



Company and Technology Overview

Hydrogen Technical Advisory Committee

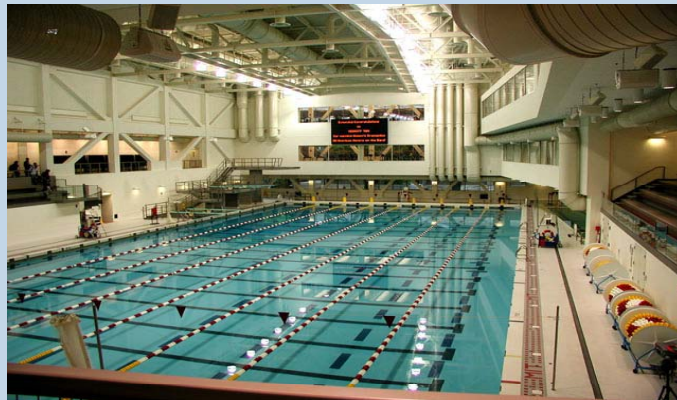
14 October 2010

Transforming the Global Energy Paradigm

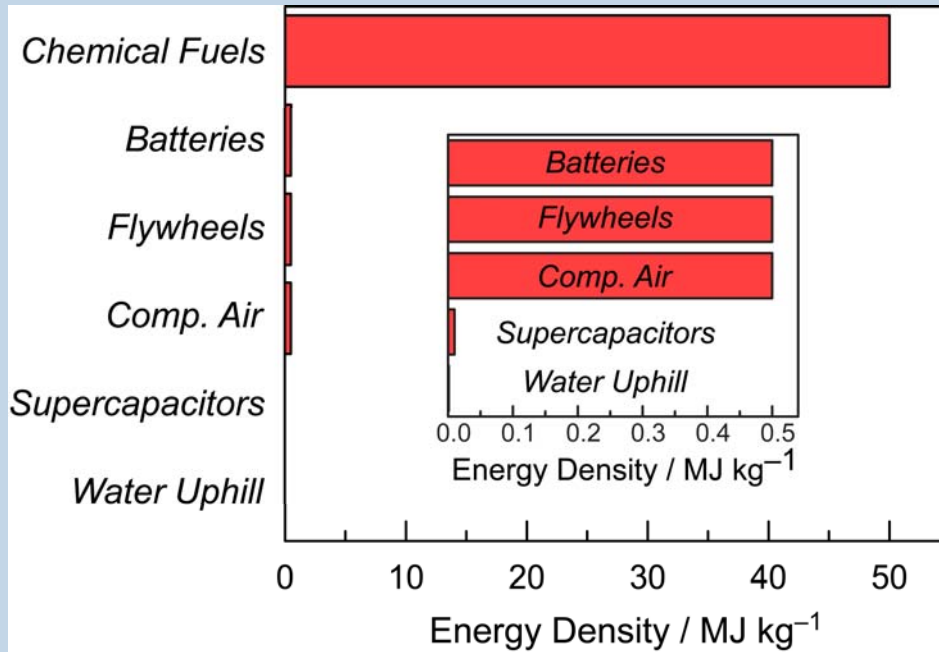
- **Vision: Sunlight + Water = Affordable, Accessible Energy**
 - *Sunlight*: Most Abundant Energy Source
 - *Water*: Most Abundant Liquid
 - *Sun Catalytix Technology*: Accessible and Inexpensive
- **Impact**
 - *Affordable Energy Storage*
 - *Distributed Fuel Production*
 - *Every Home and Community Becomes an Energy Production Center*
 - Personalized energy paradigm consistent with prior shifts
 - Land lines to cell phones
 - Mainframes to personal computers

Sunlight + Water: Near Limitless Energy Supply

- Global energy usage likely to double to ~30TW by 2050
- 1.2×10^5 TW of Solar at Earth's surface; 800 TW practical
- Other renewables limited in scale
 - Biomass: 5-7 TW gross (with all arable land not used for food)
 - Wind: 2-4 TW total extractable
 - Geothermal: 12 TW gross over land small fraction recoverable
- *Volume of MIT pool $\rightarrow H_2$ and O_2 per second = 43 TW*

