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Enantioselective Intramolecular Hydroacylation of Unactivated Alkenes: An NHC-Catalyzed Robust and Versatile Formation of Cyclic Chiral Ketones  

Intramolecular Hydroacylation of Unactivated Alkenes by NHC Catalysis

**Significance:** Glorius and co-workers report a highly enantioselective intramolecular N-heterocyclic carbene catalyzed hydroacylation of unactivated alkenes 1 to afford ketones 2 which bear an \( \alpha \)-quaternary stereocenter. By employing pre-catalyst A, both aliphatic and aromatic aldehydes reacted smoothly to afford the desired products, by 5- or 6-exo-trig cyclization, in good to excellent yields and enantioselectivities. The scalability of the reaction was also proven by running the reaction on a two-gram scale (99% yield, er = 99:1).

**Comment:** Enantiopure ketones bearing an \( \alpha \)-quaternary stereocenter (2) are important motifs in biologically active molecules. The authors report an asymmetric approach to synthesize these compounds by employing simple unactivated olefins as starting materials. The methodology shows a strong functional group tolerance, proven by a ‘robustness screening’ and a broad substrate scope. A current limitation of the protocol is represented by the instability of products bearing an \( \alpha \)-tertiary stereocenter; they racemize under the basic reaction conditions. Previous mechanistic studies (Angew. Chem. Int. Ed. 2011, 50, 4983) from the same group on salicyaldehyde-derived substrates suggest a protonation of the alkene from the Breslow intermediate followed by C–C bond formation.