## Manganese Alkyl Carbonyl Complexes: From Iconic Stoichiometric Textbook Reactions to Catalytic Applications

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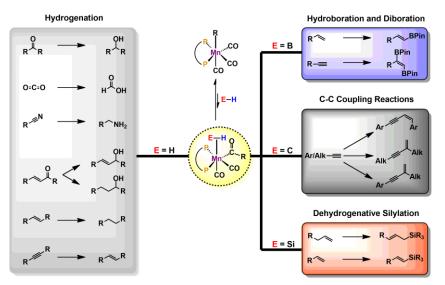
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The utilization of base-metal catalysts represents an emerging field in homogeneous catalysis. Among others, manganese-based complexes be highly were proven to competitive catalysts for (de)hydrogenation several reactions.

This lecture outlines the potential of alkylated Mn(I)carbonyl complexes for the activation of non-polar and moderately polar E-H (E = H,



C, Si, B) bonds and disclose our successful approach for the utilization of complexes in the field of homogeneous catalysis.<sup>1-5</sup> This will involve the rational design of manganese-complexes for hydrogenation reactions, including alkenes, which was not possible with defined manganese complexes before. Furthermore, the potential of our Mn-based catalysts in the field of hydrofunctionalization reactions for carbon-carbon multiple bonds will be discussed. Our investigations unveiled novel insights in reaction pathways of dehydrogenative silylation of alkenes and allowed *trans*-1,2-diboration of terminal alkynes, which was not reported for transition metals before.

## References

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