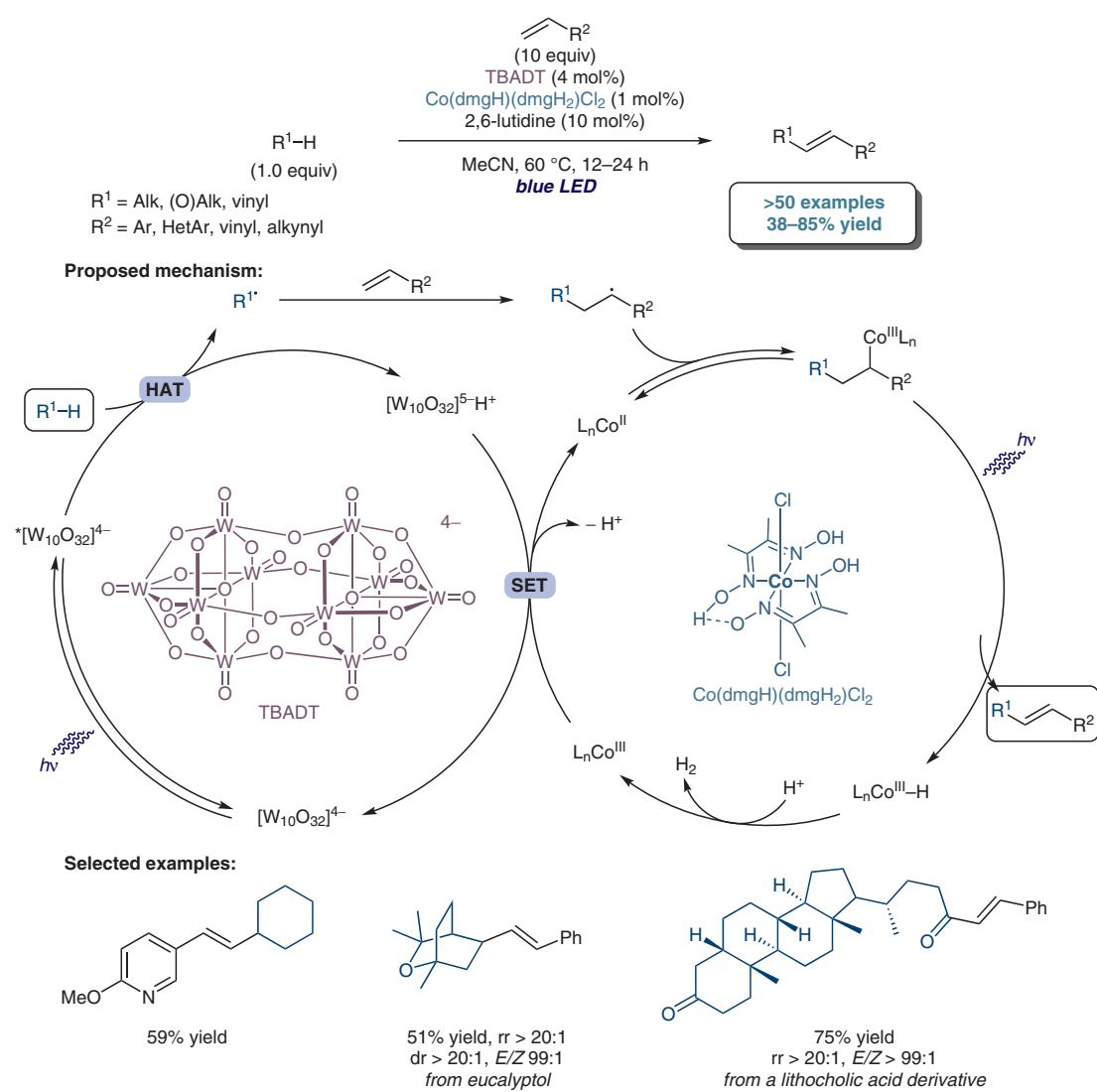


Dual-Catalyst Photoinduced Alkenylation of Alkanes and Aldehydes via C–H Activation



Significance: The authors report a site-selective photoinduced alkenylation of various alkanes and aldehydes with aryl and heteroaryl alkenes. This dehydrogenative reaction utilizes two photoactive catalysts (tetra-*n*-butylammonium decatungstate (TBADT) as HAT agent and $\text{Co}(\text{dmgH})(\text{dmgH}_2)\text{Cl}_2$), which eliminates the need for an external oxidant.

Comment: This mild alkenylation was used for the late-stage functionalization of diverse complex molecules, while still ensuring a high level of site-selectivity for the sterically most accessible and most electron-rich C–H bond. Wu and co-workers propose a mechanism that involves cooperation of the tungstate and the cobalt catalysts.