



Bioinspired Composite Nanomaterials for Photocatalytic Hydrogen Production

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Pleotint LLC

DOE Project ID#: PDP36



Overview



Timeline

- Start - Oct. 2005
- End - Sept. 2009

Budget

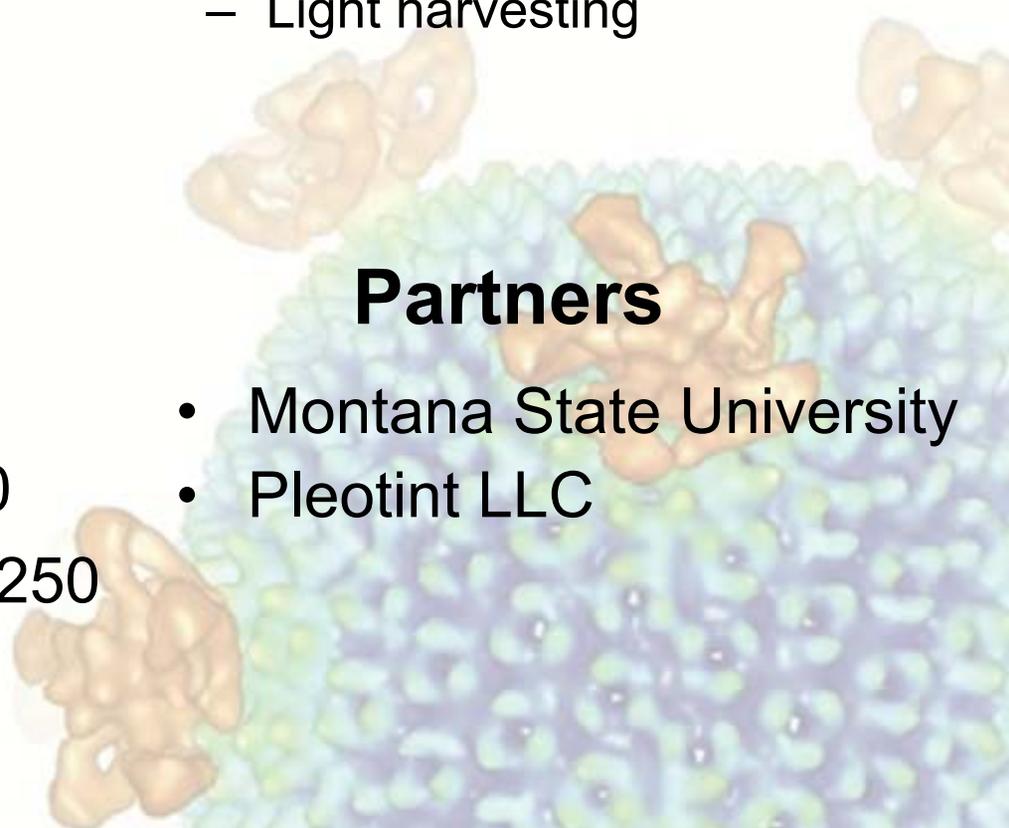
- Total project funding
 - \$1,491,250
 - DOE \$1,193,000
 - Contractor \$298,250

• Barriers addressed

- Enzyme stability/durability
- Oxygen sensitivity
- Light harvesting

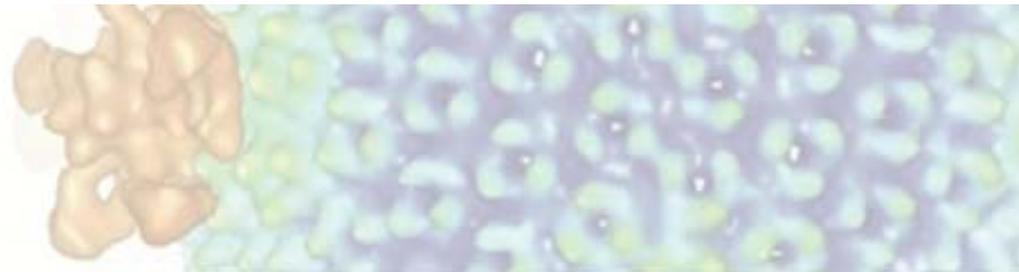
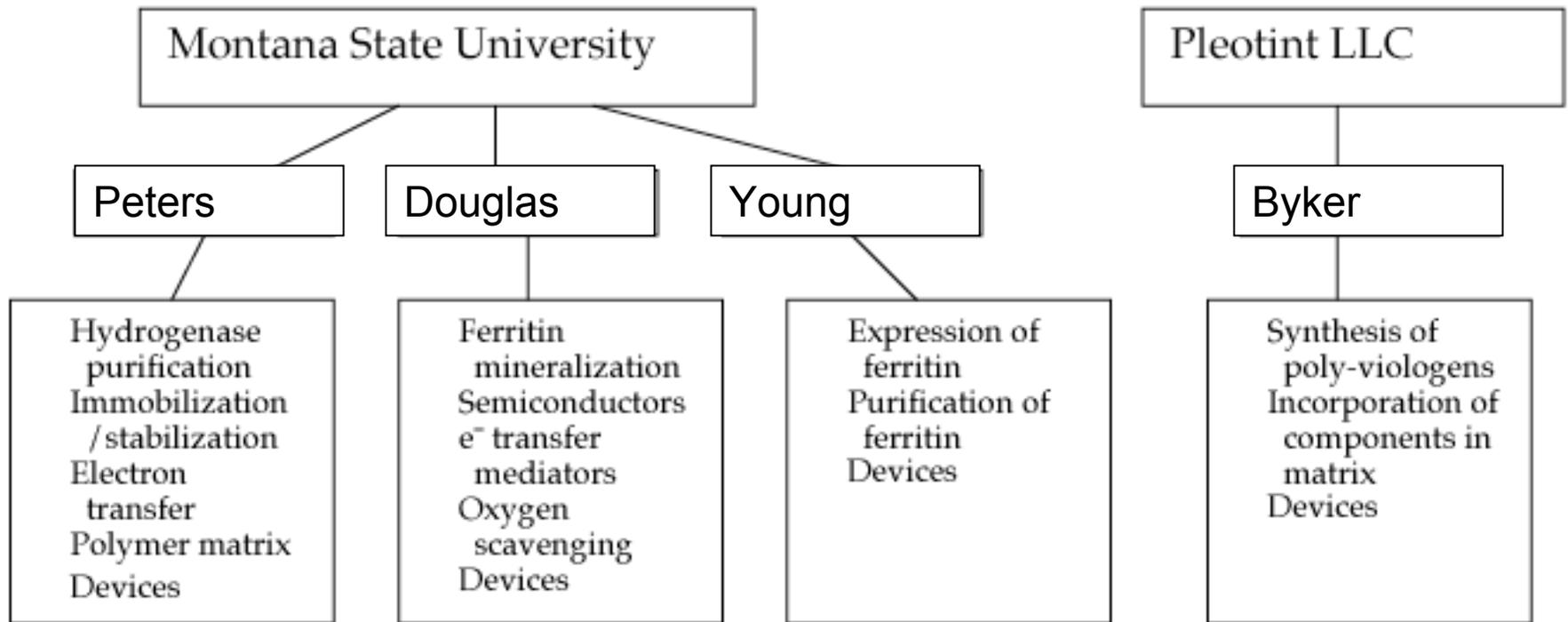
Partners