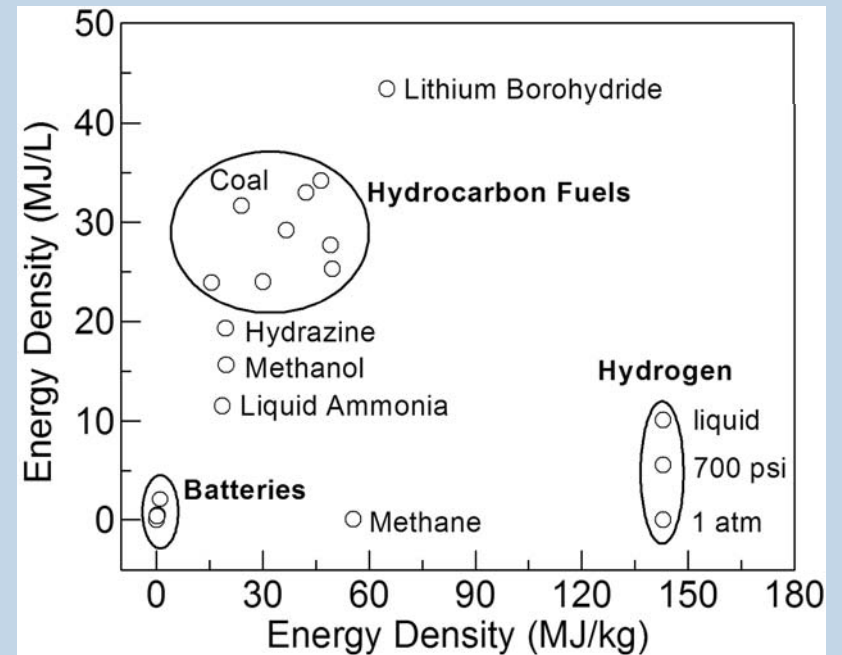


Energy Storage in Chemical Fuels



Chemical fuels possess energy densities unmatched by any other storage means

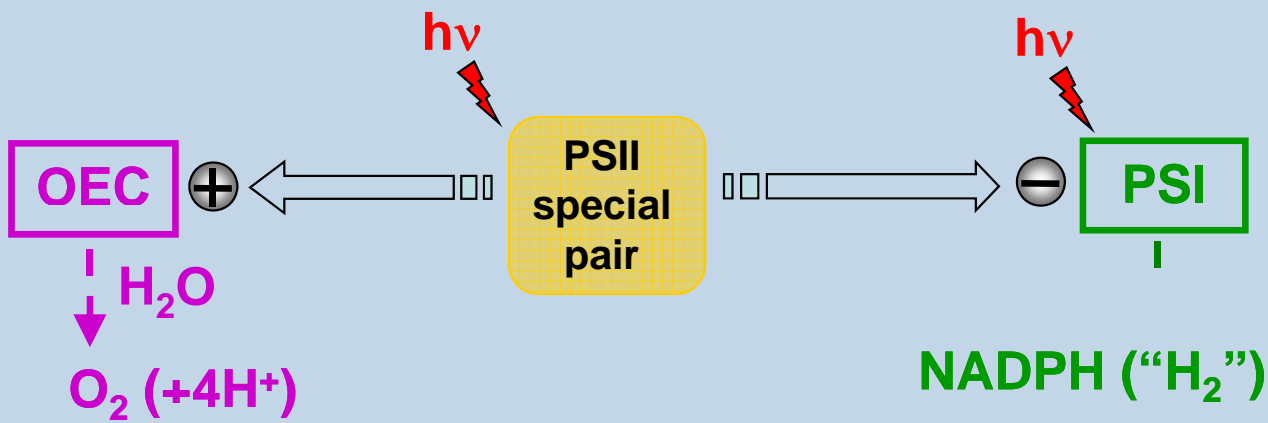
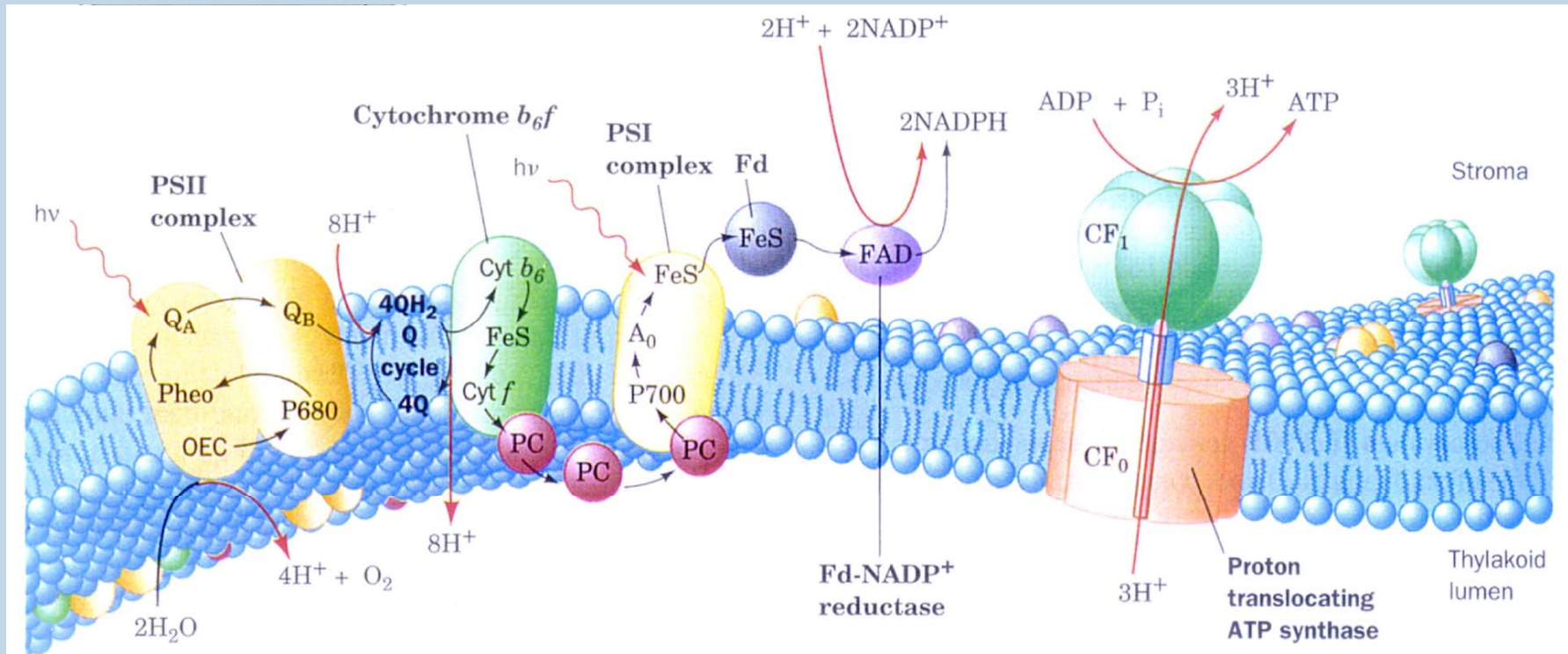
Hydrogen is the most energy dense fuel by mass



