

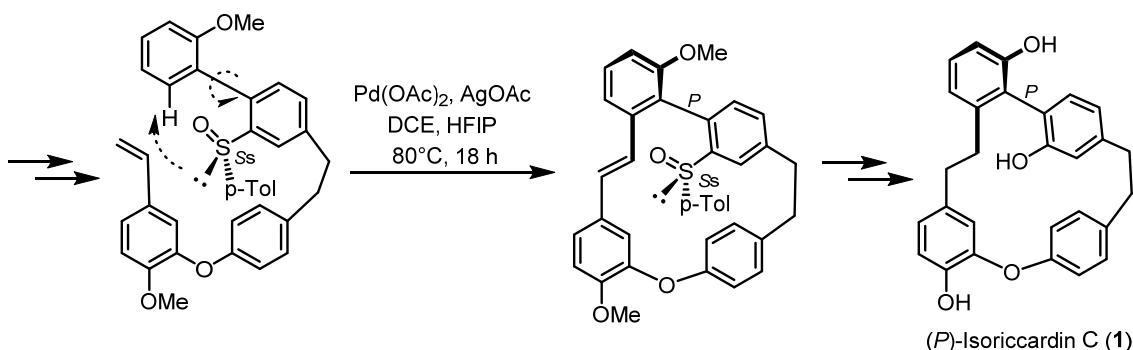
SYNTHESIS OF AXIALLY CHIRAL BIS(BIBENZYL) DERIVED FROM PERROTTETIN E VIA C–H ACTIVATED HECK TYPE CYCLIZATION

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Isoriccardin C (**1**) is one of six cyclic and axially chiral natural products isolated from liverworts and derived from the acyclic precursor Perrottetin E [1]. Regarding a variety of pharmacological interesting properties, different – but up to now not atroposelective – approaches for total syntheses were developed [2], but for **1** only in our group [3] using cyclization methods like Wittig or McMurry reactions. Atroposelective syntheses of these bis(bibenzyls), however, is a more challenging task [4]. Inspired by our promising results in the atroposelective synthesis of a structurally related isoplagiochin through Mizoroki-Heck reaction [5] we now investigated for **1** the C–H activated oxidative Fujiwara-Moritani option [6].



The sulfinyl group was used as an auxilliary with excellent metal coordination and ortho' directing properties during the C–H activation [7]. Finally, we obtained enantiopur Isoriccardin C (**1**) replacing the sulfinyl moiety by the hydroxyl group in the natural compound.

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