- Montana State University
- Pleotint LLC





Overall Project Structure





Objectives



1. Optimize the hydrogenase stability and electron transfer
2. Optimize the semiconductor nano-particle photocatalysis, oxygen scavenging, and electron transfer properties of protein nano-cages
3. Gel/Matrix immobilization and composite formulation of nano-materials and hydrogenase
4. Device fabrication for H₂ production

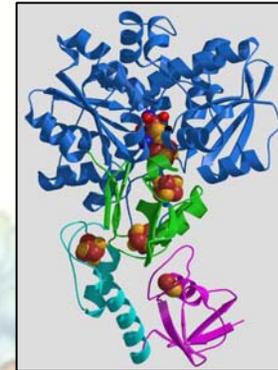


Approaches

Couple Different Catalyst Systems for Light Driven Hydrogen Generation

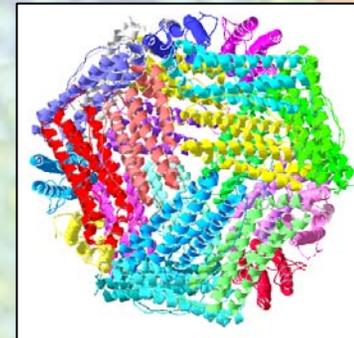
Biological catalysts (Hydrogenases)

- stabilization/immobilization
- electron transfer



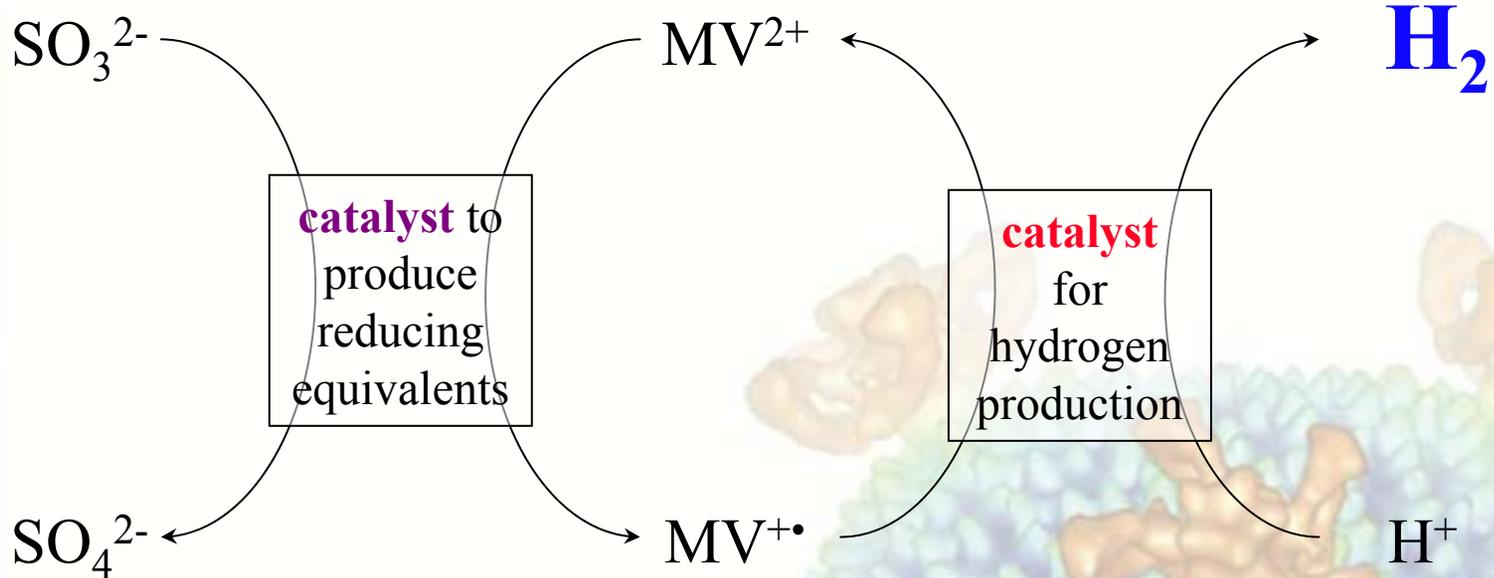
Nanoparticle Photocatalysts

- light harvesting
- O₂ scavenging

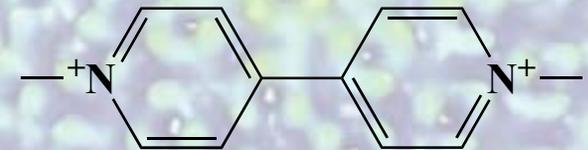




Hydrogen from Water: Coupled Reactions



GOAL: use a **catalyst** to reduce an electron mediator (methyl viologen, MV^{2+}) with SO_3^{2-} as electron donor. Another **catalyst** then uses $\text{MV}^{+\bullet}$ to produce H_2 .



