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**Tetrahedron Young Investigator Award 2009  
Recent Advances in Catalysis and Green Chemistry**  
**Michael J. Krische**

*Guest editor:* Stephen F. Martin*The Department of Chemistry and Biochemistry, The University of Texas at Austin, 1 University Station A5300, Austin, TX 78712, USA***Contents**

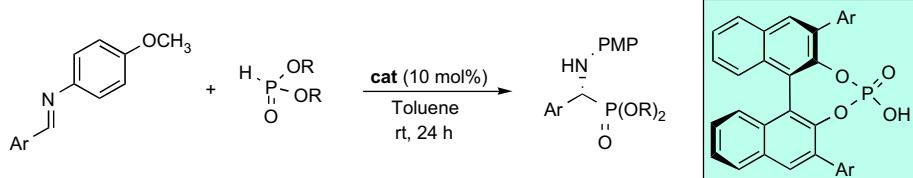
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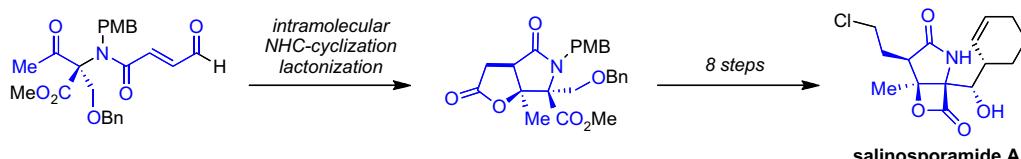
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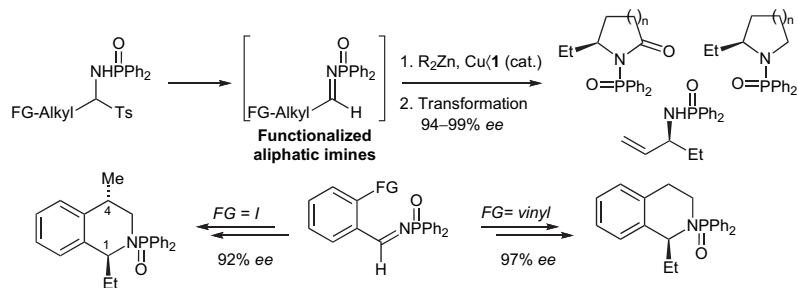
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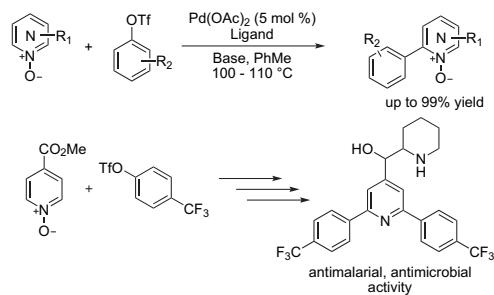
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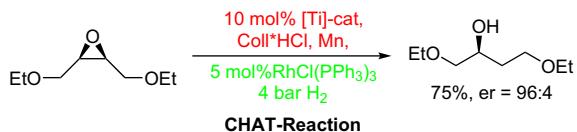
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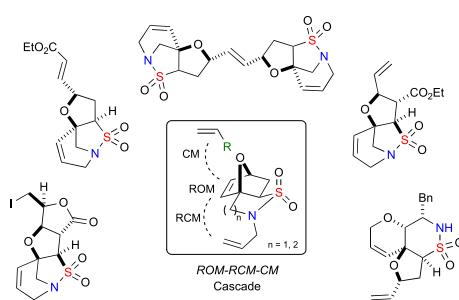
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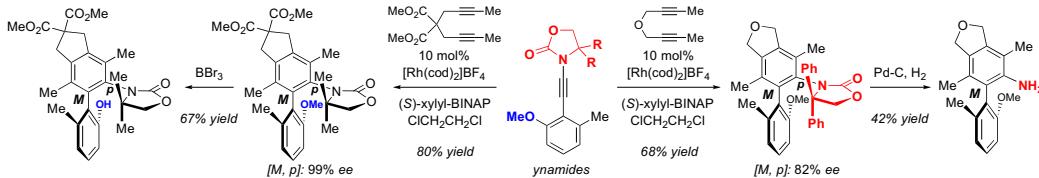
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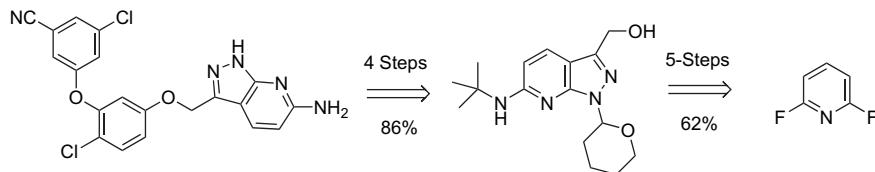
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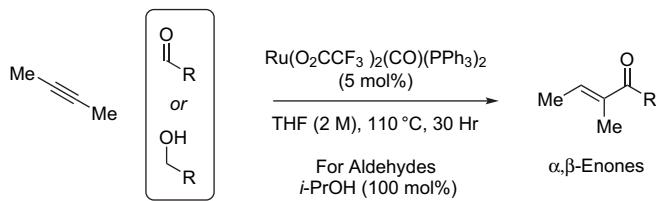
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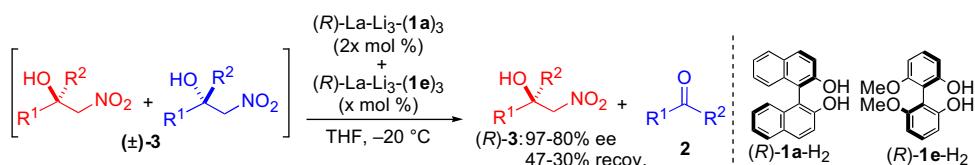
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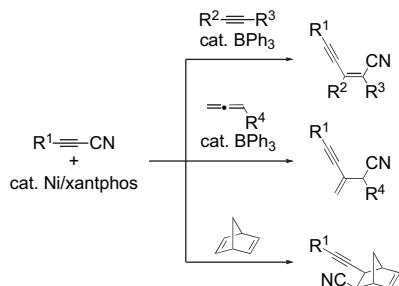
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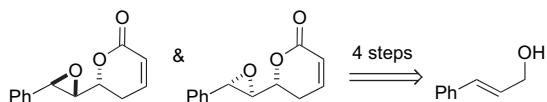
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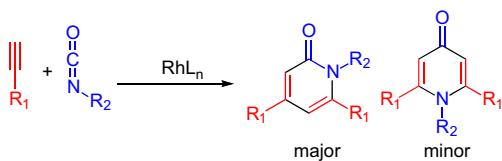
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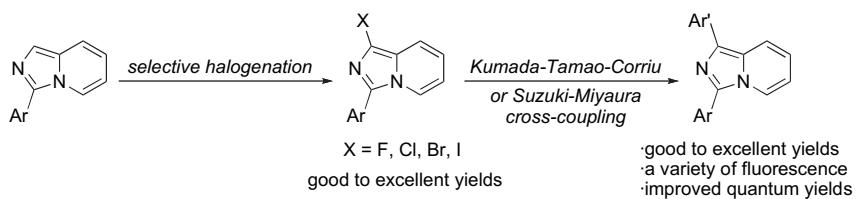
