

# Portland Section Meeting Notice

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## Electrifying the Chemical Industry

a talk by

### Prof. Paul Kempler

Research Assistant Professor, Chemistry and Biochemistry  
Associate Director, Oregon Center for Electrochemistry  
Director of the Electrochemistry Master's Internship Program (EMIP)  
at University of Oregon

**Thursday January 9, 2025, 6:30-9 PM**

Our meeting day returns to Thursday. This meeting features a catered dinner from a local Portland eatery, a short talk by University of Oregon electrochemistry masters student Andrew Goldman, and a keynote lecture from Dr. Kempler. This event is all ages welcome and encouraged.

## Dinner Reservations

**Venue:** Helioterra Winery  
2025 SE 7th Ave.  
Portland, OR 97214

**Schedule: Doors open at 6:30pm on Wednesday December 11th.**

**Helioterra Winery** is a women-owned and operated small scale winery.  
Wines are available from Helioterra for purchase,  
as well as beer and non-alcoholic beverages.

**Prof. Kempler Bio/Abstract next page.**  
**also, Jonathan Xu Abstract and graphic from December 11 ACS meeting presentation.**

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## Electrifying the Chemical Industry



Paul Kempler

**Talk Summary:** Electrochemistry is well-positioned to connect inexpensive and clean electricity to the production of commodity chemicals and materials. In this talk, I will talk about career opportunities for students with degrees in electrochemistry based on the first 75 graduates from the Electrochemistry Masters Internship Program at the University of Oregon. Next, I will present recent research results from my group related to the production of green hydrogen and the reduction of iron ore to metal for green steel.

**Speaker Bio:** Paul Kempler is an Assistant Professor of Chemistry and Biochemistry and the Director of the Electrochemistry Master's Internship Program (EMIP). His current research efforts include studying interfacial ion-transport processes and developing electrochemical cells for producing iron for emissions-free steelmaking. He received the UO Sustainability Award for Research and Scholarship in 2023.

## DpdC in *Salmonella* Montevideo: Potentially Novel Catalytic Mechanism for Nitrile Hydratase Activity

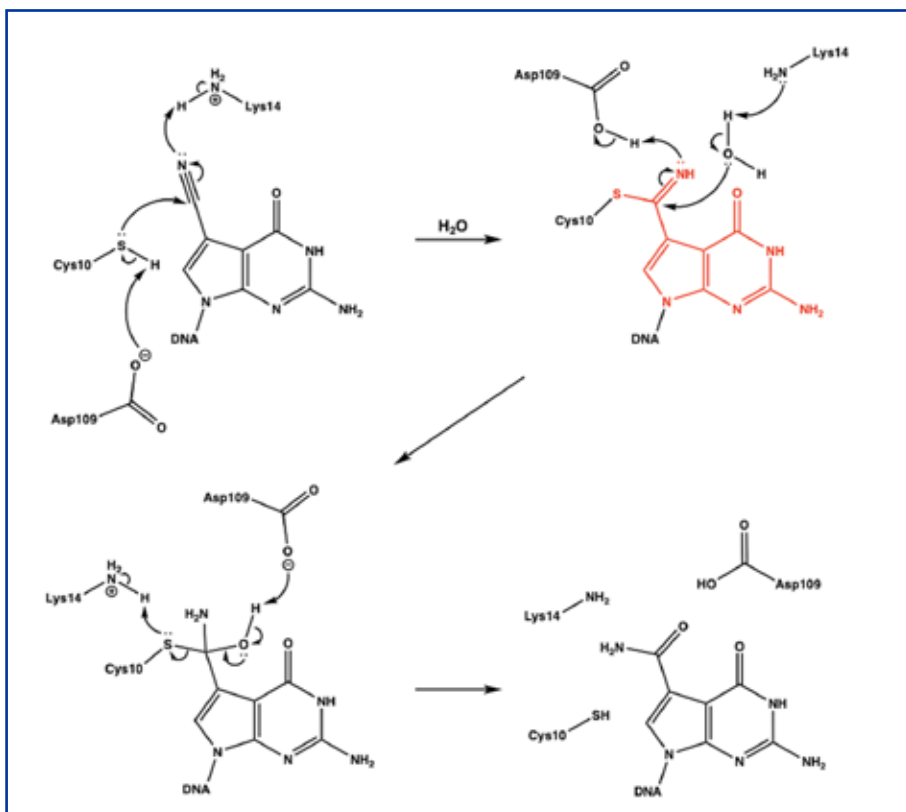
Jonathan Xu, Andrew Buckley, Dirk Iwata-Reuyl

*Ed: Jonathan Xu presented findings from his research during the December 11 ACS meeting. He provides this Abstract and graphic of his project.*



Jonathan Xu (Photo Dibblee).

Chemical modification of nucleotides in DNA is vital to restriction-modification (RM) in bacterial and phage systems. While this typically involves the methylation of bases in recognition sequences, recently we reported the presence of structurally diverse 7-substituted-7-deazaguanines in bacterial and phage DNA that constitute previously unknown RM systems. These studies established the formation of one such modified nucleoside, 2'-deoxy-7-amido-7-deazaguanosine (dADG), as the hydrolysis product of 7-cyano-7-deazaguanine (preQ<sub>0</sub>) modified DNA by the enzyme DpdC. Notably, the enzyme's lack of active site metal-binding sequences characteristic of nitrile hydratases suggests that DpdC follows an alternative, potentially novel mechanistic pathway. Here we propose the nucleophilic formation of a thioimide intermediate preceding hydrolysis involving a conserved cysteine residue (Cys<sup>10</sup>) in the putative active site of DpdC enzymes, and demonstrate significantly reduced dADG synthesis following mutagenesis of Cys<sup>10</sup> to Ser<sup>10</sup>. This attenuation in activity is consistent with the residue's involvement in catalysis, warranting further mechanistic investigation.



Proposed DpdC Catalytic Mechanism.png