methyl viologen, MV^{2+}



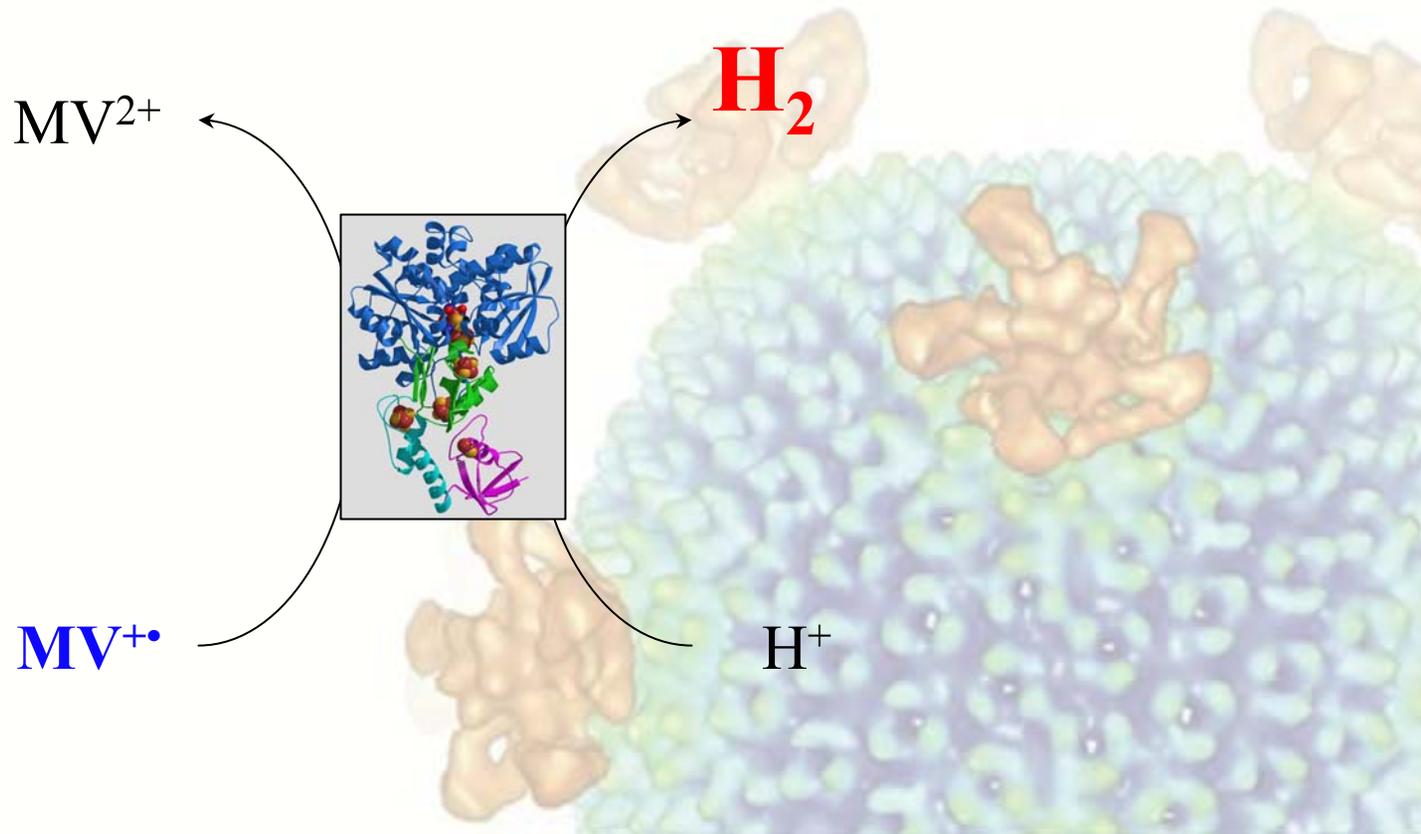
Enzymatic H₂ Formation



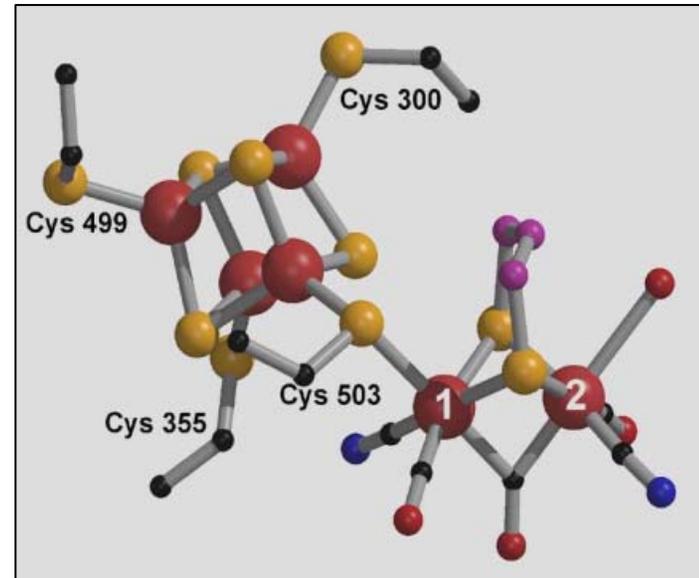
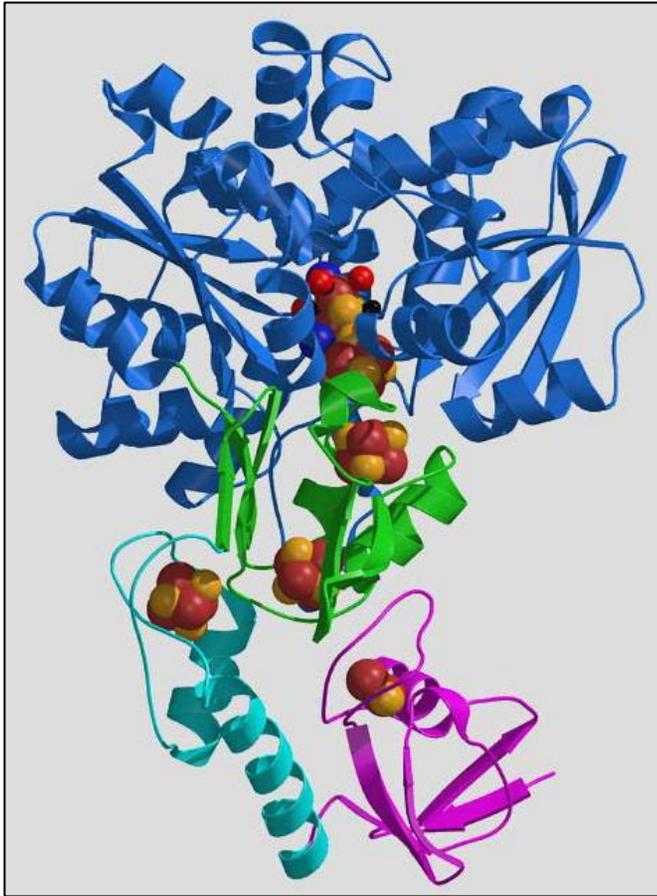
Hydrogenase enzymes

Highly active catalysts (9,000 H₂/enz/sec)

Utilize MV⁺ as reducing equivalents



Biological Hydrogen Production



Hydrogenase

Peters et al, *Science* (1998)



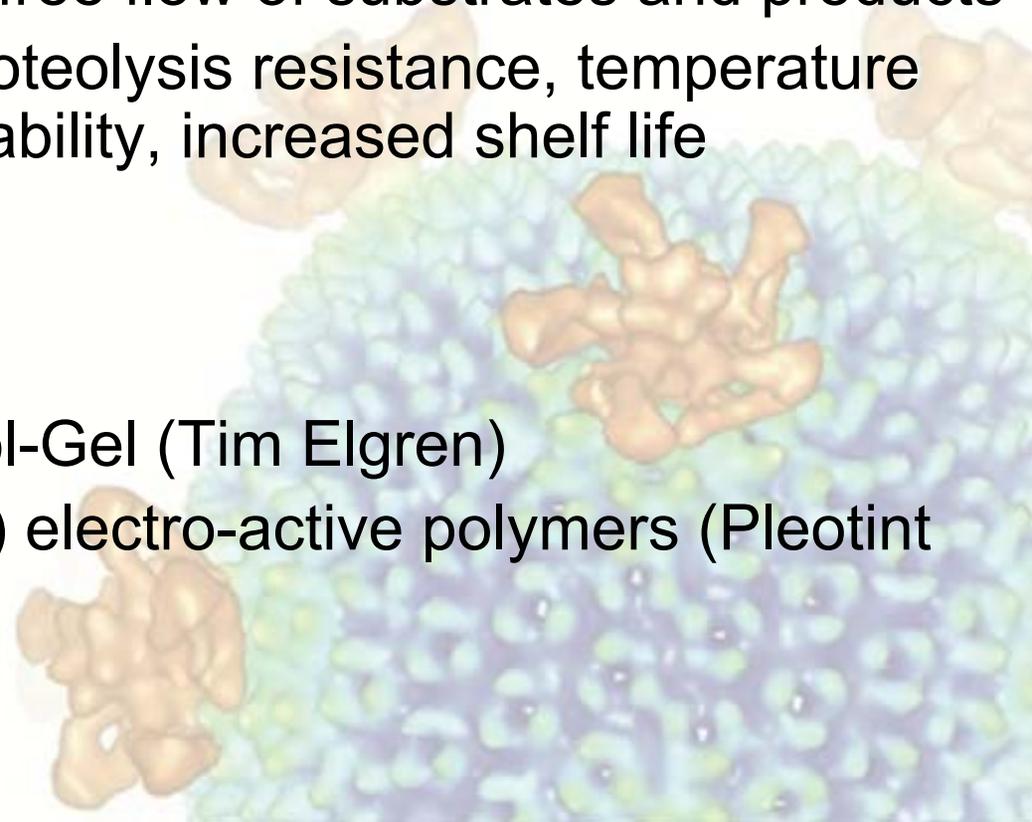
Hydrogenase Immobilization

Advantages

- Solid Phase – free flow of substrates and products
- Durability – Proteolysis resistance, temperature stability, pH stability, increased shelf life

