

# **Novel, Combinatorial Method for Developing Cathode Catalysts for Fuel Cells**

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This presentation does not contain any  
proprietary or confidential information

Project ID: # FCP30

# Overview

## Timeline

- Start Date: October, 2004
- End Date: December, 2007
- 75% Completed

## Budget

- Phase II SBIR
- Total Project Funding
  - \$750,000
- 2005 Funding: \$246,000
- 2006 Budget: \$420,000

## Barriers

- Low activity of non-Pt catalysts
  - 2004 Status: 8 A/cm<sup>3</sup>
  - 2010 Target: >130 A/cm<sup>3</sup>

## Partners

- Illinois Institute of Technology

# Need for New Fuel Cell Cathode Catalyst

- **Automotive Applications:**
  - Order of magnitude improvement over current Pt alloy based MEA's.
  - Cost – \$10/kW MEA Cost
  - High Efficiency – 0.2 g/peak kW total anode/cathode loading.
  - Long Life – 10-15 years life

# Project Objectives

- **Develop a controlled method for accurate high-throughput evaluation of new catalyst materials.**
- **Scale up combinatorial approach: Sample preparation, screening system and data processing.**
- **Evaluate several families of catalysts for oxygen reduction activity.**
- **Scale up new, low-cost high-activity catalysts for evaluation in fuel cells.**
- **Develop instrument for efficient evaluation of multiple fuel cell components (catalysts, membranes, MEA's, etc) for general use in process development and manufacturing quality control.**

# Why Combinatorial Approach for Catalyst Development?

- **Barriers to rational design.**
  - Complex surface chemistry.
  - Lack of a complete understanding of the reaction processes involved.
- **Many possible catalyst permutations (not confined by equilibrium phases).**
- **Screening in parallel allows for better evaluation of relative performance.**
- **Can potentially greatly reduce the cost of optimization and accelerate the discovery of new catalysts.**



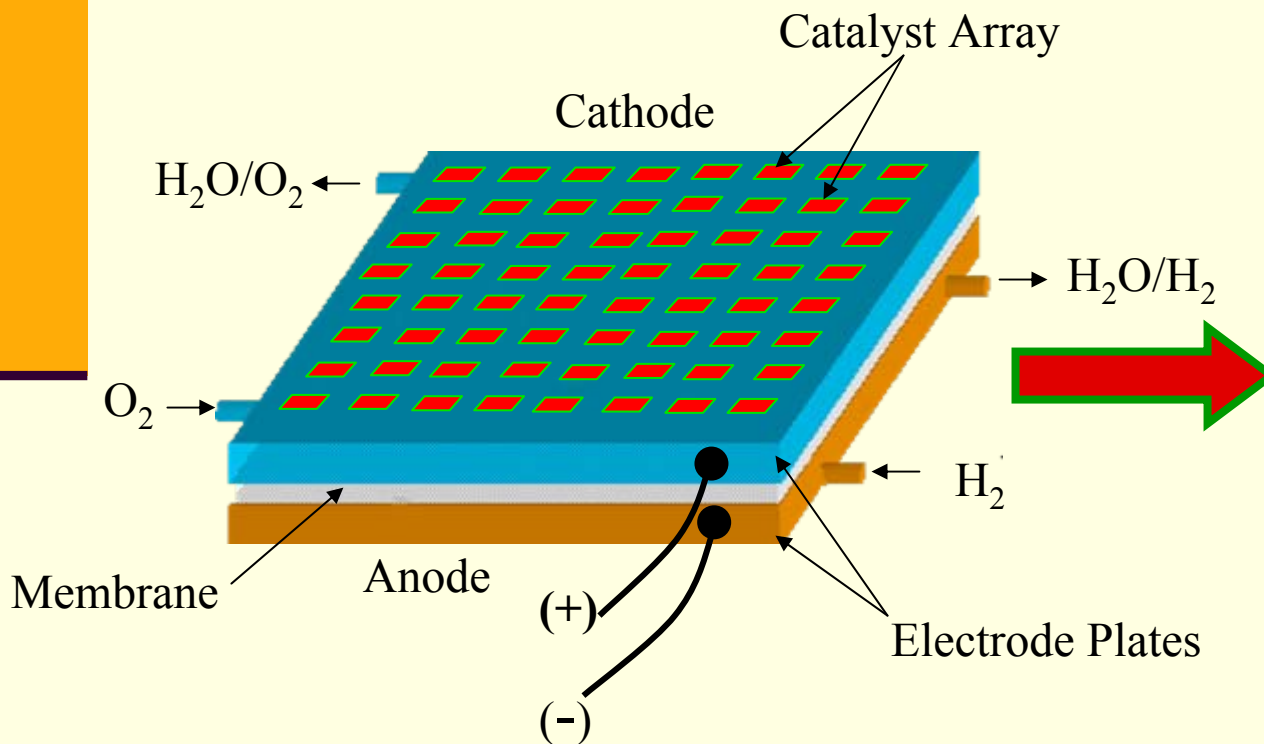
# Phase II Project Catalyst Development Strategy

