper 63.546(3)	30 Zn zinc 65.38(2)	47 Ag silver 107.87	48 Cd cadmium 112.41	49 In indium 114.82	50 Sn tin 118.71	51 Sb antimony 121.76	52 Te tellurium 127.603(8)	53 I iodine 126.90
37 Rb rubidium 85.468	38 Sr strontium 87.62	39 Y yttrium 88.906	40 Zr zirconium 91.224(2)	41 Nb niobium 92.906	42 Mo molybdenum 95.95	43 Tc technetium	44 Ru ruthenium 91.07(2)	45 Rh rhodium 102.91	46 Pd palladium 106.42	47 Ag silver 107.87	48 Cd cadmium 112.41	49 In indium 114.82	50 Sn tin 118.71	51 Sb antimony 121.76	52 Te tellurium 127.603(8)	53 I iodine 126.90	54 Xe xenon 131.29	
55 Cs cesium 132.91	56 Ba barium 137.33	57-103 lanthanoids	72 Hf hafnium 178.49(2)	73 Ta tantalum 180.95	74 W tungsten 183.84	75 Re rhenium 186.21	76 Os osmium 190.23(3)	77 Ir iridium 192.22	78 Pt platinum 195.08	79 Au gold 196.97	80 Hg mercury 200.59	81 Tl thallium 204.38, 204.38	82 Pb lead 207.2	83 Bi bismuth 208.98	84 Po polonium	85 At astatine	86 Rn radon	
87 Fr francium	88 Ra radium	89-103 actinoids	104 Rf rutherfordium	105 Db dubnium	106 Sg seaborgium	107 Bh bohrium	108 Hs hassium	109 Mt meitnerium	110 Ds darmstadtium	111 Rg roentgenium	112 Cn copernicium	113 Nh nihonium	114 Fl flerovium	115 Mc moscovium	116 Lv livermorium	117 Ts tennessine	118 Og oganesonium	

Substitutional variation:

- Sb^{III} can be substituted with 2+ and 3+ ions
- Sb^{V} can be substituted with 5+ ions
- Miscible with other rutile materials (4+ ions)

High entropy oxides:

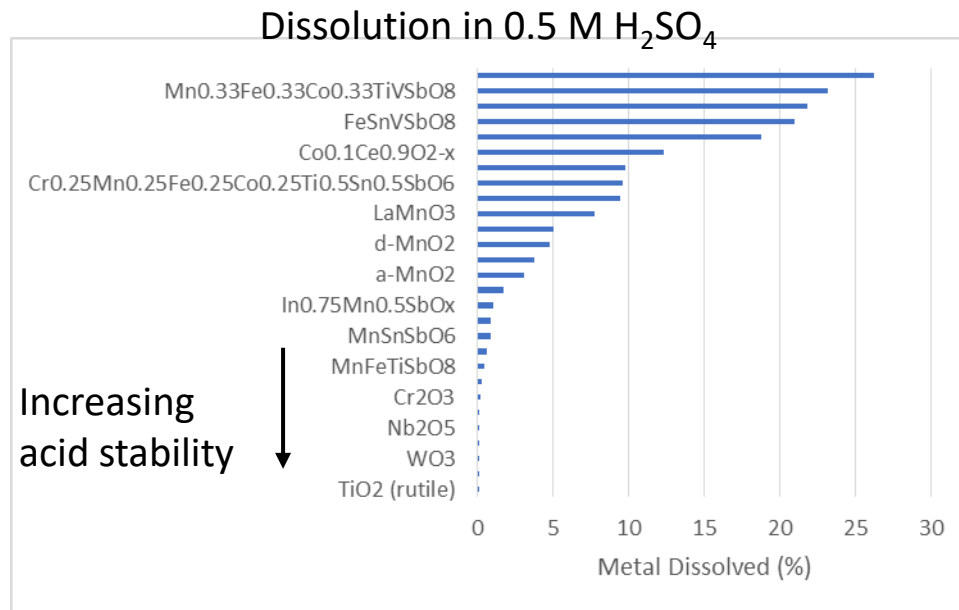
- Structure can stabilize disordering of up to 7 elements (demonstrated) at once
- Structure can accommodate wide range of ionic radii (68 – 94 pm)