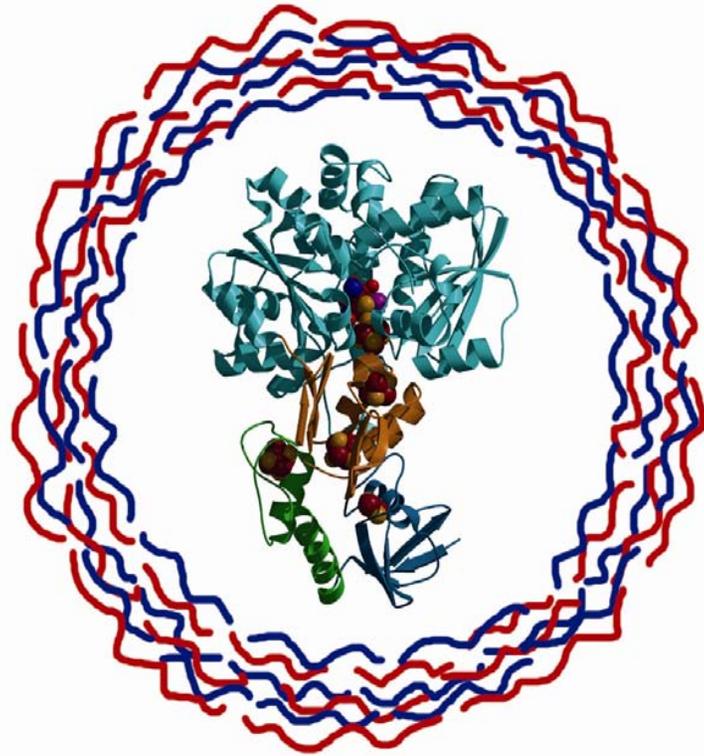
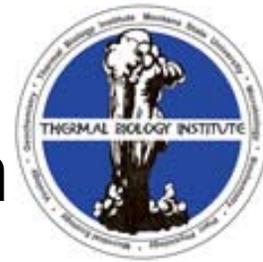
Approaches

- Silica oxide Sol-Gel (Tim Elgren)
- Poly(viologen) electro-active polymers (Pleotint LLC)

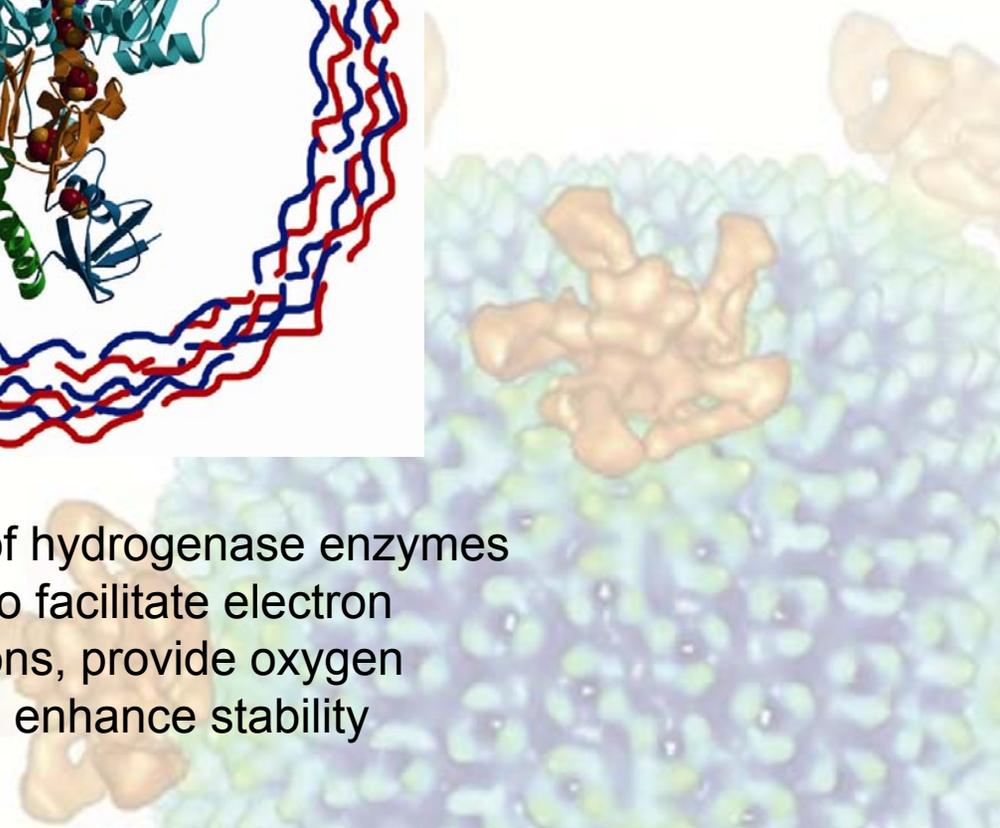




Encapsulation/Immobilization



Incorporation of hydrogenase enzymes into materials to facilitate electron transfer reactions, provide oxygen protection, and enhance stability



Procedure for making Sol-Gel hydrogenase materials



Prepared Sol-Gel mixture

1.57 ml Tetramethyl-ortho-silicate (TMOS)
350 μL H_2O
11 μL 0.04 M HCl

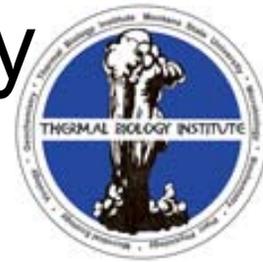
Sonicated solution for 30 min in cold bath
(with degassing)

Making Sol-Gel hydrogenase materials

*100 μL hydrogenase (100 μg of protein
in 50 mM Tris- HCl pH 8,0) :*
100 μL Sol-Gel mixture