- **Identify best chemistry first then optimize for utilization.**
- **Control all critical parameters to determine inherent catalyst activity.**
- **Use systematic DOE techniques to design catalyst array compositions and testing condition variables.**

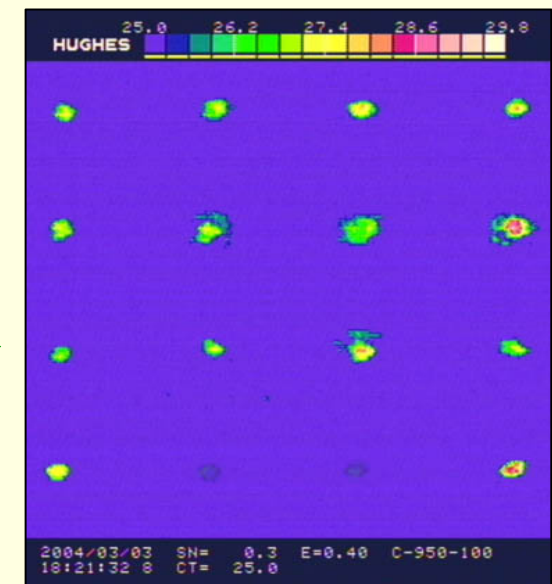
# Technical Approach

## Thermal Sensing

Thermal sensing allows for in-situ monitoring of individual catalysts samples in a closed fuel cell system.

