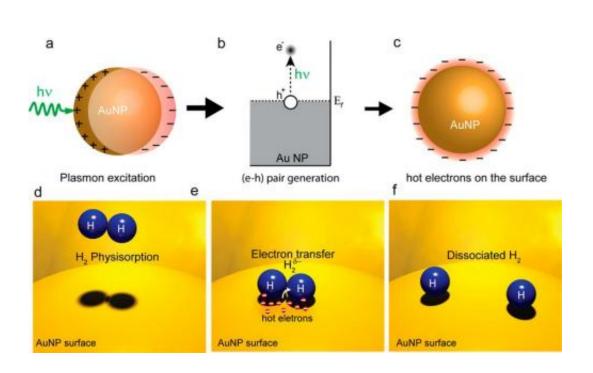


December 17 2012, by Lisa Zyga

Hot electrons do the impossible in catalytic chemistry



In (a), (b), and (c), surface plasmons excited by a laser generate hot electrons on a gold nanoparticle surface. In (d), (e), and (f), hot electrons transfer to H2 molecules and cause them to dissociate. Since dissociation is the rate-limiting step in many chemical reactions, this hot electron process can increase the reaction efficiency. Image credit: Shaunak Mukherjee, et al. ©2012 American Chemical Society

(Phys.org)—From petroleum refining to food processing, the vast majority of commercial chemical applications involve catalysts to control the rate of chemical reactions. Anything that can increase the



efficiency of catalysts could have a widespread impact in these areas. In a new study, scientists have shown that an efficiency improvement may be possible by creating "hot" electrons (those with a very high energy) from a metallic surface on which the chemical reaction occurs, and using the electrons to control the chemical reaction and make it more efficient.

The researchers, led by Professors Peter Nordlander and Naomi J. Halas of Rice University in Houston, Texas, and Professor Emily A. Carter of Princeton University, New Jersey, have published their study on the technique to improve catalytic chemistry in a recent issue of <u>Nano Letters</u>.

In their method, the researchers used a visible-light laser to illuminate a gold nanoparticle surface (the catalyst surface) at room temperature. The incoming laser light optically excites <u>surface plasmons</u> on the <u>metal</u> <u>surface</u>, and the plasmons then decay into <u>hot electrons</u>. Because of their high energies, the hot electrons extend further away from the nanoparticles than electrons with lower energies do. If another atom or molecule that can accept the electron is nearby, the hot electron can transfer into that acceptor's electronic states.

In these experiments, the researchers adsorbed H_2 molecules on the gold nanoparticle surface, a procedure that is commonly performed in <u>heterogeneous catalysis</u>, in which the adsorbed molecules act as <u>reactants</u>. The researchers found, as the main result of their study, that some of the hot electrons could transfer into the closed shells of the H_2 molecules and cause the two <u>hydrogen atoms</u> to separate, or dissociate. This process, called "plasmon-induced dissociation of H_2 on Au," could improve the efficiency of certain <u>chemical reactions</u>.

"In many important chemical reactions, the rate limiting step is the dissociation of closed shell molecules like a hydrogen molecule," Nordlander told *Phys.org.* "If this reaction step can be 'dealt with' (using



our demonstrated hot electron process), the reaction may proceed much faster and require less energy."

To measure the dissociation, the researchers exposed the nanoparticle substrate to both H_2 and D_2 (two atoms of deuterium, a hydrogen isotope containing one neutron). If the hot electrons dissociated these molecules, it would result in the formation of HD molecules. The researchers found that, as soon as they turned the laser on, the rate of HD formation on the nanoparticle surface increased by a factor of 6. They also measured that the rate was strongly dependent on the concentration and size of the gold nanoparticles. The researchers explained that, in a sense, the electrons "do the impossible" because there would be no dissociation without them.

"A hydrogen molecule adsorbed on a macroscopic gold surface does not dissociate spontaneously, even at elevated temperatures," Nordlander said. "It is an impossible chemical reaction. The energy for dissociation is simply too large. We picked the hydrogen-gold system exactly for this reason. We wanted a system where the interpretation of the effect was straightforward. On other metals, like transition metals, a hydrogen molecule can dissociate spontaneously, in particular near defects and at elevated temperatures. But not on gold."

These results demonstrate how the field of plasmonics – which involves interactions between photons and nanostructures – can have applications in the field of catalytic chemistry, particularly photocatalysis, The researchers hope that this demonstration of the ability to control chemical reactions using light could be extended for specific uses. By using the laser frequency to tune the surface plasmon resonances, it may be possible to create hot electrons with specific energies that populate specific electronic states of molecules adsorbed on the nanoparticle <u>surface</u>.



"The immediate application is to improve the efficiency of metallic catalysts by illuminating them with light," Nordlander said. "Our groups have a very 'green agenda' and our intent is to design novel types of catalysts that are driven by sunlight. This is something we have an abundance of here in Houston.

"Another important application is to investigate to what extent we can steer and control chemical reactions by having hot electrons of particular energies transferring into specific molecular orbitals. This is a relatively unexplored research topic. Normal electrons have positive energies and are much 'hotter' that our hot electrons. Such high-energy electrons typically do not stay on a molecule sufficiently long for the atoms to start to move."

More information: Shaunak Mukherjee, et al. "Hot Electrons Do the Impossible: Plasmon-Induced Dissociation of H_2 on Au." *Nano Letters*. DOI: 10.1021/nl303940z

Copyright 2012 Phys.org All rights reserved. This material may not be published, broadcast, rewritten or redistributed in whole or part without the express written permission of Phys.org.

Citation: Hot electrons do the impossible in catalytic chemistry (2012, December 17) retrieved 26 April 2024 from https://phys.org/news/2012-12-hot-electrons-impossible-catalytic-chemistry.html

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.