Polymerization of Sol- Gel material for 3-5 min

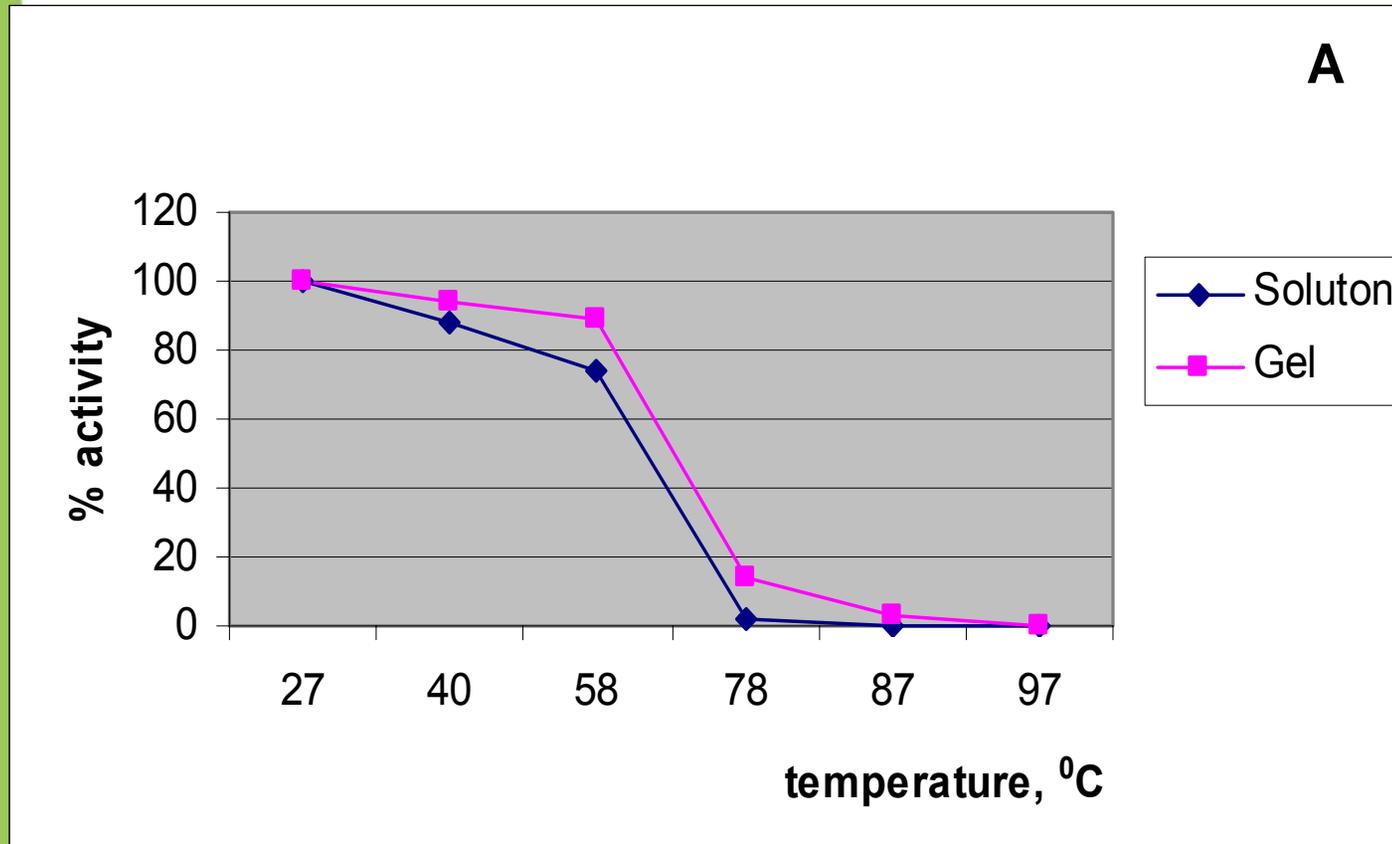
Recovery of hydrogenase activity encapsulated in Sol-Gel



Hydrogenase		% activity	
		Solution	Sol-Gel
1.	<i>Clostridium pasterianum</i>	100	63.8±15.8
2.	<i>Lamprobacter modestogalophilus</i>	100	67.5±8.8
3.	<i>Thiocapsa roseopersicina</i>	100	70.1±2.5



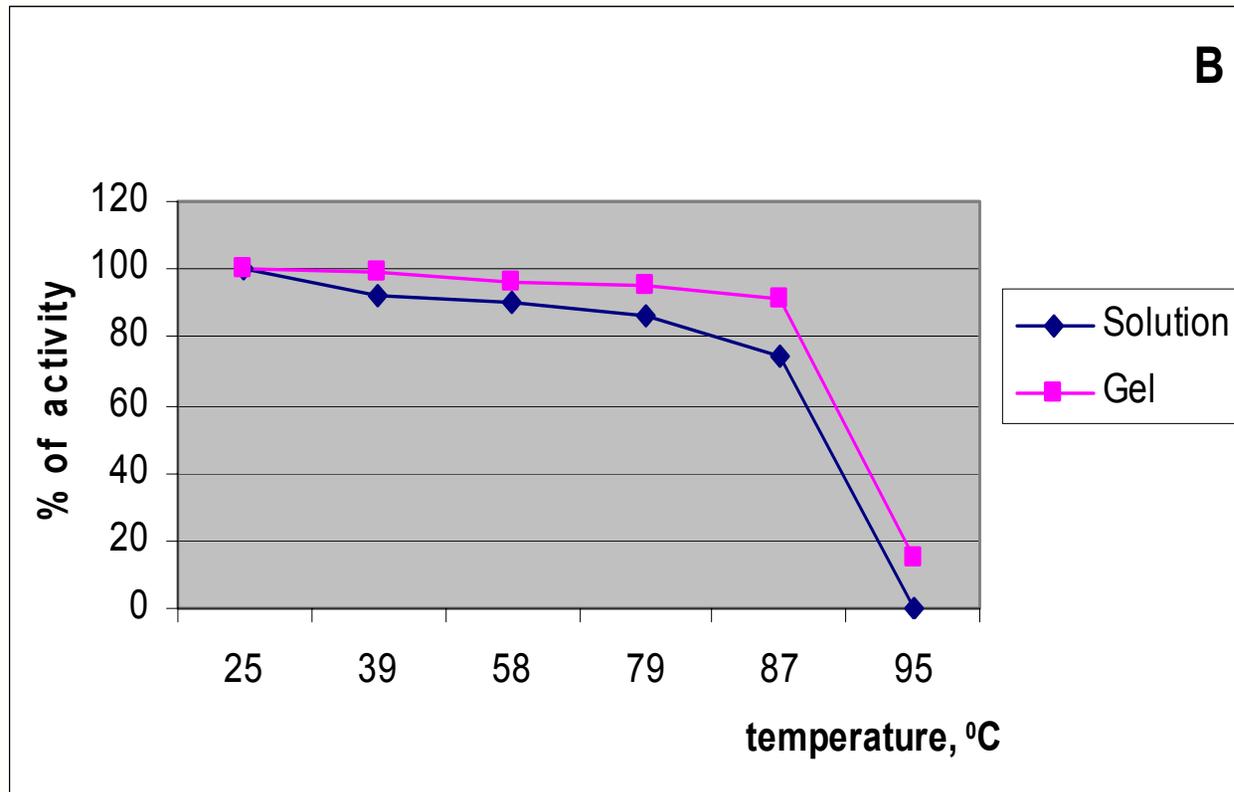
Thermal stability of hydrogenases encapsulated in Sol-Gel materials



A - Clostridium pasterianum hydrogenase



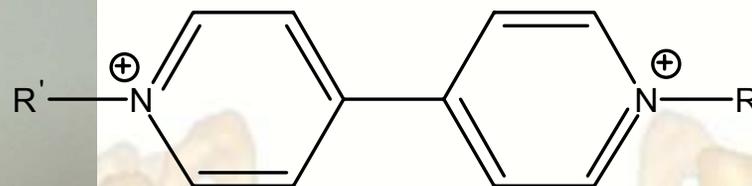
Thermal stability of hydrogenases encapsulated in Sol-Gel materials