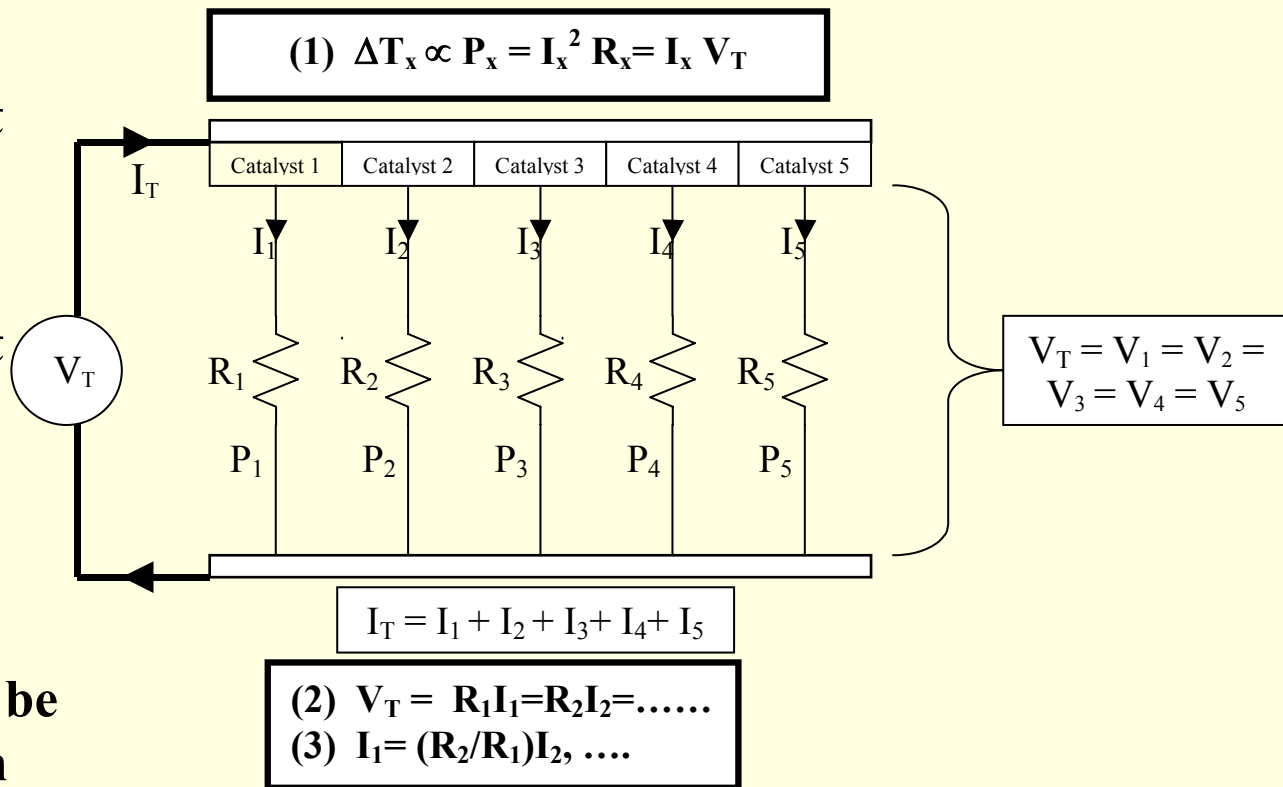


Thermal Image



# Heat Generation and Catalyst Efficiency

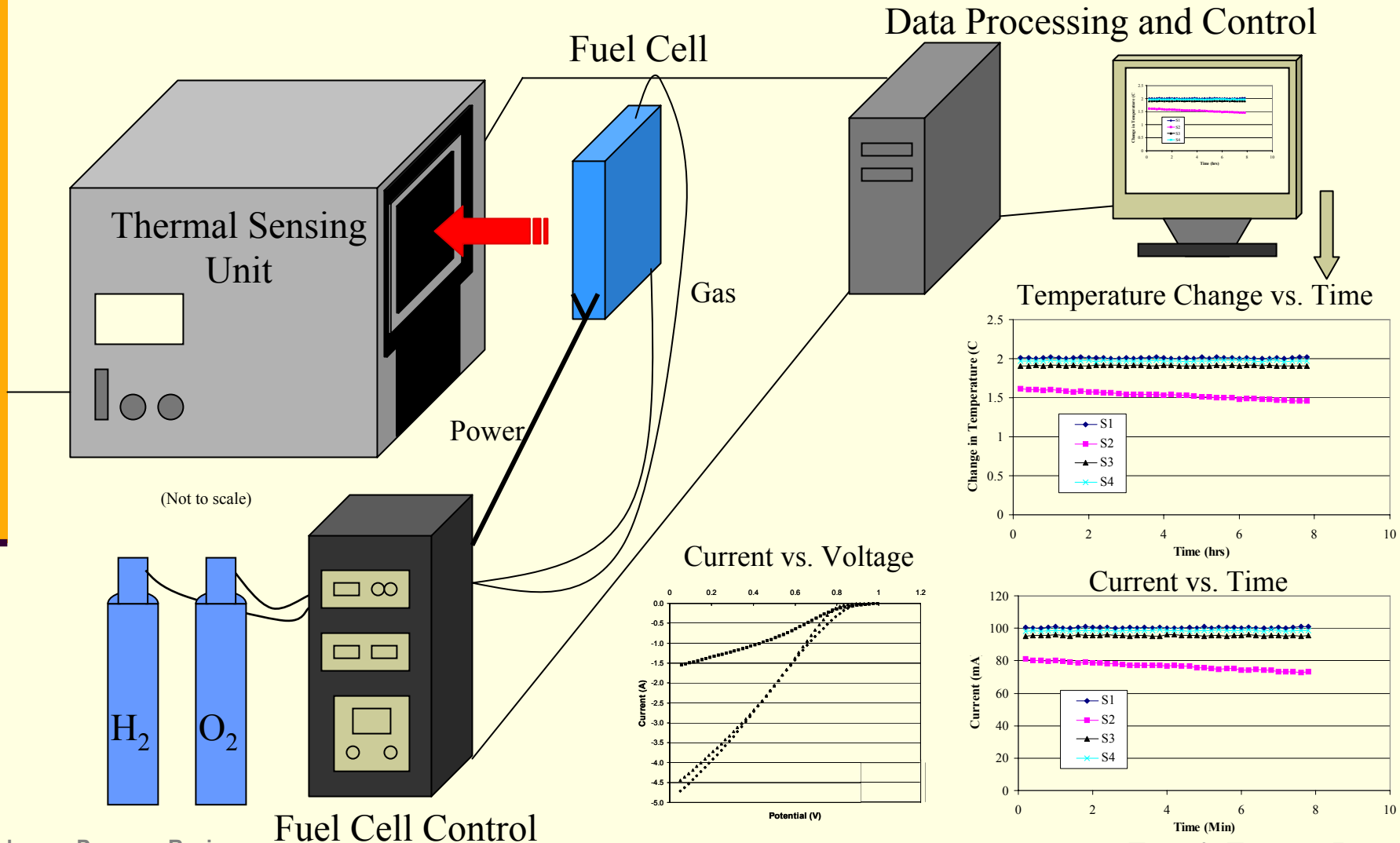
- Correlation between  $i^2R$  heat generated and current density.
- The best catalyst will generate the most heat.
- The current passing through each sample can be determined from  $dT$ .