Target Applications

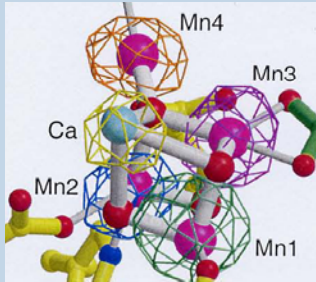
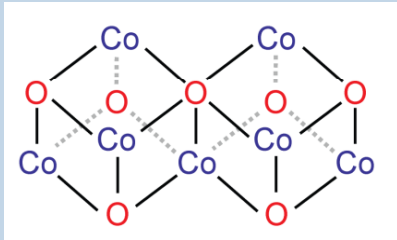
- **On-site industrial hydrogen**
 - For various applications: semiconductor fabrication, metal annealing, generator cooling, etc.
- **Energy storage**
 - Off grid power
 - On-grid storage
- **Fuels**
 - Automotive traction
 - Hydrogen as a fuel precursor (*e.g.*, renewable diesel)

Natural Photosynthesis

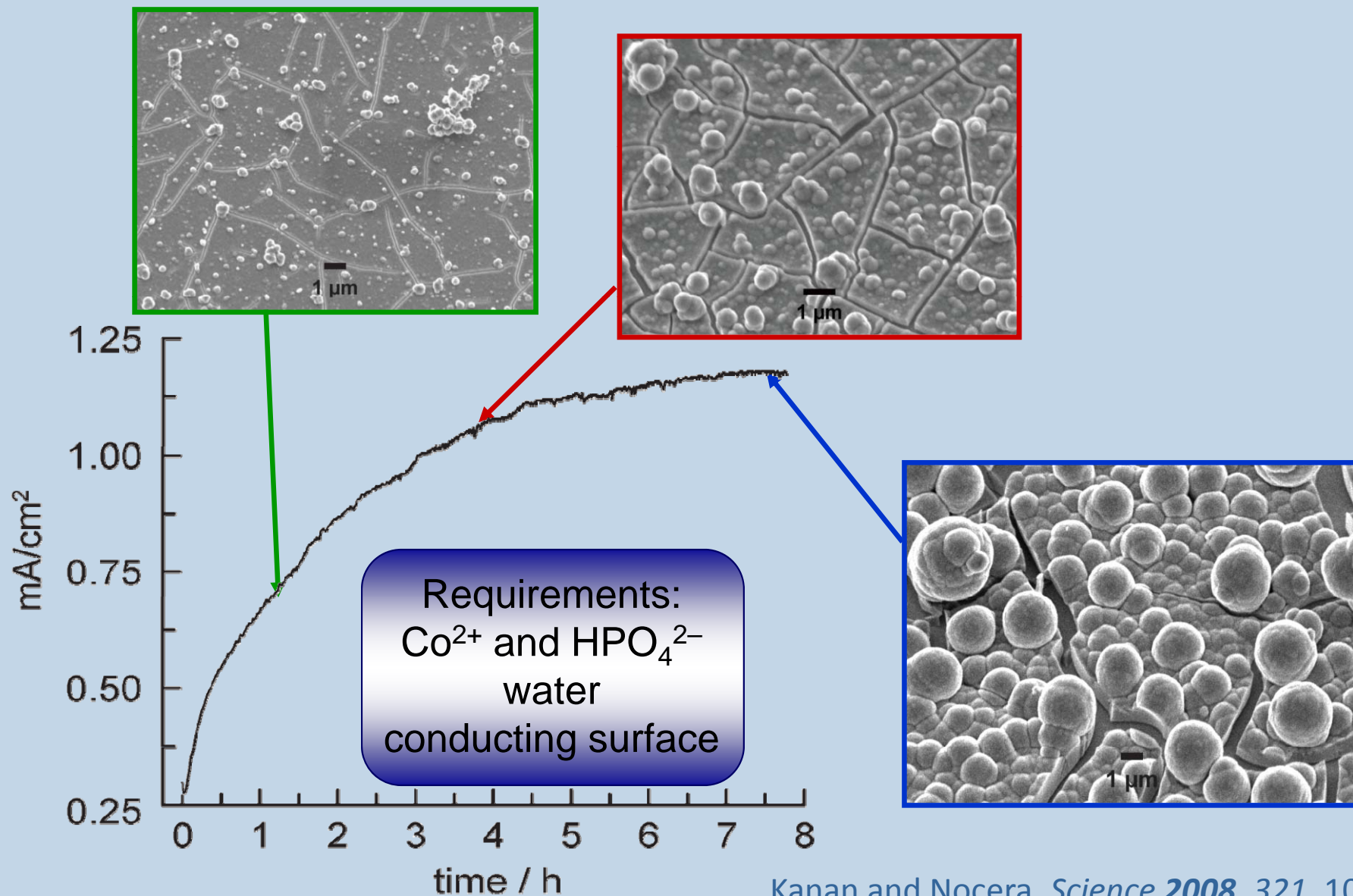


- (1) catalysis is performed under benign conditions at neutral pH
- (2) generation of OEC for water oxidation is key evolutionary step
- (3) photon sets up a (wireless) current within the leaf
- (4) separate light harvesting from energy storage