B - Lamprobacter modestogalophilus hydrogenase

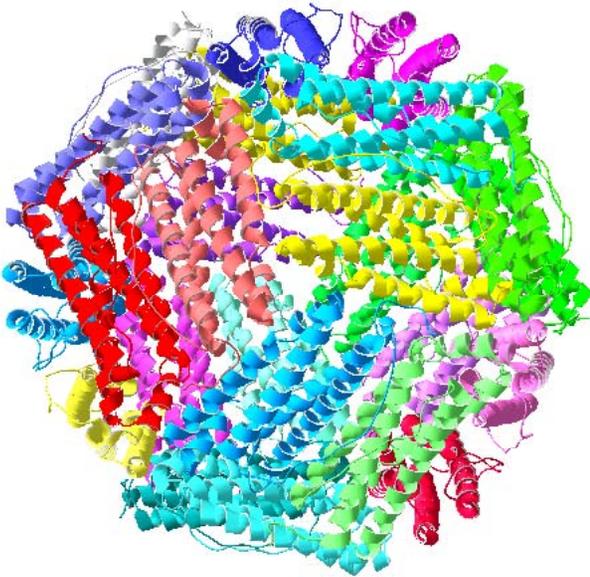


Association of Redox Mediator with Gel Matrix

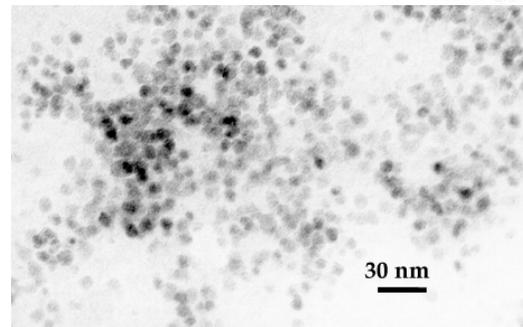
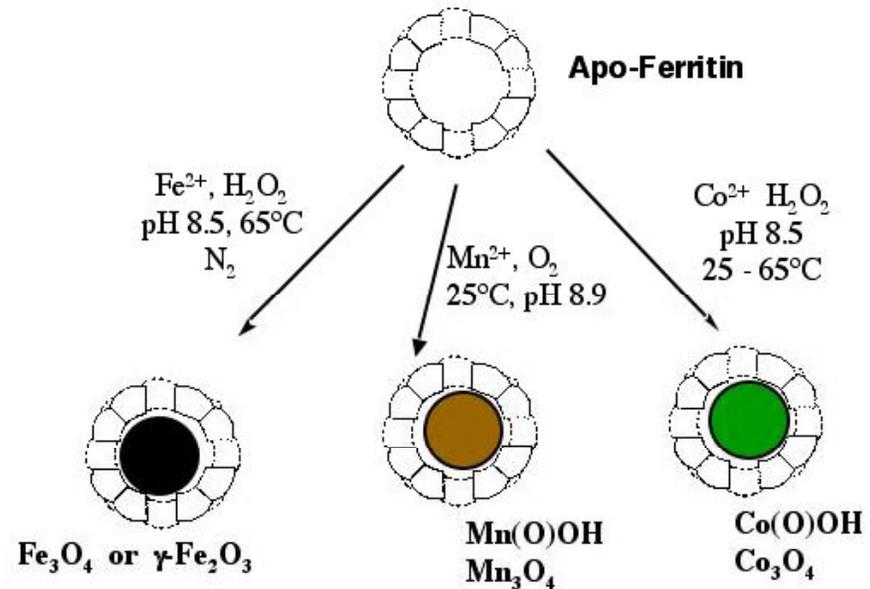


- $\text{R}, \text{R}' = \text{CH}_3$
 $\text{CH}_2\text{CH}_2\text{CH}_2\text{-SO}_3^-$
 $\text{CH}_2\text{CH}_2\text{CO}_2^-$
 $\text{CH}_2\text{CH}_2\text{NH}_3^+$
 $(\text{CH}_2)_n\text{-PO}_3\text{H}_2$

Nanoparticle synthesis within the Ferritin Protein Cage

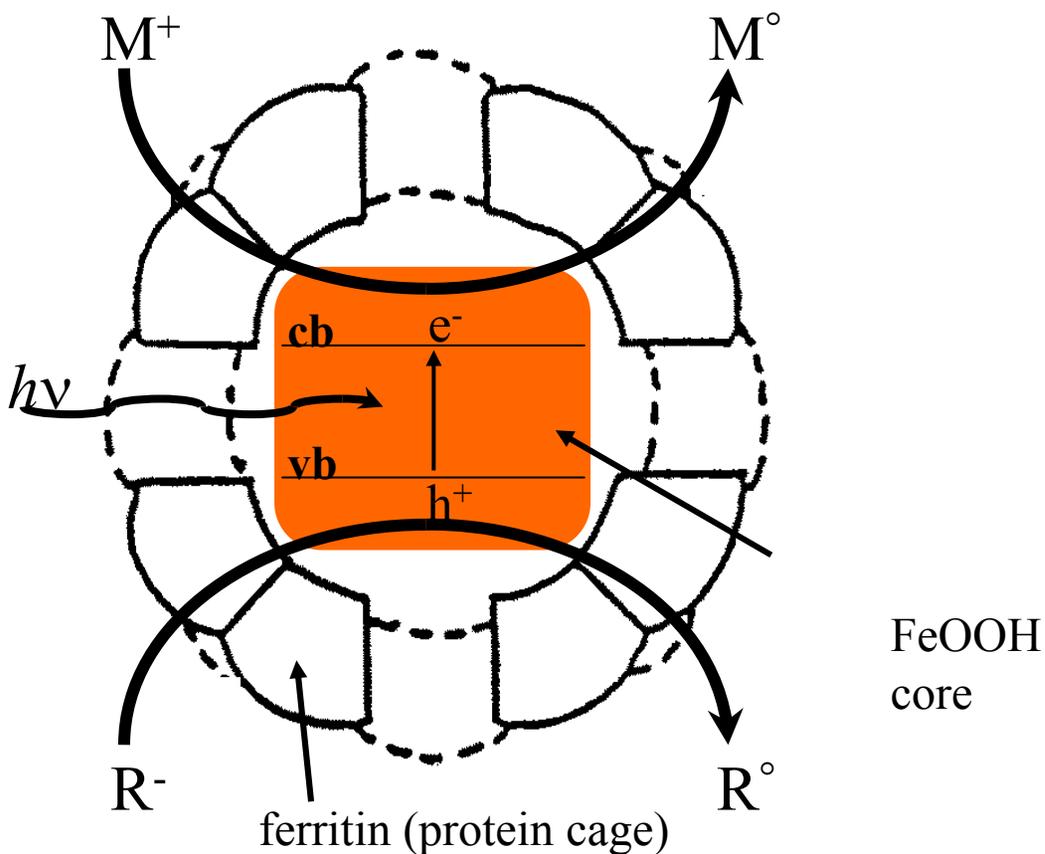


Ferritin protein cage
24 subunits - 12 nm diam



TEM of metal oxide
nanoparticles

Protein Cage Photocatalysts



Light absorption by ferritin core (FeOOH) causes charge separation oxidizes R^- and reduces M^+ catalytically.

Examples:

- Reduction of CrO_4^{2-} to Cr(III) using tartrate as electron donor (Kim et al., *Chem. Mater.*, 2002).
- Reduction of Cu(II) to Cu(0) particles using citrate as electron donor (Ensign et al., *Inor. Chem.*, 2004).

Current: use this photocatalytic system (or an analogue) to reduce MV^{2+} to MV^{+} using sulfite as electron donor.