Progress: Acid Stability Rutile Structures

Acid Stability:

- Rutile antimony oxides stabilize 3+ ions towards acidic attack
- Stabilized ions include Mn^{3+} , Fe^{3+} , In^{3+}
- Other stable elements can be added (Ti^{4+} , Cr^{3+} , Sn^{4+})



Electronic Conductivity:

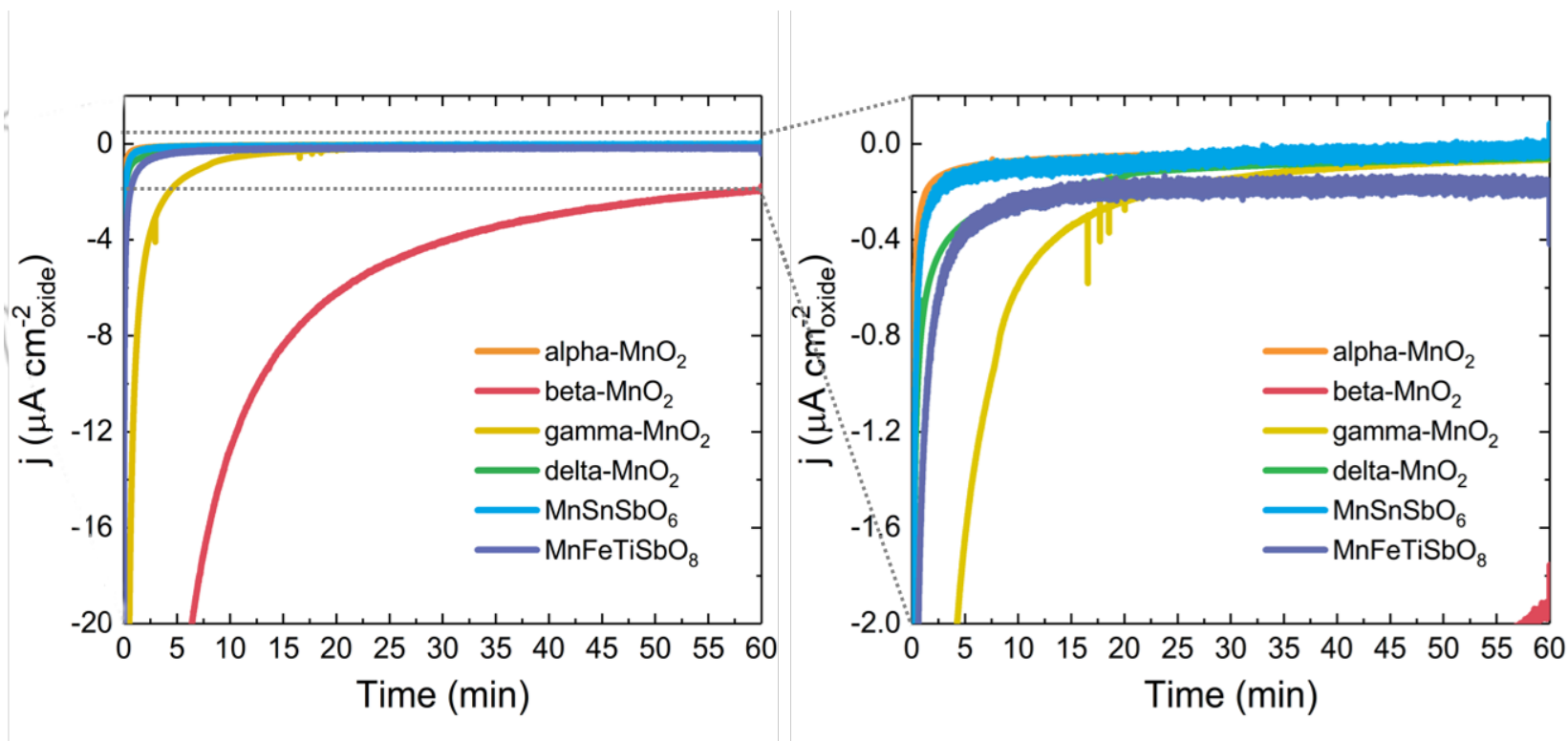
- Conductivity can be enhanced by addition of In^{3+} or Zn^{2+}

