Category

Metals in Synthesis

Key words

ketones

organotin

palladium catalysis

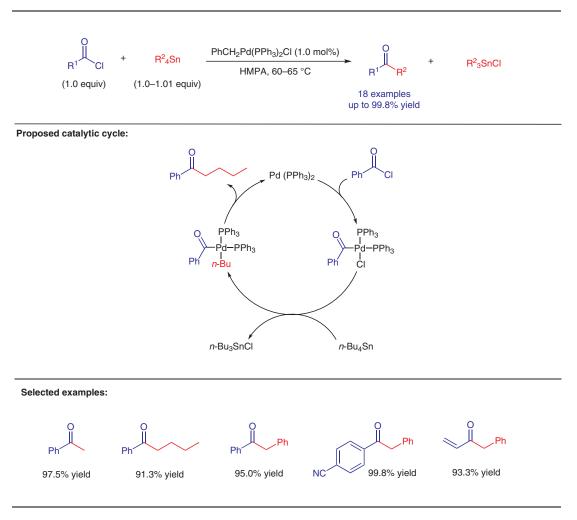
Stille-carbonylative cross-coupling



D. MILSTEIN, J. K. STILLE^{*} (COLORADO STATE UNIVERSITY, FORT COLLINS, USA) A General, Selective, and Facile Method for Ketone Synthesis from Acid Chlorides and Organotin Compounds Catalyzed by Palladium

J. Am. Chem. Soc. **1978**, 100, 3636–3638.

The Stille Carbonylative Cross-Coupling Reaction



Significance: Milstein and Stille reported a highyielding palladium-catalyzed carbonylative coupling reaction of acid chlorides and organotin reagents. The reaction is very mild, does not require inert atmosphere and shows tolerance to a wide scope of functional groups. **Comment:** Notably, aryltin groups are transferred in preference to alkyltin groups. The authors also showed that a second organic group attached to the tin can be transferred, however at a notably slower rate.

Review: J. K. Stille *Angew. Chem., Int. Ed. Engl.* **1986**, 25, 508–524.

SYNFACTS Contributors: Mark Lautens, Randy Sanichar Synfacts 2020, 16(03), 0314 Published online: 18.02.2020 **DOI:** 10.1055/s-0039-1690358; **Reg-No.:** L01420SF