Thermodynamics and Kinetics

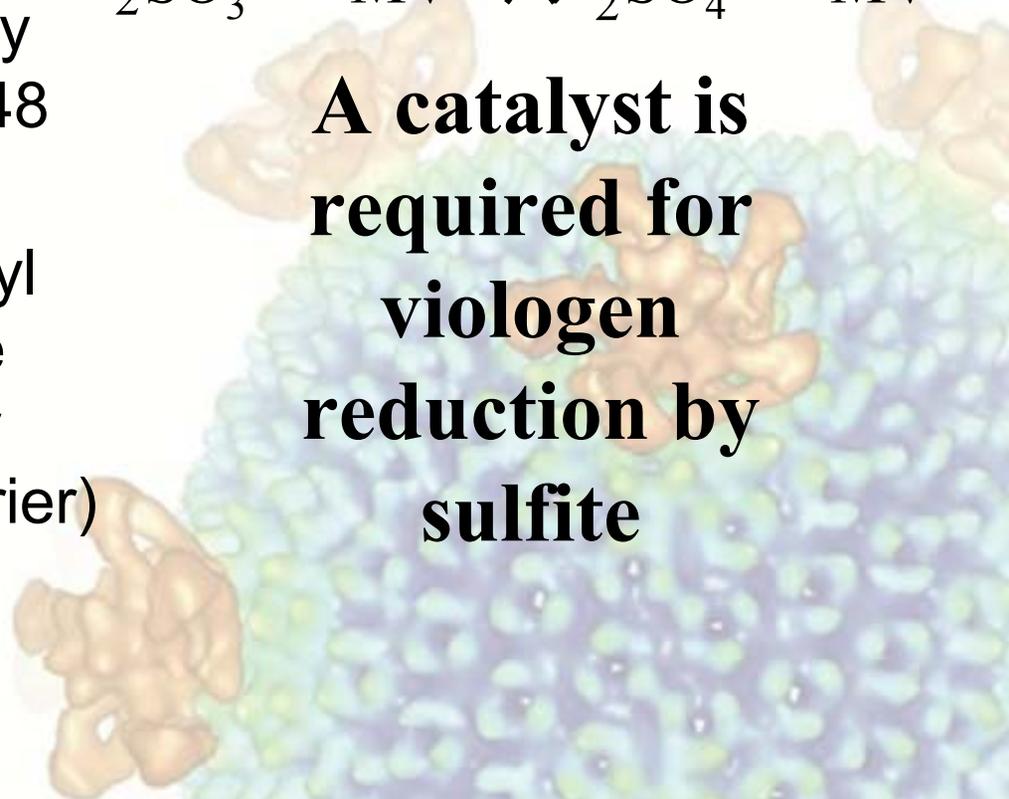


- electron transfer from sulfite to methyl viologen is thermodynamically favorable, $\Delta G = -48$ kJ/mol
- reduction of methyl viologen by sulfite does not normally occur (kinetic barrier)

viologen reduction

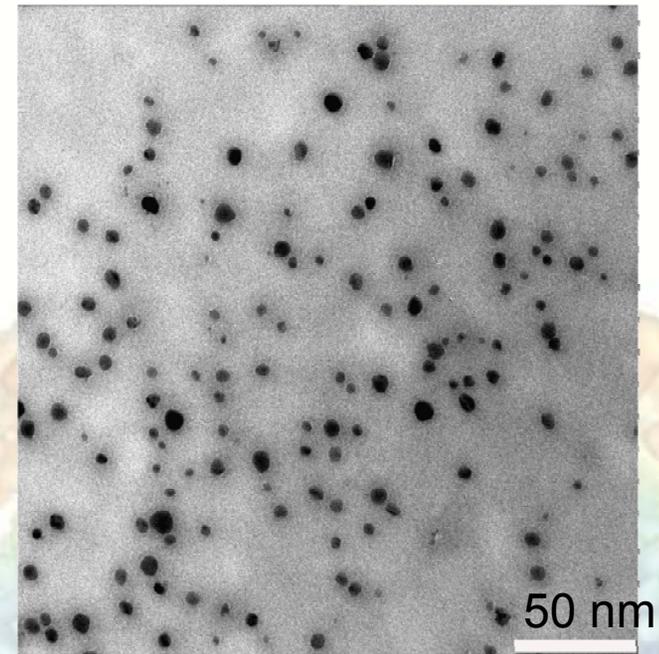
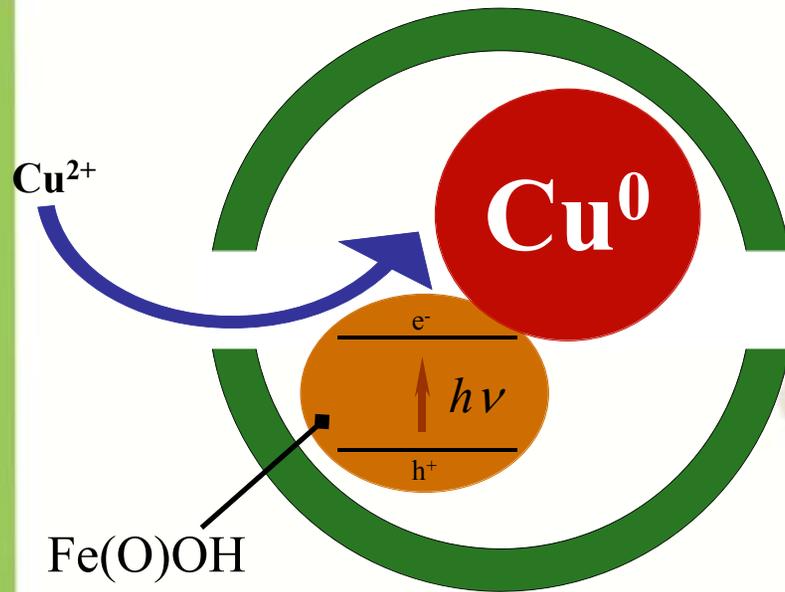


A catalyst is required for viologen reduction by sulfite





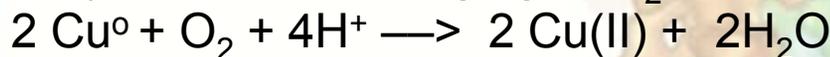
Photoreduction of Cu(II)



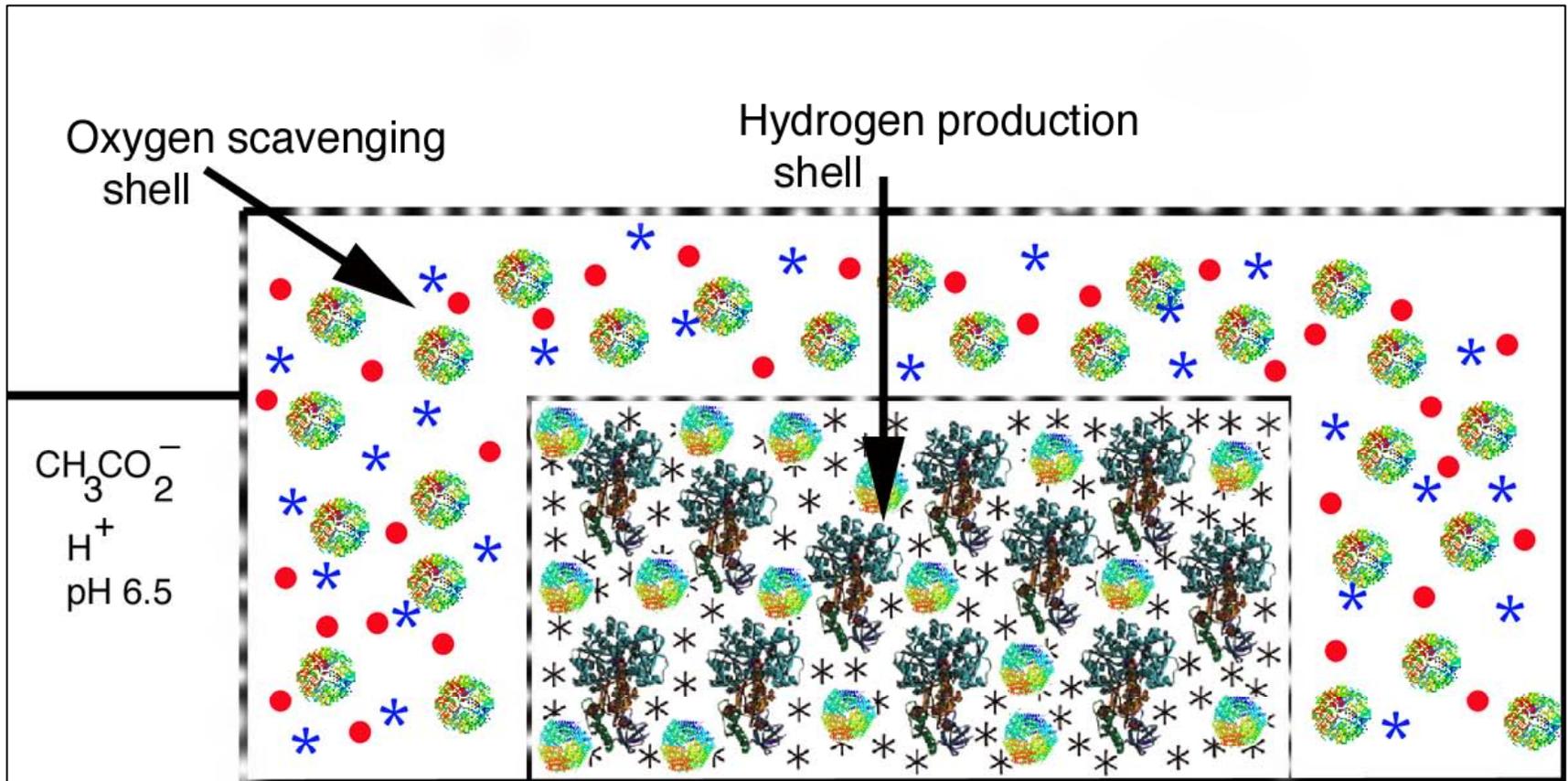
TEM of Ferritin encapsulated Cu nanoparticles

