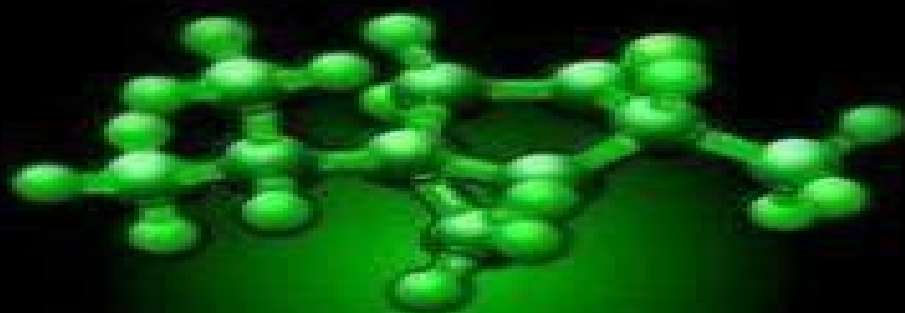


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of Reagents for
Organic Synthesis

**Reagents, Auxiliaries
and Catalysts
for C-C Bond Formation**

Edited by
Robert M. Coates and Scott E. Denmark

Reagents Auxiliaries And Catalysts For C C Bond Formation

Tse-Lok Ho



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June 2008 Metal Catalyzed Reductive C-C Bond Formation Michael J. Krische, 2007-07-20

The prototypical catalytic reductive C-C bond formations the Fischer-Tropsch reaction 1 and alkene hydroformylation 2 were discovered in 1922 and 1938 respectively 3 4 These processes which involve reductive coupling to carbon monoxide have long been applied to the industrial manufacture of commodity chemicals 5 Notably alkene hydroformylation also known as the oxo synthesis has emerged as the largest volume application of homogeneous metal catalysis accounting for the production of over 7 million metric tons of aldehyde annually Despite the impact of these prototypical reductive C-C bond formations this field of research lay fallow for several decades Eventually the increased availability of mild terminal reductants in particular silanes led to a renaissance in the area of catalytic reductive C-C bond formation For example the first catalytic reductive C-C couplings beyond hydroformylation which involve the hydrosilylative dimerization of conjugated dienes 6 12 appeared in 1969 approximately 16 years after the first reported metal catalyzed alkene hydrosilylation 13 Following these seminal studies the field of catalytic reductive C-C bond formation underwent explosive growth

culminating in the emergence of an evergrowing body of research encompassing a powerful set of transformations To our knowledge no thematic volumes devoted solely to metal catalyzed reductive C-C bond formation have been assembled For the first time in this issue of Topics in Current Chemistry we present a compilation of monographs from several leaders in this burgeoning area of research This collection of reviews serves to capture the diversity of catalytic reductive C-C couplings presently available and in turn the remarkable range of reactivity embodied by such transformations There is no indication that this field has reached its zenith and it is the hope of the present author that this volume will fuel further progress Fieser's

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provides a comprehensive overview of the rapidly developing field of asymmetric synthesis Using easy to understand graphical abstracts it presents 348 important catalytic and stoichiometric reactions leading to the synthesis of optically active chiral compounds The first part of the book covers reactions related to reductions oxidations carbon-carbon bond formation and carbon-heteroatom bond formation Each graphical abstract is accompanied by a list of important keywords and references to assist the reader The second part concentrates on experimental aspects describing synthetic procedures for selected chiral reagents and chiral auxiliaries and provides an invaluable reference tool for laboratory work Written with both the graduate student and professional organic chemist in mind this book will serve as an important resource for the

synthetic organic chemist **Organic Chemistry** Jonathan Clayden, Nick Greeves, Stuart Warren, 2012-03-15 A first and second year undergraduate organic chemistry textbook specifically geared to British and European courses and those offered in better schools in North America this text emphasises throughout clarity and understanding *Organic Chemistry* Mr. Rohit Manglik, 2024-05-07 Focuses on structure synthesis mechanisms and reactions of organic compounds Chemistry and Industry, 1999 **The Chemical Engineer**, 1999 Handbook of Organopalladium Chemistry for Organic Synthesis Ei-ichi Negishi, Armin de Meijere, 2003-11-24 Organized to provide maximum utility to the bench synthetic chemist The editor is well known for his work in exploring developing and applying organopalladium chemistry Contributors include over 24 world authorities in the field *Stereoselective Synthesis of Drugs and Natural Products* Vasyl Andrushko, Natalia Andrushko, 2013-08-16 Brings together the best tested and proven stereoselective synthetic methods Both the chemical and pharmaceutical industries are increasingly dependent on stereoselective synthetic methods and strategies for the generation of new chiral drugs and natural products that offer specific 3 D structures With the publication of *Stereoselective Synthesis of Drugs and Natural Products* researchers can turn to this comprehensive two volume work to guide them through all the core methods for the synthesis of chiral drugs and natural products *Stereoselective Synthesis of Drugs and Natural Products* features contributions from an international team of synthetic chemists and pharmaceutical and natural product researchers These authors have reviewed the tremendous body of literature in the field in order to compile a set of reliable tested and proven methods alongside step by step guidance This practical resource not only explores synthetic methodology but also reaction mechanisms and applications in medicinal chemistry and drug discovery The publication begins with an introductory chapter covering general principles and methodologies nomenclature and strategies of stereoselective synthesis Next it is divided into three parts Part One General Methods and Strategies Part Two Stereoselective Synthesis by Bond Formation including C C bond formation C H bond formation C O bond formation C N bond formation Other C heteroatom formation and other bond formation Part Three Methods of Analysis and Chiral Separation References in every chapter serve as a gateway to the literature in the field With this publication as their guide chemists involved in the stereoselective synthesis of drugs and natural products now have a single expertly edited source for all the methods they need *Catalytic Asymmetric Conjugate Reactions* Armando Cordova, 2010-12-01 This unique and long awaited handbook on this important topic in the hot field of stereoselective organic synthesis covers several types of nucleophiles Top international authors deal with modern forms of achieving stereoselective conjugate additions based on the use of chiral auxiliaries or asymmetric catalysis such as P N ligands organocatalysis domino reactions Lewis acid and base catalysis There is also a discussion of the employment of enantioselective conjugate addition transformations in total synthesis of important molecules With its reliable and previously unpublished experimental procedures this is a true source of high quality information

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