

Guest Colloquium of the CRC 1452 CLINT ~ Catalysis at Liquid Interfaces



Single-Atom Alloy Catalysts: Born in a Vacuum, Tested in Reactors, and Understood In Silico

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09 December 2021 15:30 Zoom Online Meeting



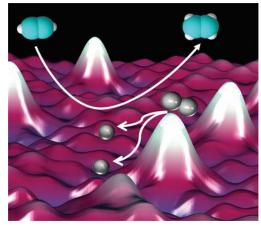


Single-Atom Alloy Catalysts: Born in a Vacuum, Tested in Reactors, and Understood In Silico

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In this talk I will discuss a new class of heterogeneous catalysts called Single-Atom Alloys in which precious, reactive metals are utilized at the ultimate limit of efficiency. 1-6 These catalysts were discovered by combining atomic-scale scanning probes with more traditional approaches to study surface-catalyzed chemical reactions. This research provided links between atomic-scale surface structure reactivity which are key to understanding and ultimately controlling important catalytic processes. In collaboration with Maria Flytzani-Stephanopoulos these concepts derived from our surface science and theoretical calculations have been used to design Single-Atom Alloy nanoparticle catalysts that are shown to perform industrially relevant reactions at realistic reaction conditions. For example, alloying



Scanning tunneling microscope (STM) image showing atomically-dispersed palladium atoms in a copper surface. The palladium atoms activate hydrogen enabling the industrially important acetylene-ethylene conversion with 100% selectivity.

elements like platinum and palladium with cheaper, less reactive host metals like copper enables 1) dramatic cost savings in catalyst manufacture, 2) more selective hydrogenation and dehydrogenation reactions, 3) reduced susceptibility to CO poisoning, and 4) higher resistance to deactivation by coking. I go on to describe very recent theory work by collaborators Stamatakis (UCL) and Michaelides (Cambridge University) that predicts reactivity trends for a wide range of *Single-Atom Alloy* combinations for important reaction steps like H-H, C-H, N-H, O-H, and CO₂ activation. Overall, I hope to highlight that this combined surface science, theoretical, and catalyst synthesis and testing approach provides a new and somewhat general method for the a priori design of new heterogeneous catalysts.

References:

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- [2] Marcinkowski et al. Nature Materials 12, 523 (2013).
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- [4] Liu et al. JACS 138, 6396 (2016).
- [5] Marcinkowski et al. Nature Chemistry 10, 325 (2018).
- [6] Hannagan et al. Science 372, 1444 (2021).