	Conductivity ($\text{mS}\cdot\text{cm}^{-1}$)
InSbO_4	220
$\text{In}_{0.75}\text{Mn}_{0.5}\text{SbO}_x$	230
MnO_2	0.005

Challenges:

- Incorporation of Mn^{4+} preferred but synthesis is challenging due to reduction
- Current Sb-based materials have low ORR activity

Progress: Electrochemical Activity



Activity of $\beta\text{-MnO}_2$ is significant over a 1-hour hold at 0.9 V
($\sim 2 \mu\text{A}\cdot\text{cm}^{-2}$ vs. initial target $0.5 \mu\text{A}\cdot\text{cm}^{-2}$, final target $4.4 \mu\text{A}\cdot\text{cm}^{-2}$)

Further durability not yet demonstrated

Mn-doped Sb-oxides are acid stable, but have low electrochemical activity to date.

Increasing Mn content (doped Mn oxide) is expected to demonstrate higher electrocatalytic activity

Progress: Computed Doped Mn Oxides

Materials Computed:

B-MnO₂ with the following substituents:

Active metals: Co, Fe, Ni

Stabilization: Ti, Cr, Sb, Sn, W, Nb, Cd, Ga

Conductivity: Zn, In

Predicted stable materials:

Likely ($\Delta\phi_{\text{hull}} < 0.1$ eV/atom):

Sb_xMn_{1-x}O₂ (x = 0.25, 0.5)

Nb_xMn_{1-x}O₂ (x = 0.25, 0.5)

Possible ($0.1 < \Delta\phi_{\text{hull}} < 0.2$ eV/atom):

Ti_xMn_{1-x}O₂ (x = 0.25, 0.5, 0.75)

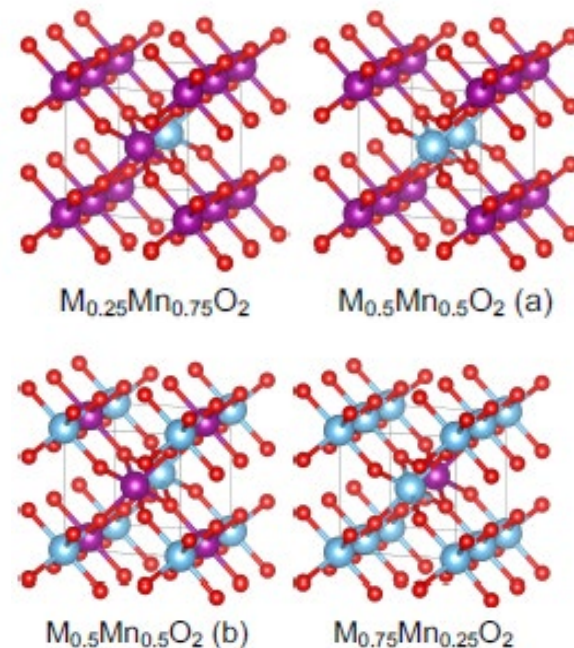
Fe_xMn_{1-x}O₂ (x = 0.25, 0.5)

Sn_xMn_{1-x}O₂ (x = 0.25, 0.5, 0.75)

W_xMn_{1-x}O₂ (x = 0.25, 0.5)

Co_xMn_{1-x}O₂ (x = 0.25, 0.75)

Structures Considered:



DFT stability calculations for a range of ternary manganese oxides were performed to determine further acid stable synthesis targets.