Our Technology Mimics Nature's Processes

Design Criteria	Photo-system II	Cobalt-Phosphate
<i>Conditions of operation</i>	Neutral pH and STP Low over-potential	Neutral pH and STP Low over-potential
<i>Repair</i>	D1 protein	$\text{HPO}_4^{2-}/\text{Co}^{3+}$ equilibrium
<i>Water Supply</i>	Fresh or Salt	Fresh or Salt
<i>Stability</i>	Regenerative	Regenerative
<i>Abundant materials</i>	Manganese	Cobalt
<i>Structure</i>		

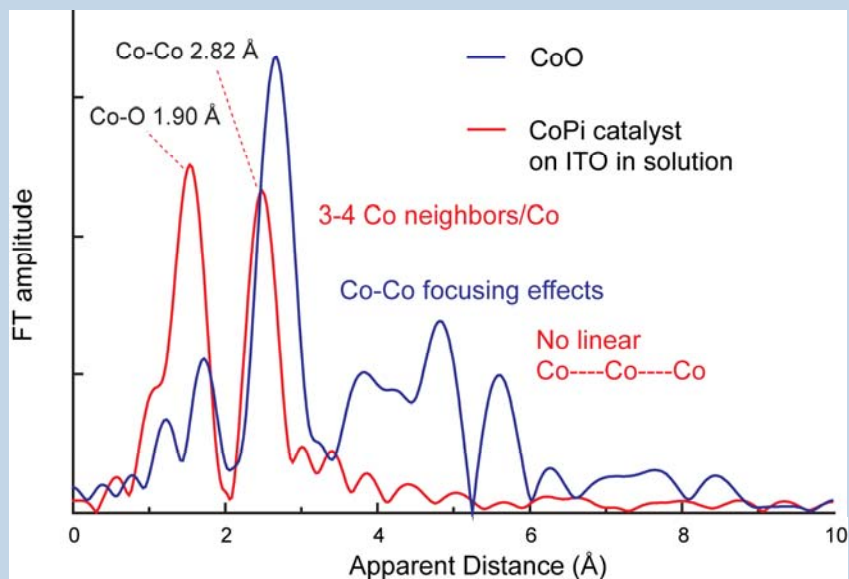
In Situ Formation of an Oxygen-Evolving Catalyst



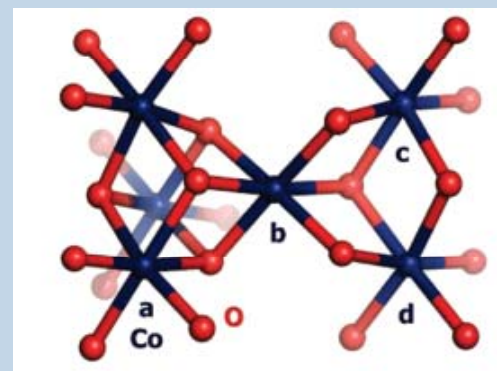
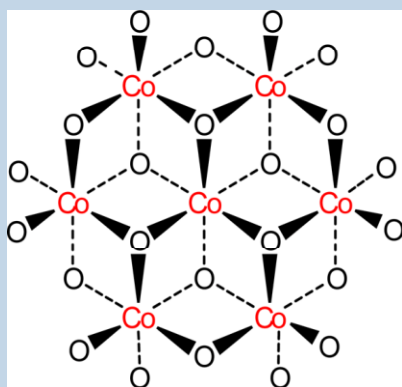
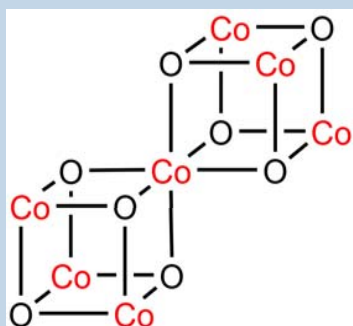
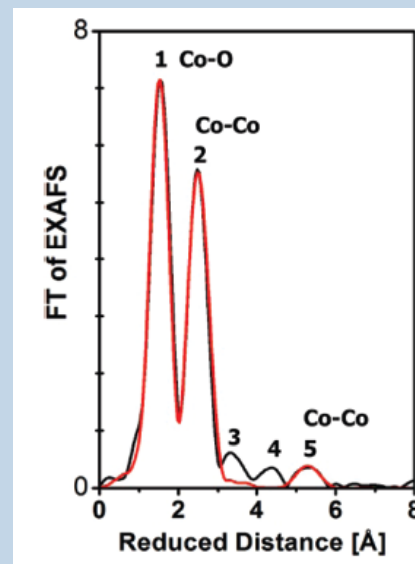
Kanan and Nocera, *Science* **2008**, 321, 1072.

Structural Insight from EXAFS

Not a typical Oxide material



Confirmed by a German group



With Junko Yano and Vittal Yachandra at LBNL

Dau and coworkers *JACS* **2009**, *131*, 6936

Two Modalities for Sun Catalytix Technology

1. Electrolyzer Modality

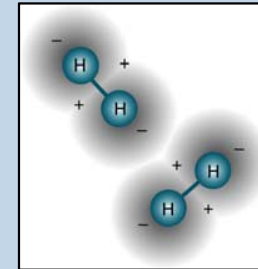
Solar, Wind or other electricity source



Sun Catalytix Electrolyzer

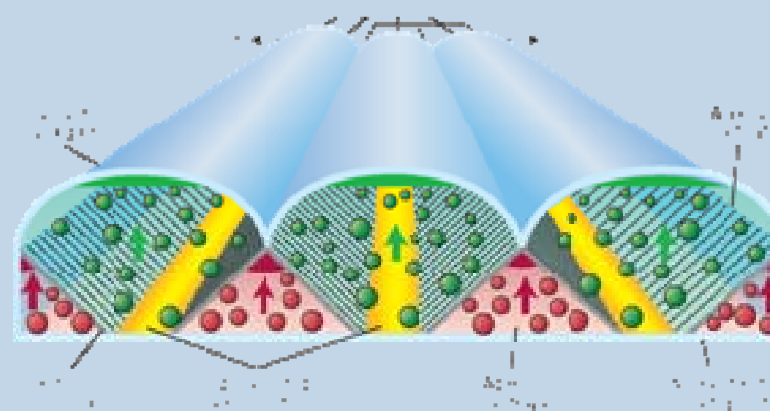
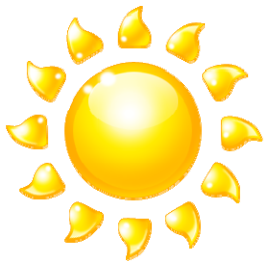


