Trans-Selective Radical Silylzincation of Ynamides
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Category
Metal-Mediated Synthesis

Key words


Significance: The authors report a regio- and stereoselective silylzincation reaction of terminal ynamides using $\left(\mathrm{Me}_{3} \mathrm{Si}\right)_{3} \mathrm{SiH}$ and diethyl zinc. The resulting vinylzinc intermediates are trapped by a copper(l)-mediated substitution reaction to obtain $Z-\beta$-silylenamides in high yields.

Comment: The radical-chain process involves an addition of the $\left(\mathrm{Me}_{3} \mathrm{Si}\right)_{3} \mathrm{Si}$ radical to the ynamide to provide a $Z$-configured $\alpha$-amino vinylic radical which reacts with the dialkylzinc reagent by homolytic substitution to afford a $\alpha$-zincated $\beta$-silylenamide.
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