Platinum/0.2V  $\sim 10^{-3}$  W/cm<sup>2</sup>  
 Carbon/0.2V  $\sim 10^{-6}$  W/cm<sup>2</sup>



# Fuel Cell Catalyst Screening System



(Not to scale)

Fuel Cell Control



# Technical Approach Advantages

- In-situ screening under real operating conditions.
- Good control of critical parameters that affect performance.
- Great flexibility to screen any catalyst type for any fuel cell system.
- Simple, low-cost system scale-up.

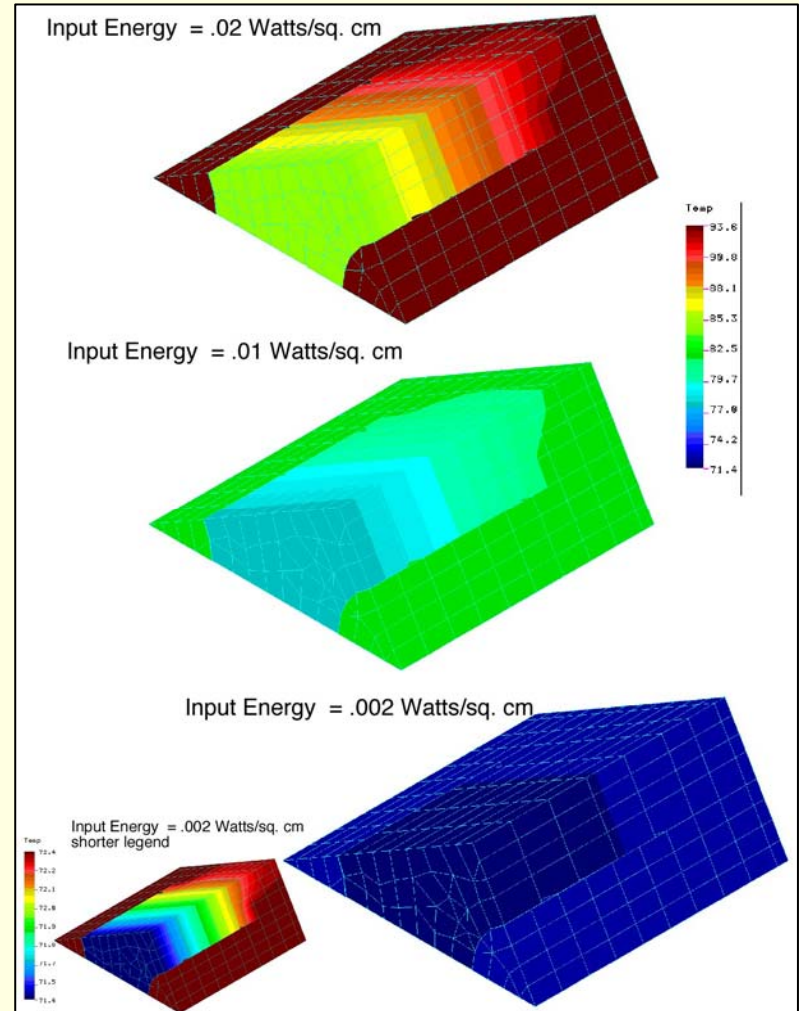


# Technical Accomplishments/ Progress/Results

- **Finalized Gen 1 screening system design and verified performance.**
  - Uniform stack pressure.
  - Uniform fuel distribution.
  - Uniform heat signal.
- **Prototype Gen 2 screening system developed.**
- **Developed high-throughput sample preparation system.**
- **Exploration of catalyst families.**
- **Detailed characterization of binary Pd-Co catalyst system.**

# Thermal Modeling to Aid System Design

- **Developed design in miniature before scale up.**
  - 35 cm<sup>2</sup> array cell.
  - Accelerates development cycle.
  - Lowers cost of development.
    - Smaller MEA's
    - Fewer Samples.
    - Less Labor.