Hydrogen

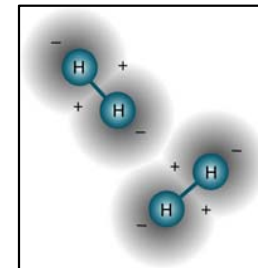


2. Photoelectrochemical (PEC) Modality

Sunlight

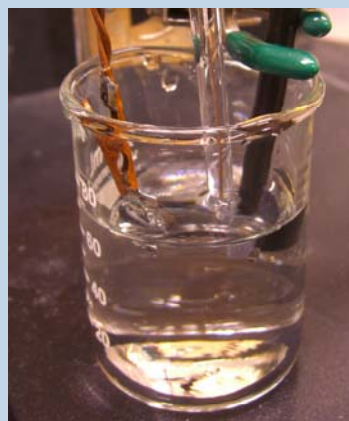


Hydrogen

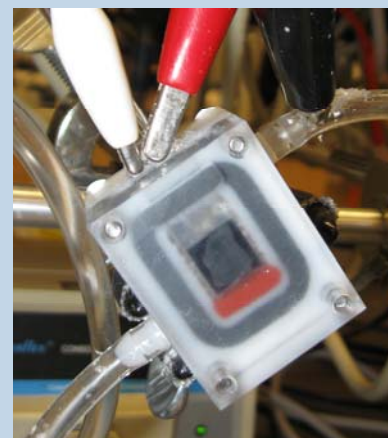


Electrolyzer Technology Status

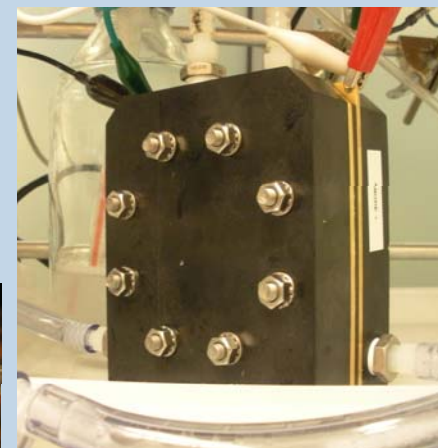
- Anode catalyst performance meets initial target
- Cathode catalyst performance meets initial target
- Membrane / separator criteria defined, initial materials selected
- Integration of cell components, electrode design, performance enhancement underway



2008



2009

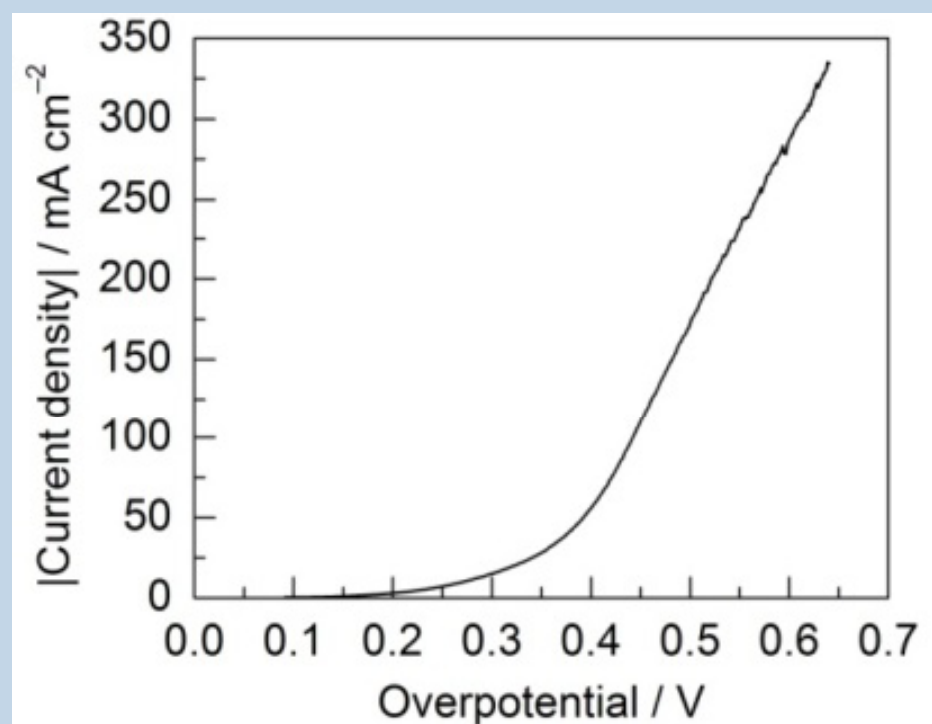
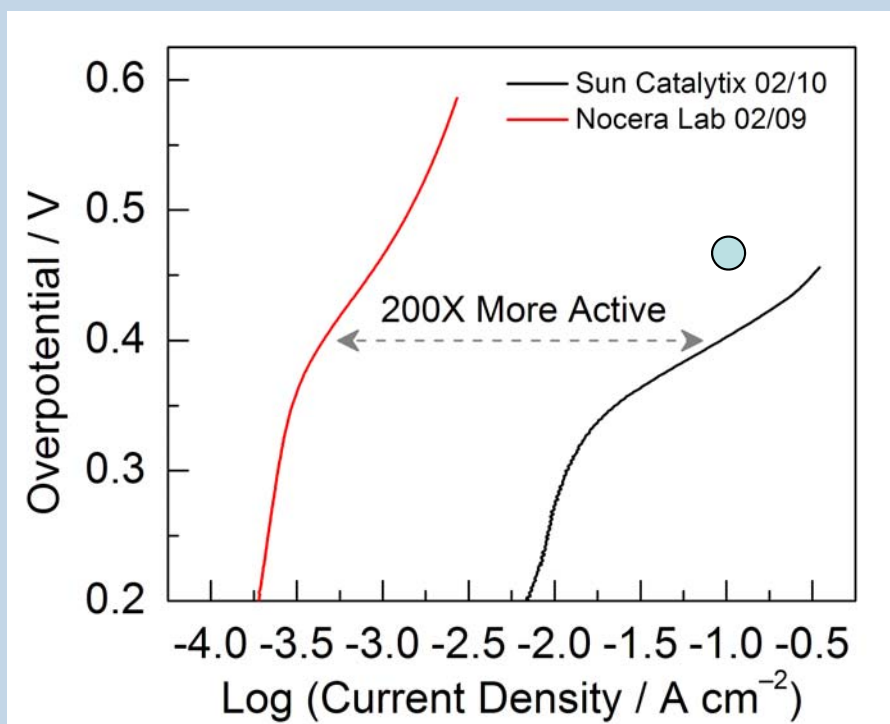


