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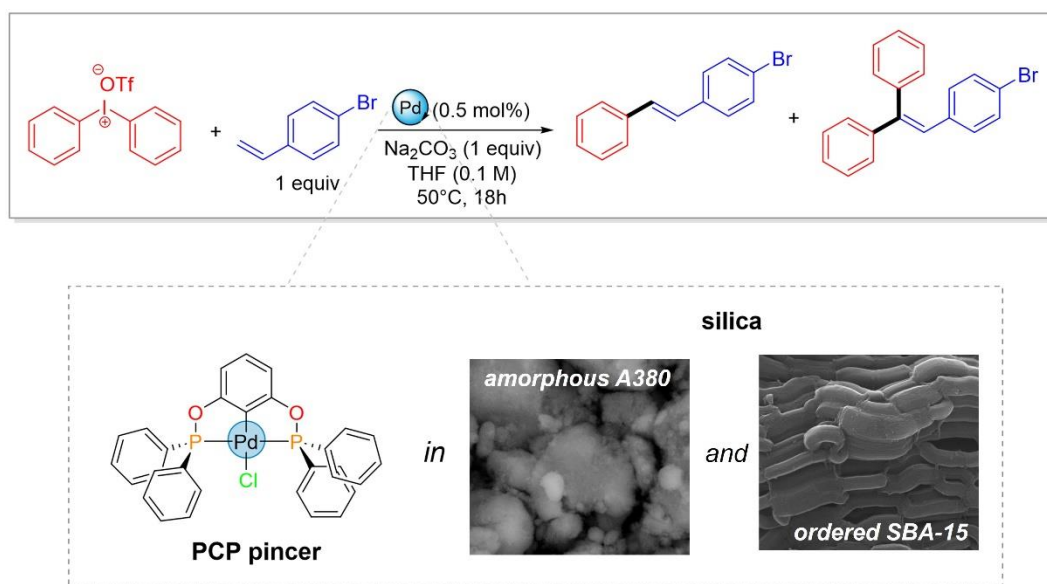
Heterogenized palladium pincer complexes for the cross-coupling reaction of styrenes and iodonium salts

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ABSTRACT

Phosphine-based palladium pincer complexes were initially thought of as readily steric- and electronically controllable, conferring a suitable balance between thermal stability and reactivity¹. When submitted to the mild reductive conditions of typical Heck and Suzuki couplings, however, these were shown to easily decompose to metal nanoparticles (NP), the active form of the catalyst². To take advantage of the NP formation, air- and moisture-stable phosphinite PCP Pd-pincer complexes were supported at 1 wt.% Pd in both amorphous and ordered silica by wet impregnation and evaluated in the coupling of styrenes and iodonium salts, which would follow a Pd(II)-Pd(IV) cycle in a homogeneous phase³. The best heterogeneous catalyst afforded 97% conversion and 17:3 selectivity to the monoarylated product by GC analysis, while both homogeneous and physical mixtures of support + complex yielded only traces of products. Further support pore structure and charge of the complex are also under evaluation and will be presented.



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