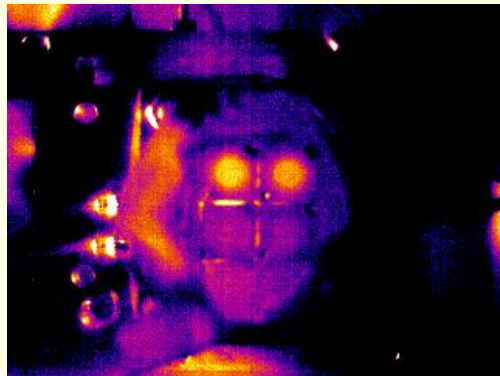
# Qualifying Gen 1 System Design

- **Qualify on 4-sample array apparatus before scale up.**
- **Qualification Procedure.**
  - Demonstrate correlation between current and temperature.
  - Verify uniform fuel flow.
  - Verify uniform stack pressure.
  - Verify evaluation of constant catalyst surface area across array.
- **Scale-up to 25-sample array apparatus.**
- **Catalyst Screening.**

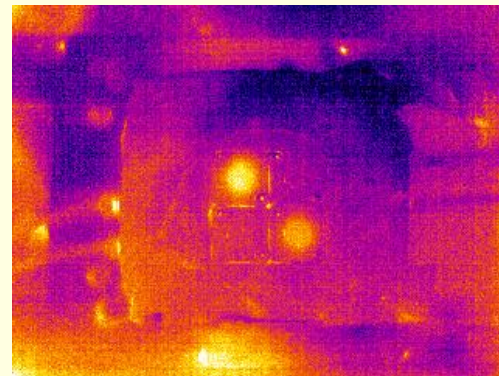
# Gen 1 System: Thermal Signal Correlations

## 4-Sample Array

2x Pt (top) vs. 2x Carbon (bottom)

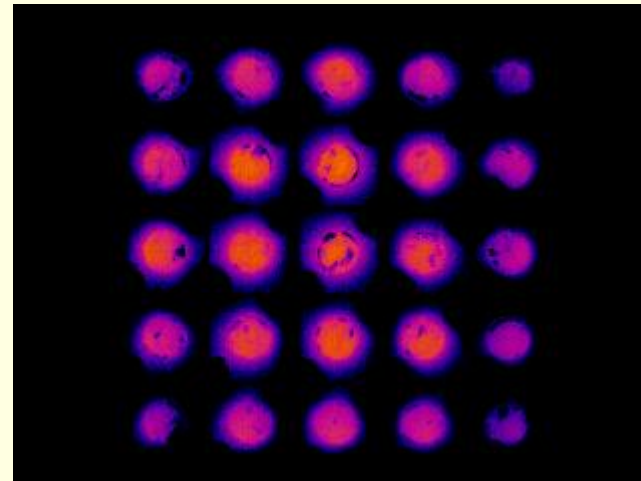


After switching right-side samples



## 25-Sample Array

- 55 °C Operation
- H<sub>2</sub>/O<sub>2</sub>
- Binary array
- $\Delta T \sim 2^\circ\text{C}$
- Hot spots – highest activity



# Stage 2 Catalyst Development

- **Best identified catalyst families are further characterized by conventional methods.**
- **Electrodeposited catalyst samples – CV's, Rotating Disk.**
- **Carbon supported catalyst – MEA's, H<sub>2</sub> fuel cell.**

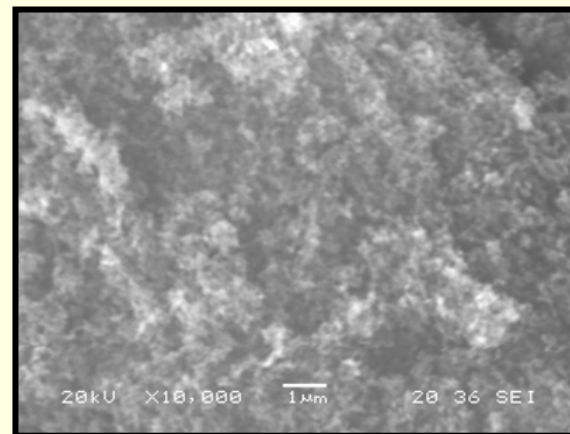
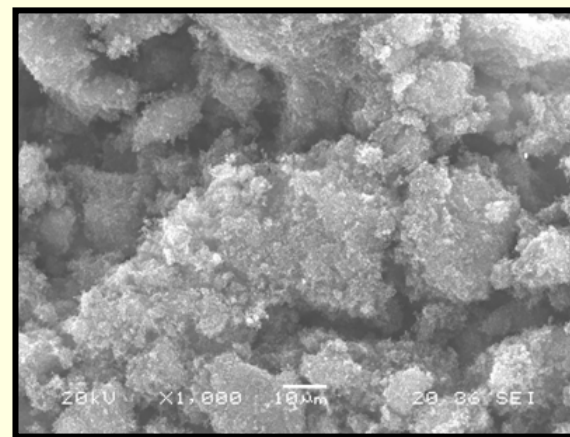
# Preparation and Characterization of Carbon Supported $\text{CoPd}_x$ Electrocatalysts

