

Efficient and selective synthesis of *E*-vinylamines *via* tungsten(0)-catalyzed hydroamination of terminal alkynes.

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Streszczenie

CH= phenylacetylene, 4-methylphenylacetylene, 4-fluorophenylacetylene, 1-hexyne, methyl 2-propynyl ether, prop-2-yn-1-ol) with secondary amines (piperidine, pyrrolidine, morpholine, piperazine, methylpiperazine, 4-methylpiperidine and 3-methylpiperidine) was achieved in high yield (up to 99%), regioselectivity (only anti-Markovnikov product) and stereoselectivity (only *E*-isomers) within a maximum of 5 h in reactions catalyzed by the tungsten tetracarbonyl complex $\text{cis-[W(CO)}_4(\text{piperidine})_2\text{]at } 90^\circ\text{C}$ without any additional solvent."

Słowa kluczowe

E-enamines, hydroamination reaction, secondary amines, terminal alkynes, Tungsten catalyst

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