

0.0 0.2 0.4 0.6 0.8 1.0 1.2 1.4 1.6

(d) 1 M NaOH, 0.02 M HMF

Potential vs RHE

0.4 0.6 0.8 1.0 1.2 1.4 1.6 1.8

Potential (V vs RHE)

# **Novel Approach to the Synthesis of Supported Gold Nanoparticles for HMF Oxidation**



Potential (V vs RHE)

Potential (vs RHE)

—— Au

1.2

0.8

(MA)

— Au/X

0.4 0.6

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# **Step 1: Gold Oxidation** Potentiostat | RE (Hg/HgO | - - <del>]</del> - | I\_ \_ \_ \_ I 2.0 M KOH

# **Results and Discussion**



References





## Figure 5: Cyclic Voltammetry (CV) Scans of Au/C, Au, and Au/X vs. RHE

Figure 5a: CV scan of Au/C (Au NPs on carbon) on fluoride tin-oxide (FTO) glass, revealing first observation of unique peak during the reverse

• Figure 5b and 5c: CV scans of the Au wire in alkaline media without HMF (5b), and with HMF (5c). A peak is also observed only in the reverse scan around 1.1V vs RHE, where oxidized gold

• Figure 5d: CV scan of the Au wire with HMF conducted below the potential range for gold oxide formation, with no reverse peak observed. Figure 5e-f: Enhanced catalytic activity observed for Au/X in HMF oxidation (5e) and ethanol



- investigation.

## **Conclusion and Future Work**

. David and Lorraine Freed Undergraduate Research Symposium, Lehigh University 2. Clare Boothe Luce Research Scholars Program







Figure 4: SEM and HR-TEM Images of Au Surfaces

• Figure 4a-b: SEM images showing the surfaces of the oxidized gold wire and the synthesized Au/X with CV treatment, respectively.

Figure 4c: SEM image of the Au/X wire without CV, highlighting the less uniformly distributed and rough, porous structure compared to **4b** with CV.

Figure 4d-e: HR-TEM images of the Au/X wire that display the formation of Au nanoparticles (Au NPs). The nature of surrounding film/layer (X) is under

Au/X layer significantly enhances oxidation processes in HMF and other reactants like EtOH, demonstrating the efficacy of gold as an electrocatalyst.

A simple and reproducible method was established for synthesizing Au nanoparticles.

Future work will focus on detailed characterization of the catalyst layer and its underlying mechanism by advanced spectroscopy and electrochemical analysis.

# Acknowledgements