## ■ Preparation

- Deposit Salts using Sodium Bicarbonate as reducing agent.  $\text{Co}[\text{NO}_3]_2 \cdot 6\text{H}_2\text{O}$ ,  $\text{Pd}[\text{NO}_3]_2$
- Catalyst filtered, rinsed, vacuum dried and activated in a 2% $\text{H}_2$ , 98% $\text{Ar}$  atmosphere for 24 hours.
- Loading 10 wt %  $\text{CoPd}_x$  on Vulcan XC72R

## ■ Characterization

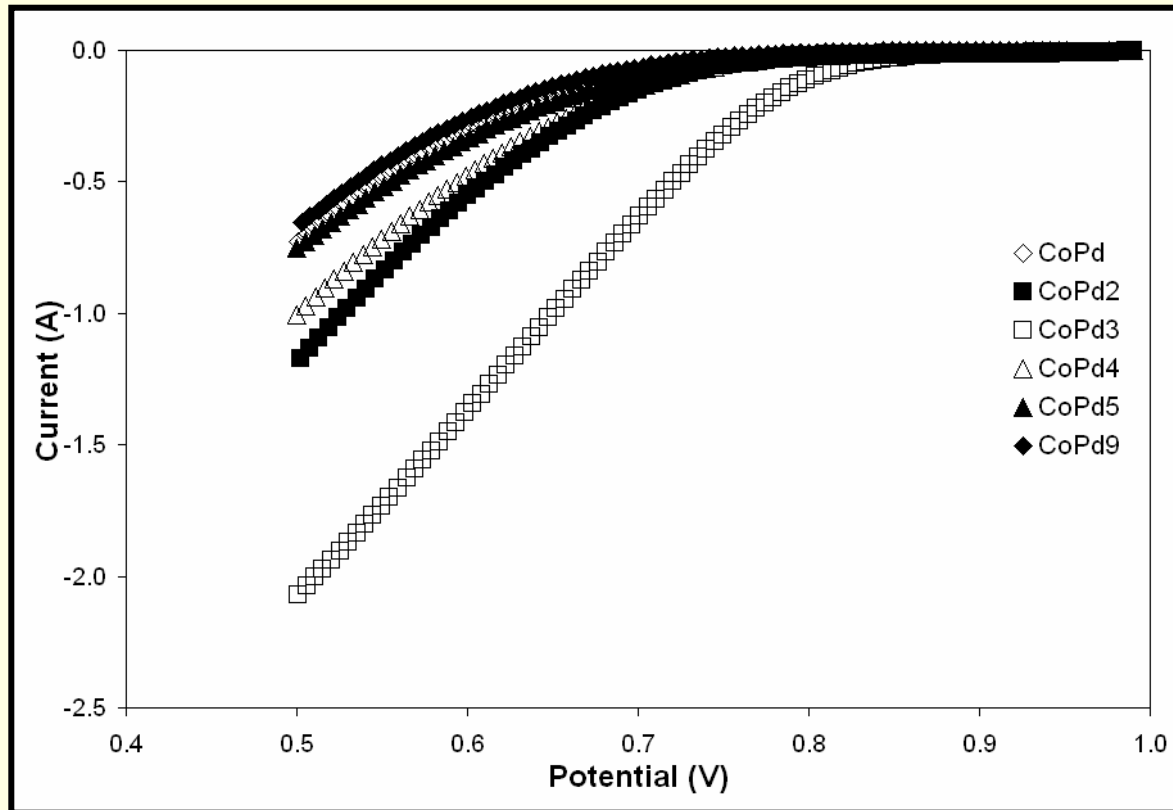
- SEM/EDX, BET





# Co-Pd Catalyst Family with High Activity

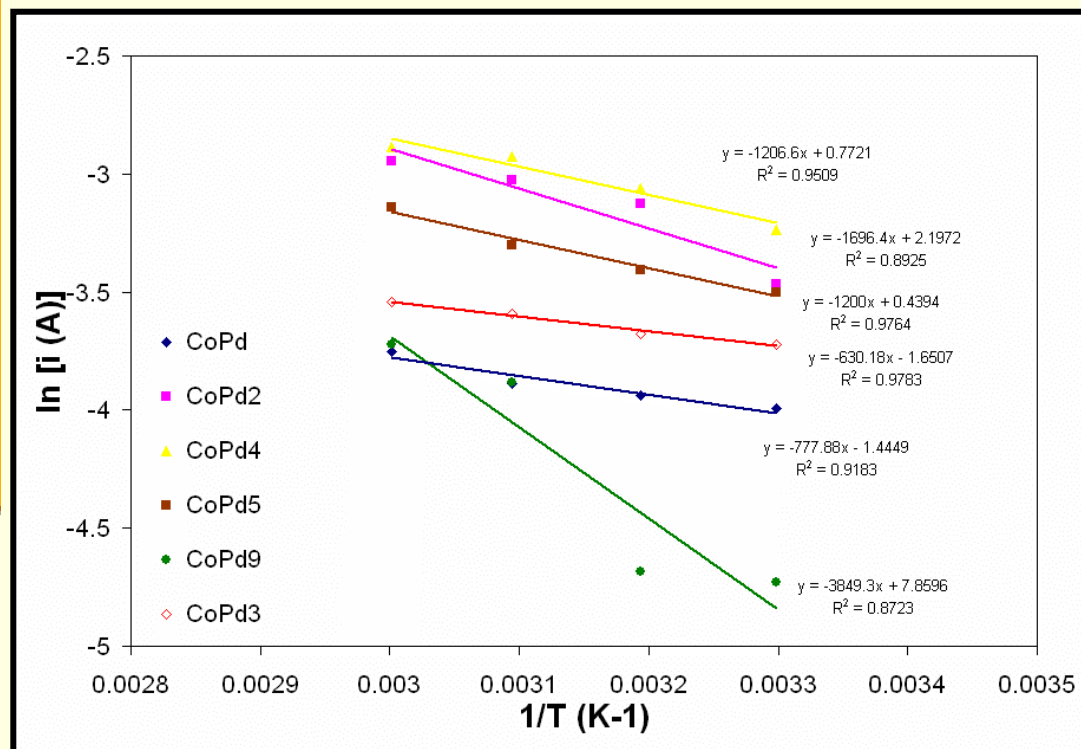
## Polarization Curves for Co-Pd compositions



