M. EISOLD, D. DIDIER* (LUDWIG-MAXIMILIANS-UNIVERSITÄT MÜNCHEN, GERMANY) Stereoselective Access to Alkylidenecyclobutanes through γ-Selective Cross-Coupling Strategies *Org. Lett.* **2017**, *19*, 4046–4049.

Palladium-Catalyzed Synthesis of Alkylidenecyclobutanes

Significance: Alkylidenecyclobutanes are useful strained building blocks that can also be found in a number of natural compounds. In this work, the authors disclose the use of a γ -selective Suzuki–Miyaura cross-coupling for the synthesis of this class of compounds containing a quaternary stereocenter.

Comment: Achiral cyclobutenylmethylboronic esters reacted with aryl halides in the presence of a palladium catalyst and potassium hydroxide. The corresponding products were obtained in moderate to excellent yields and essentially with perfect γ -selectivity. In the case of δ -substituted substrates, high levels of diastereoselectivity were obtained. The use of (chiral) α -substituted starting materials generated the corresponding products containing a trisubstituted double bond functionality in high stereoselectivity (and enantioselectivity).

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Category

Metal-Catalyzed Asymmetric Synthesis and Stereoselective Reactions

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