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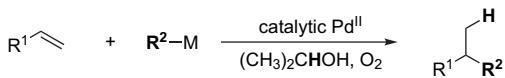
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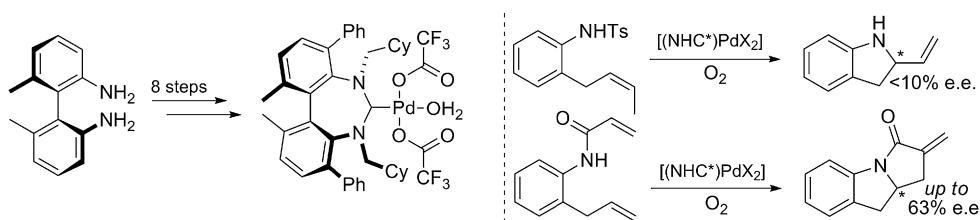
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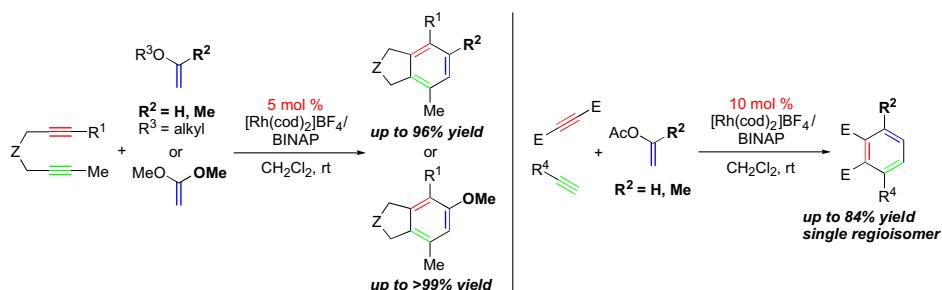
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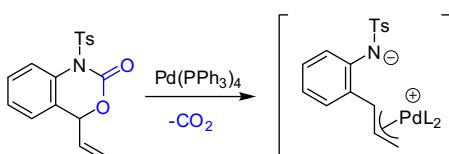
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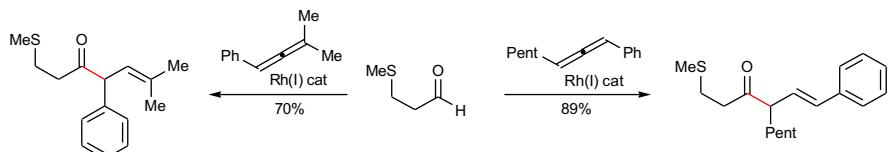
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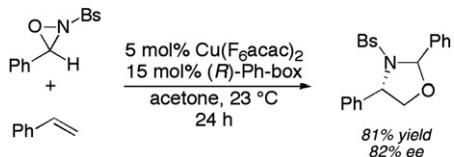
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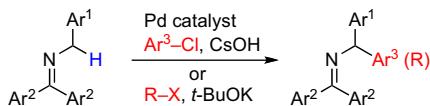
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Takashi Niwa, Takafumi Suehiro, Hideki Yorimitsu\*, Koichiro Oshima\*



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\*Corresponding author

† Supplementary data available via ScienceDirect

**COVER**

By combining the fields of hydrogenation and carbonyl addition, a broad new family of C-C bond forming hydrogenations and transfer hydrogenations is evoked, and the chemistry of carbonyl addition evolves beyond the use of stoichiometric organometallic reagents. Under hydrogenation conditions, unsaturates serve as synthetic equivalents to non-stabilized carbanions, enabling byproduct-free carbonyl addition. Under transfer hydrogenation conditions, alcoholic reactants serve as both hydrogen donors and aldehyde precursors, enabling byproduct-free carbonyl addition directly from the alcohol oxidation level.

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