2010

Electrolysis Performance Enhancement

Electrode design has enabled $\sim 200\times$ current enhancement relative to results published in Kanan and Nocera, *Science* **2008**, 321, 1072.

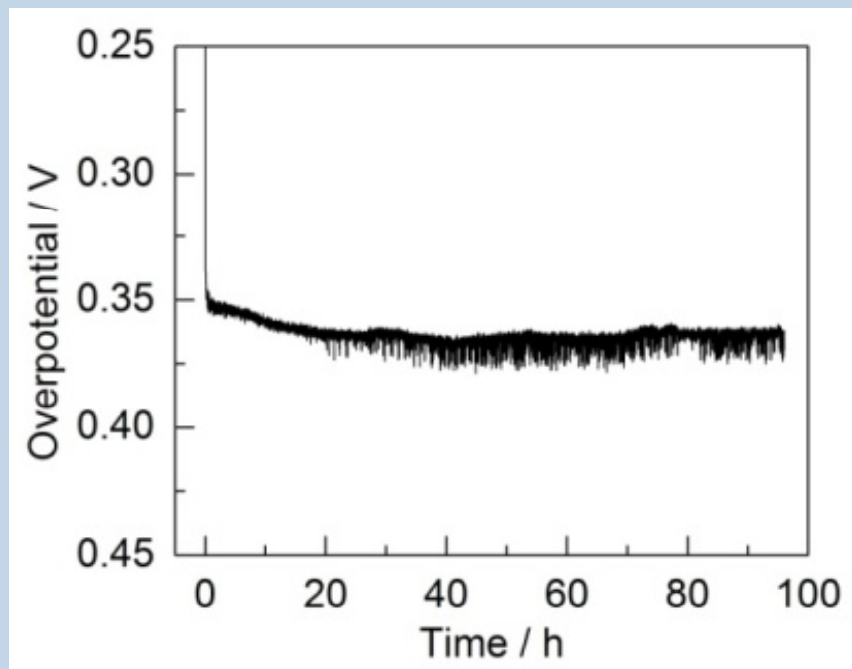
Anode Activity Enhancement



Esswein et. al., Submitted to *Energy and Environmental Science*

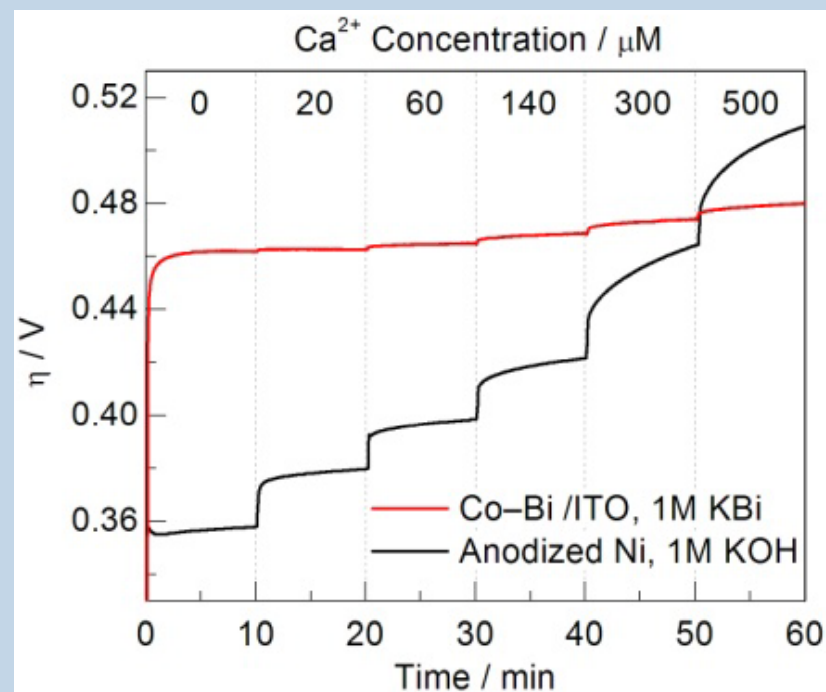
Anode Catalyst Capability

Anode Stability



Engineered Co-OEC catalysts exhibits high stability in untreated river water

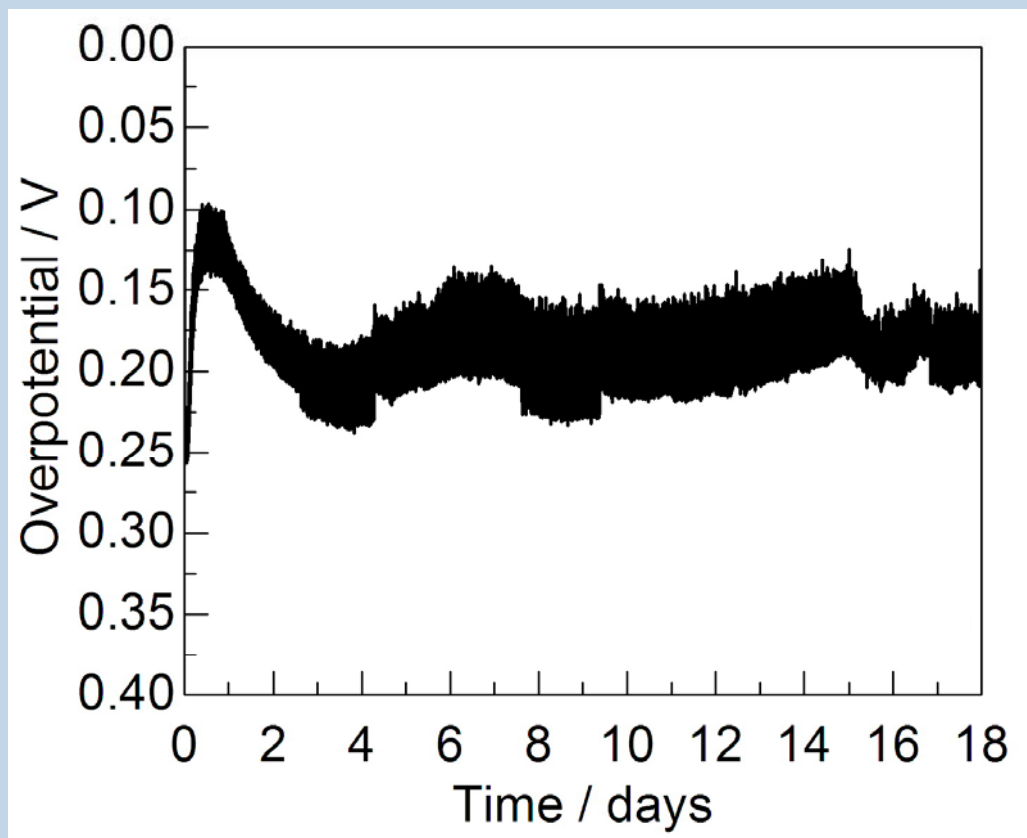
Esswein et. al., Submitted to *Energy and Environmental Science*



Engineered Co-OEC catalysts / conditions enable high resistance to poisoning

Earth-Abundant Cathode Catalyst

Cathode Stability