Responses to Reviewer's Comments

“The project offers an alternative to the highly unstable metal nitrogen-doped carbon catalyst. If successful, this project might have significant impact on DOE technical targets.”

“The Go/No-go should include simultaneously demonstrating voltage performance and acid stability with the same formulation.”

- The project team agrees with this assessment.

“There is a lack of evidence that complex oxides can be active, stable, and conductive under the relevant conditions for PEMFCs”

- The project team has identified oxides that are both stable and conductive under relevant conditions and are working to increase the activity.

Collaborations

First principles design to membrane-electrode assembly

Subcontractor, University



PI: Prof. Yang Shao-Horn

- Oxide optimization for acid stability
- ORR Electrocatalytic performance optimization of acid-stable oxides

Electrocat Consortium:

Collaborator: Dr. Deborah Myers

- High-throughput synthesis of doped oxides

Prime, Industry



PI: Tim Davenport

- Catalyst Layer Optimization
- MEA Fabrication
- MEA Performance and Durability Testing

Challenges and Barriers

- **Challenge:** Identifying an oxide with sufficient electrocatalytic activity to meet the project goals has not yet been achieved. Successful identification of such a material is necessary to meet the Go/No-go milestone.
- **Planned Resolution:** The project team obtained a no-cost extension to meet the electrocatalyst targets.
- **Challenge:** Integration of the electrocatalyst material into a membrane-electrode assembly has not begun.
- **Planned Resolution:** The project team will leverage the high-throughput capabilities of the Electrocat consortium to optimize the MEA fabrication.

Proposed Future Work

Project Milestones	Schedule
Demonstrate intrinsic ORR activity $\geq 4.4 \mu\text{A}\cdot\text{cm}^{-2}_{\text{oxide}}$ at 0.9 V (iR-free) with an acid stable oxide	8/31/2020
Demonstrate MEA with performance of $0.025 \text{ A}\cdot\text{cm}^{-2}$ at 0.9 V (iR-free) under 1 atm O_2 and 80 °C	8/31/20
Demonstrate MEA with PGM-free ORR catalyst that meets targets FC-4 and FC-5 under “Electrocatalyst Cycle” AST protocol.	2/28/21
Demonstrate MEA with PGM-free ORR catalyst that meets targets FC-6 and FC-7 under “Catalyst Support Cycle” AST protocol.	5/31/21

- High-throughput techniques will be leveraged to synthesize targeted acid-stable oxide materials
- As identification of a highly active electrocatalyst is critical, alternative electrocatalyst materials will be considered in place of an oxide.

Any proposed future work is subject to change based on funding levels

Summary

- Discovery of acid-stable metal oxides with catalytically active elements has the potential for breakthrough ORR electrocatalytic performance
- Descriptors for acid stability have been developed to aid acid-stable oxide electrocatalyst identification
- A family of antimony-based rutile oxides has been developed exhibiting acid-stability and a wide composition space. Conductive oxide materials were identified.

Go/No-Go Technical Target:

Demonstrate MEA with performance of 0.025 A-cm⁻² at 0.9 V under 1 atm O₂ and 80 °C

MYRD&D Targets Addressed:

Target I.D. #	Characteristic	Units	2020 Targets
FC-4	Loss in initial catalytic activity	% mass loss	< 40
FC-5	Loss in performance at 0.8 A cm ⁻²	mV	< 30
FC-8	PGM-free catalyst activity	A cm ⁻² at 900 mV _{iR-free}	> 0.044

Develop durable MEAs with PGM-free metal oxide ORR catalysts

Acknowledgements

Jiayu Peng, MIT
Dr. Livia Giordano, MIT
Prof. Yang Shao-Horn, MIT



Donna Ho, DOE
John Kopasz, ANL

Simon Thompson, DOE
Dimitrios Papageorgopoulos, DOE

Electrocat Consortium

