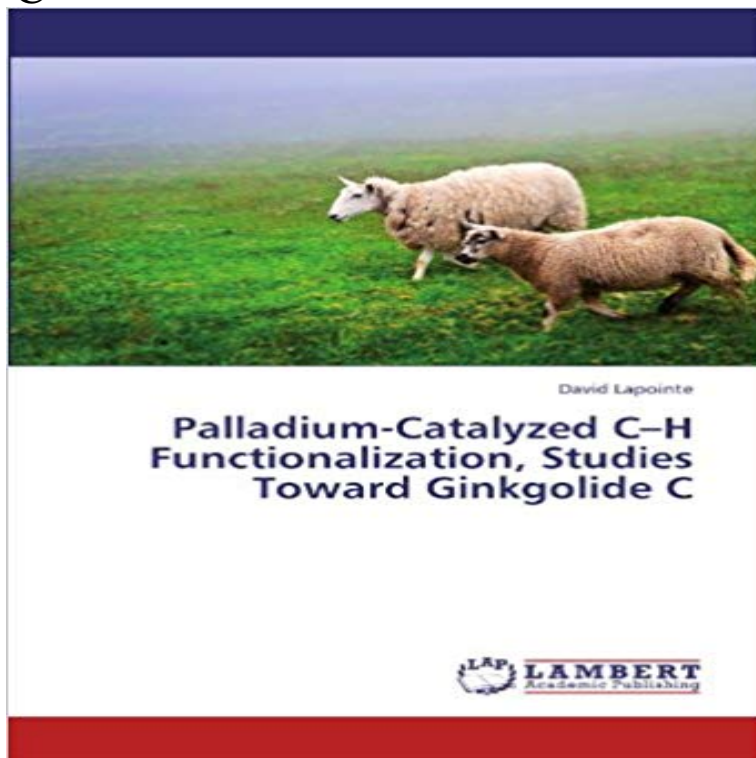


Palladium-Catalyzed C-H Functionalization, Studies Toward Ginkgolide C



The field of metal-catalyzed CH bond functionalizations is an incredibly vibrant and spans beyond the formations of biaryl motifs. The mechanistic aspects of the CH bond functionalization with metal-carboxylate complexes are highlighted. The role of additives, such as pivalic acid, is demonstrated with poorly reactive substrates. A chapter describes experimental and computational studies which suggested that a single pathway might be involved in the palladium-catalyzed CH bond functionalization of a wide range of (hetero)arenes. Afterward a general set of conditions are developed to form biaryls by direct arylation with a wide range of heteroarenes of various complexity level. The development of two new Pd-catalyzed methods based on our knowledge on the CH bond cleavage are apply to the formation of new scaffolds. Efforts toward the development of ligands to specifically promoted CH bond cleavage are presented. Lastly, most recent results on the study of the mechanism of the CH bond cleavage combining experimental and computational studies are discussed. In the second part of the thesis are presented our strategy toward the synthesis of ginkgolide C using gold catalysis.

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