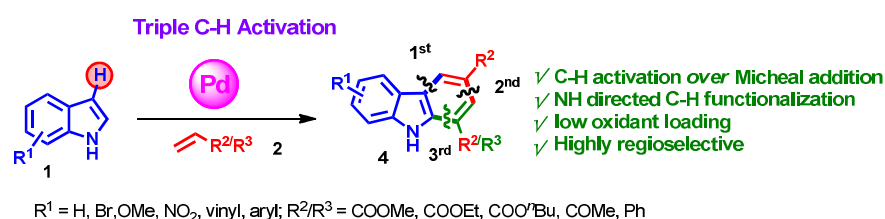


## Indole Directed C-H Activation: Direct Synthesis of Functionalized Carbazoles from Indoles via Triple C-H Activation

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Synthesis of small heterocyclic molecules in terms of selectivity, operational simplicity, functional group tolerance and environmental sustainability are in constant demand as majority of drugs; drug-like compounds contain hetero atom at their core. Since the discovery of the first intermolecular alkenylation (discovery of first C-H activation) by Fujiwara and Moritani in 1967.<sup>1</sup> Literature revealed that second successive alkenylation on alkene obtained by the first oxidative Heck is still unknown. In continuation of our ongoing work on the coupling reactions using in-house developed ligands<sup>2</sup> in this presentation I would like to discuss about our recent success on the synthesis of highly functionalized carbazoles from NH-indoles via palladium-catalyzed triple successive C-H activation.



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