Photoreduction of Cu(II) to form protein encapsulated Cu⁰ nanoparticles

Very efficient scavenging of O₂ from the media



Long-Term Goal – Device for photocatalytic hydrogen production – composite materials (nanoparticles and hydrogenase enzymes)





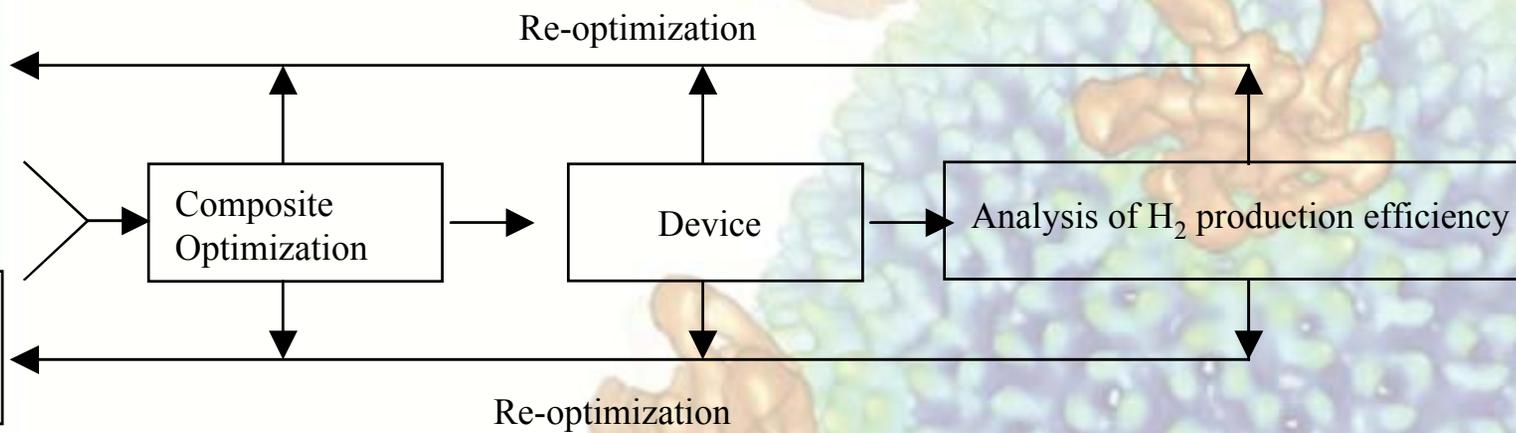
Future Goals



Development of device prototype
demonstration of light harvesting

Hydrogenase
Optimization

Semiconductor
Nanoparticle
Optimization

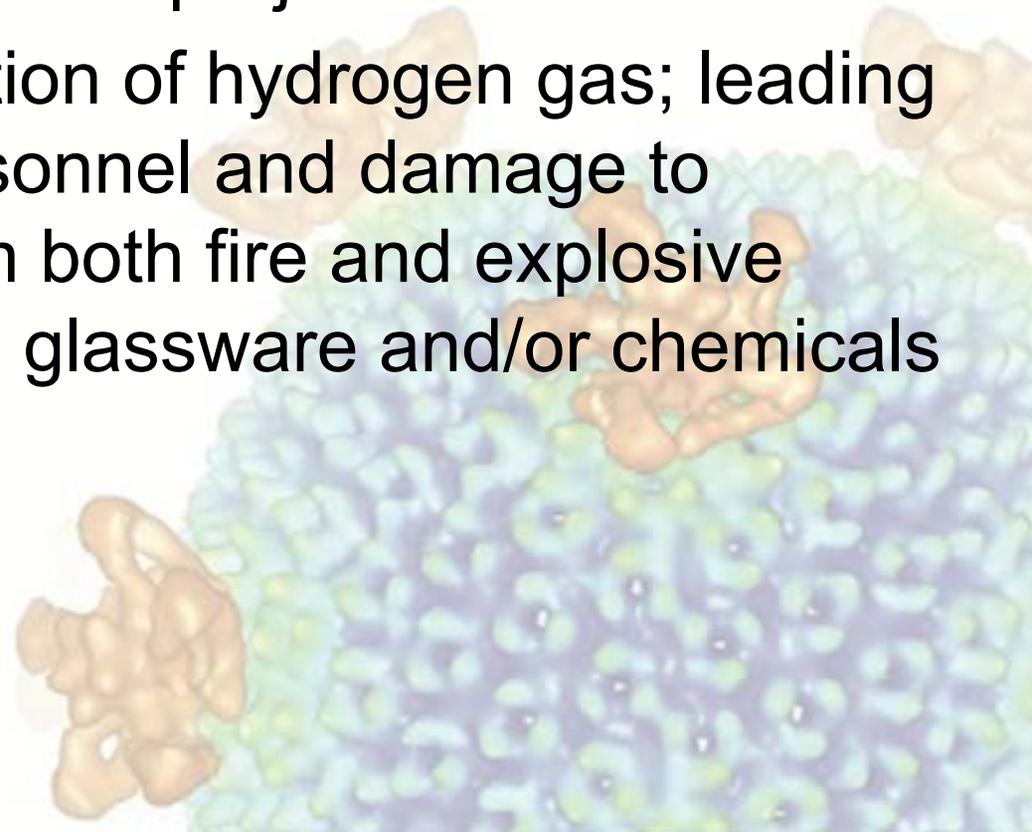




Hydrogen Safety

The most significant hydrogen hazard associated with this project is:

Accidental ignition of hydrogen gas; leading to injury of personnel and damage to equipment from both fire and explosive debris such as: glassware and/or chemicals





Hydrogen Safety

Our approach to deal with this hazard is:

Follow lab protocol of wearing safety glasses, gloves

Keep glove box H_2 level below 3%

Vent gases in fume hood

Keep away from open flame and flammable chemicals

Keep quantity of H_2 production to a minimum

In event of accidental explosion contact

Jeff Shada, Safety and Risk Management,
Advanced Tech Park, 406-994-2711