

Metal hydrides with improved stability in water

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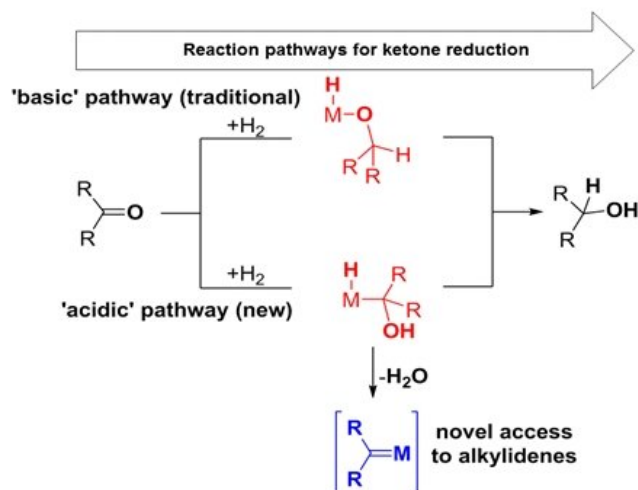


Figure shows the "basic" and "acidic" metal hydride pathways for the ketone reduction reaction. Alternative reaction pathways for known reactions add to the understanding of catalytic systems, but also offer the chance to access new reaction pathways. Credit: National University of Singapore

NUS chemists have discovered new reaction pathways using "acidic" metal hydrides to access industrially important chemical complexes.

Metal hydrides are compounds with the metal atoms bonded to hydrogen atoms, and are important for various catalytic and synthetic reactions. Mechanisms for the [reaction](#) pathways using traditional "basic" metal hydrides are well understood. However, their "basic" metal-hydrogen bond polarity causes them to react vigorously with water, often in an irreversible manner. This is undesirable for many industrial chemical reactions which are performed in aqueous solutions, or under "wet" atmospheric conditions, as a significant portion of the metal [hydride](#) catalyst will react with water molecules instead of the intended reactants. This constraint limits the use of metal

hydrides in many industrial processes. Prof Rowan YOUNG and his research team from the Department of Chemistry, NUS have developed new "acidic" metal hydrides with an inverse metal-hydrogen bond polarity that allows them to be more stable in the presence of water. This important achievement has opened up avenues to new synthetic pathways that cannot readily be accessed via conventional metal hydrides.

For example, using the newly developed "acidic" metal hydrides, the research group discovered reaction pathways to important organometallic "pincer" ligand frameworks. These "pincer" complexes are of fundamental importance to modern catalysis and are an emerging class of metal complexes that have been very successful at promoting difficult bond activations and unique catalytic reactions, but are often difficult to access through traditional synthetic methods.

The Young group also exploited the "acidic" [metal](#) hydrides to map alternative reaction pathways (to previously understood mechanisms) for the hydrogenation of ketones. The hydrogenation of ketones is a key industrial reaction carried out by many chemical companies on a large scale throughout the world.



Photo showing the research team carrying out chemical experiments using fume hoods in the laboratory. Credit: National University of Singapore

More information: Simon Sung et al. Convergent (De)Hydrogenative Pathways via a Rhodium η -Hydroxylalkyl Complex, *Organometallics* (2017). DOI: [10.1021/acs.organomet.7b00158](https://doi.org/10.1021/acs.organomet.7b00158)

Simon Sung et al. Protonolysis of an η -Hydroxyl Ligand for the Generation of a PCcarbeneP Pincer Complex and Subsequent Reactivity Studies, *Organometallics* (2017). DOI: [10.1021/acs.organomet.7b00461](https://doi.org/10.1021/acs.organomet.7b00461)

Simon Sung et al. Facile generation of iridium PCcarbeneP pincer complexes via water elimination from an alcohol proligand, *Dalton Transactions* (2017). DOI: [10.1039/C7DT03690F](https://doi.org/10.1039/C7DT03690F)

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