Sun Catalytix discovery -- cathode catalyst made of earth-abundant elements that exhibits stability in untreated river water

Full Cell Performance

- Initial results show progress toward ARPA-E goals (2V, 100 mA/cm²)
- Cells constructed using earth-abundant catalyst materials (anode is Co-OEC, cathode is proprietary)
- Ongoing efforts to improve performance by component engineering efforts

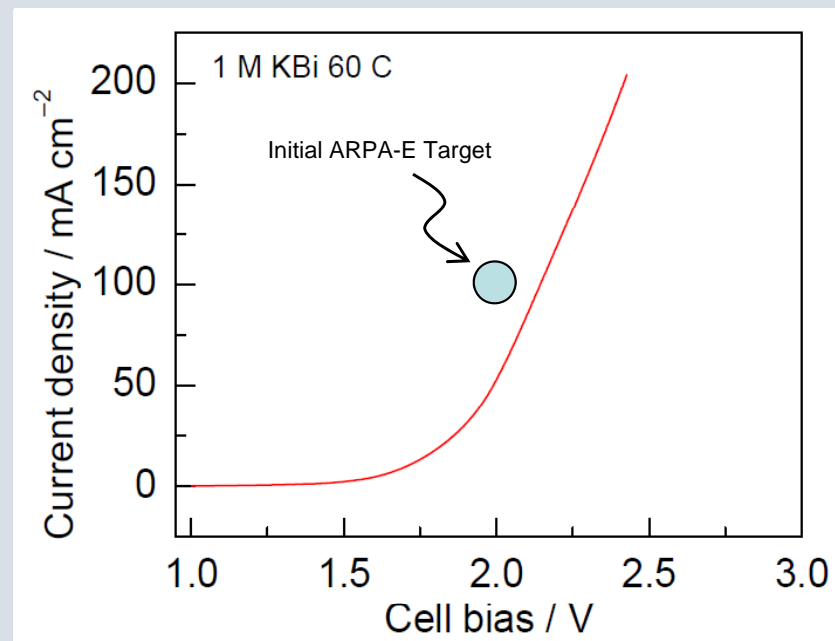


Photo-Electrochemical Cells

- Benign conditions suitable for Sun Catalytix catalysts increase the number of PV cells that can be used
- Operational current densities are limited by collecting solar photons
- Excellent match of system requirements with Sun Catalytix materials characteristics
- Several alternative configurations being explored

