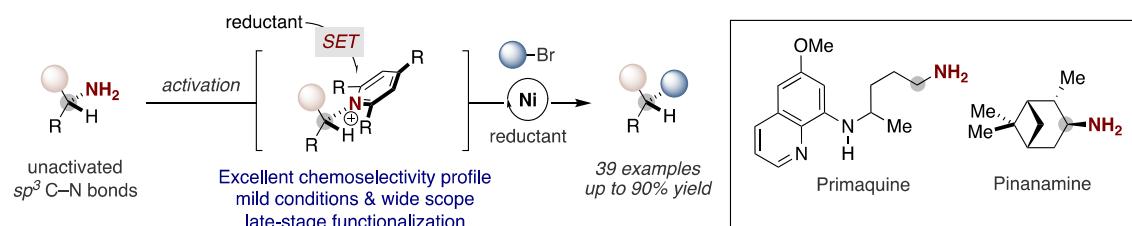


NICKEL CATALYZED REDUCTIVE DEAMINATIVE ARYLATION AT sp^3 CARBON CENTERS

R. Martin-Montero, R. Yatham, H. Yin, J. Davies, R. Martin*

Institute of Chemical Research of Catalonia (ICIQ), The Barcelona Institute of Science and Technology, Av. Països Catalans 16, 43007 Tarragona, Spain. ICREA, Passeig Lluís Companys, 23, 08010 Barcelona, Spain
e-mail: rmartin@iciq.es

Prompted by the ubiquity of aliphatic amines in a myriad of molecules that display biological relevance,¹ chemists have been challenged to design catalytic late-stage functionalization techniques by sp^3 C–N cleavage. As part of our ongoing interest in cross-electrophile coupling reactions and the recent successful implementation of pyrylium salts in cross-coupling reactions with well-defined organometallic reagents,^{2,3} we present herein a methodology for forging C–C bonds via sp^3 C–N cleavage of simple aliphatic amines with aryl halides. The protocol exhibits broad applicability with a diverse set of substitution patterns on both aryl and amine counterparts, even in the context of late-stage functionalization of advanced synthetic intermediates



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