- Temperature: 60°C
- Cathode gas: O<sub>2</sub>
- Anode gas: H<sub>2</sub>

# CoPd<sub>x</sub> Kinetic Parameters

## Arrhenius Plots for the ORR in a 5 cm<sup>2</sup> PEMFC on CoPd<sub>x</sub> Electrocatalysts

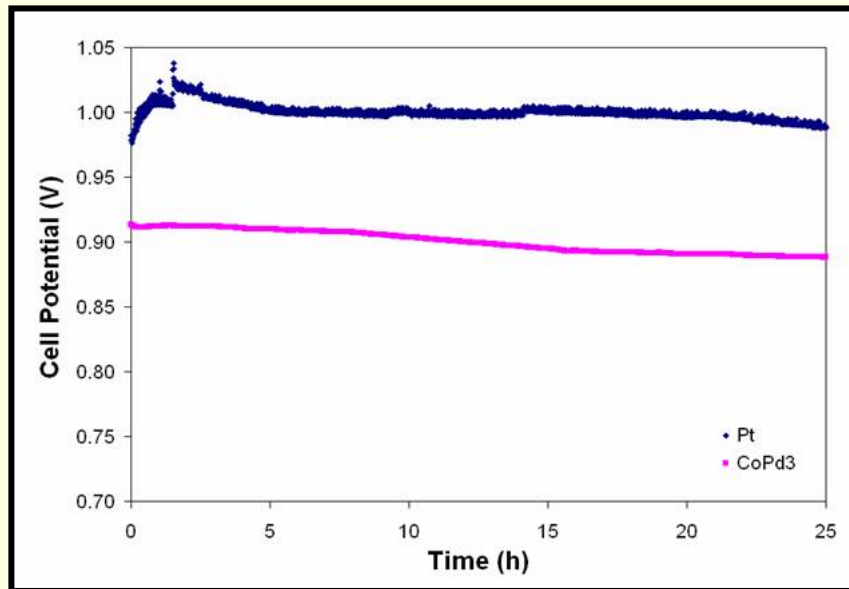


## Calculated Kinetic Parameters for ORR on CoPd<sub>x</sub> Electrocatalysts in a H<sub>2</sub>/O<sub>2</sub> PEMFC

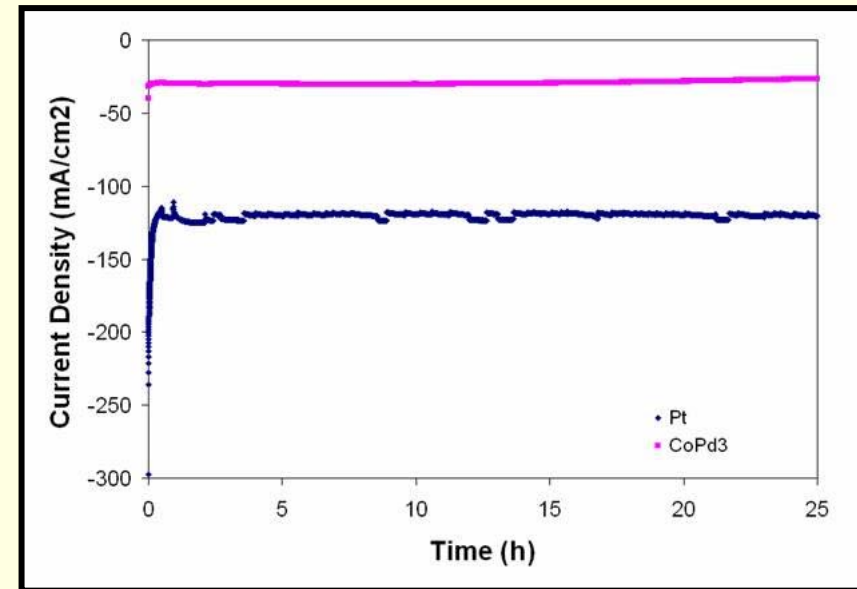
Composition	Onset Potential (V)	Activation Energy (kJ/mol)	Tafel Slope, b 60°C (mV/dec)	Exp[a/b]
CoPd	0.87	80.4	96.5	4.08E+02
CoPd <sub>2</sub>	0.88	104.0	87.1	1.23E+03
CoPd <sub>3</sub>	0.92	52.4	69.6	2.69E+04
CoPd <sub>4</sub>	0.89	100.3	90.3	7.69E+02
CoPd <sub>5</sub>	0.90	99.8	100.1	2.43E+02
CoPd <sub>9</sub>	0.89	320.0	34.8	1.72E+08

# Performance Stability of Pd<sub>3</sub>Co in Hydrogen Fuel Cell

Open Circuit Voltage Pt and CoPd<sub>3</sub>-  
cathode/ Pt-anode MEAs in a 5 cm<sup>2</sup>  
PEMFC at 60 °C.



Performance Stability of Pt and CoPd<sub>3</sub>  
cathode MEAs at 0.8 V, 60 °C.



- Some performance degradation of PdCo<sub>3</sub> catalyst observed.

# Future Work

- **Scale up Gen 1 screening system to 50-100 samples/cell.**
- **Scale up Gen 2 screening system.**
- **Continue large scale screening of non-noble metal catalysts.**
- **Verify results in standard fuel cells.**
- **Continue Pd-Co development.**

# Summary

- **We have developed an easily scalable method of combinatorially screening materials for electrochemical systems based on their efficiency related thermal signature.**
- **We are using this system to evaluate catalysts for oxygen reduction activity.**
- **Materials with the greatest potential are further characterized and optimized by conventional methods.**
- **Our combinatorial technique and development strategy greatly increase our probability of success and decrease our discovery time.**

# Publications

- Mustain, W.E.; Kepler, K. D.; Prakash, J.; “Investigations of Carbon-Supported CoPd<sub>3</sub> Catalysts as Oxygen Cathodes in PEM Fuel Cells”, *Electrochemistry Communications*, 8 (2006) 406-410.
- Mustain, W.E.; Kepler, K. D.; Prakash, J.; “CoPd<sub>x</sub> Alloys as Oxygen Reduction Electrocatalysts for Polymer Electrolyte Membrane and Direct Methanol Fuel Cells”, submitted for publication.