Photoanode/PV

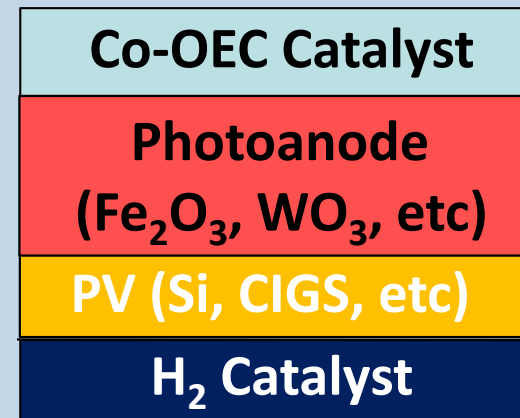
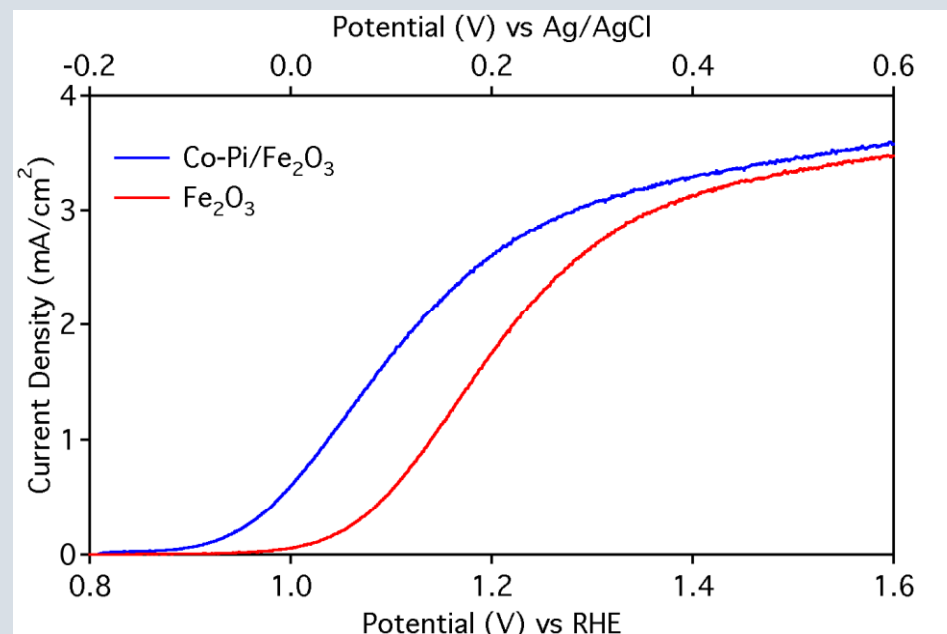
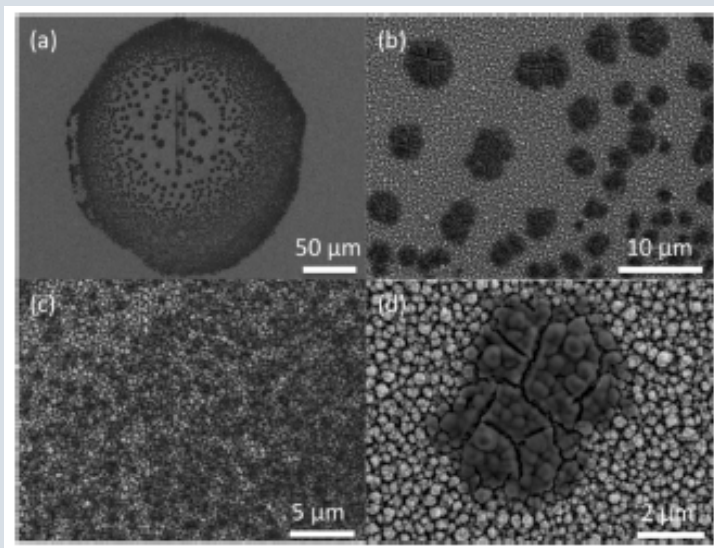


Photo-Anode Enhancement with Co-OEC



- Sun Catalytix catalysts shown to enhance photo current for Fe₂O₃
- Demonstrates proof of concept of enhanced activity via Co-OEC

Gamelin, D. R. and coworkers, Univ. of Wash.

Scientific and Strategic Leadership

Daniel Nocera
MIT
Founder



Art Goldstein
Ionics, Cabot, A123, MGH
Board Chairman



John Deutch
MIT, DOD, DOE, CIA, DARPA
SAB Chairman



Mark Wrighton
Washington U. – St Louis
SAB Member (Photo-catalytic chemistry)



Ken Smith
MIT
*SAB Member
(Chemical Engineering)*



George Whitesides
University Professor, Harvard
*SAB Member
(Materials science / chemistry)*



Henry White
University of Utah
SAB Member (Electrochemistry)



Management Team

Amir Nashat

Polaris Venture Partners, MIT

Chief Executive Officer



Tom Jarvi

United Technologies

Chief Technology Officer



Mark Barnett

CT Clean Energy Fund,
Foley Hoag Energy Technology

*VP Business Development and
General Counsel*

Company Backing

- Polaris Venture Partners
 - Leading VC firm; >\$3B assets under management
 - Founding investor; equity investments in series A and B rounds
- Tata
 - Global, visionary company and investor; \$70B group revenue
 - Equity investment in B round
- ARPA-E
 - \$4.1 M grant awarded in highly-competitive initial round



Contact:

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