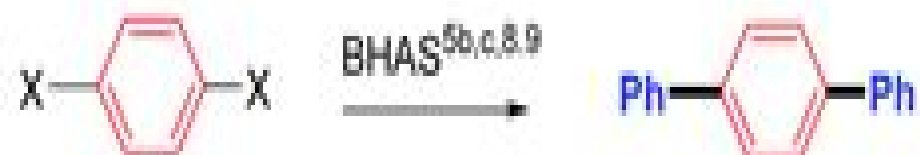


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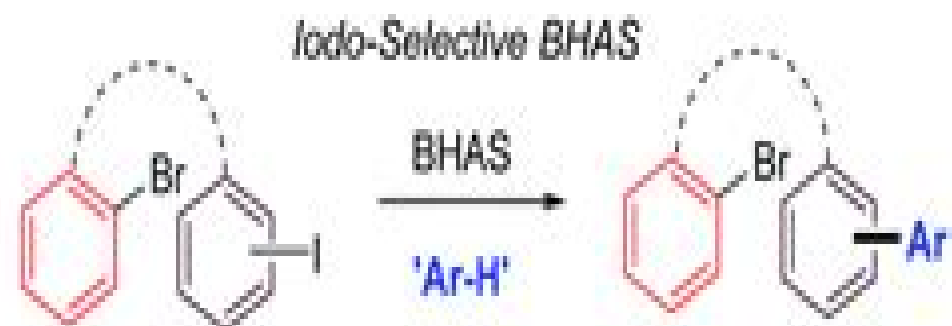
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## This Work

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# Homolytic Aromatic Substitution

**R. L. Henriquez**



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**Homolytic Aromatic Substitution** G. H. Williams, 2014-04-08 Homolytic Aromatic Substitution deals with the theoretical aspects of homolytic aromatic substitution reactions The effect of various kinds of free radicals on the substitution of atoms or groups usually hydrogen attached to aromatic nuclei is examined and the preparative use of homolytic substitution reactions is also considered This book is comprised of seven chapters and begins with an introduction to the general characteristics of homolysis along with homolytic and heterolytic aromatic substitution The discussion then turns to the various theoretical approaches used to rationalize aromatic substitution particularly those that are germane to a consideration of the problems of orientation and reactivity in homolytic substitution The following chapters explore homolytic arylation reactions including those between aryl radicals and aromatic substrates relative rates of arylation and partial rate factors for phenylation the reaction mechanism underlying intramolecular arylation and homolytic alkylation reactions The final chapter deals with hydroxylation and some other substitution reactions such as benzyloxylation acetyloxylation halogenation amination and amidation and mercuration This monograph will be of interest to organic chemists Aromatic Substitution Gareth Howel Williams, 1960 **Radicals** Nonhebel, Tedder, Walton, 1979-06-07 *Studies in Homolytic Aromatic Substitution* R. L. Henríquez, 1976 **Homolytic Aromatic Substitution** A. Nechvatal, 1952 **Studies on homolytic aromatic substitution** John Ivan George Cadogan, 1954 **Some studies in homolytic aromatic substitution** David Ian Davies, 1958 Studies on Homolytic Aromatic Substitution Reactions of Substituted Phenyl Radicals James Keith Hambling, King's College London. Department of Chemistry, 1957 Homolytic Aromatic Substitution Donald Holroyde Hey, Antony Nechvatal, T. S. Robinson, King's College London. Members, King's College London. Department of Physics, King's College London. Wheatstone Physics Laboratory, 1951 **Homolytic Aromatic Substitution--Another Issue** Gareth Howel Williams, 1960 Advances in Heterocyclic Chemistry, 1974-04-02 Advances in Heterocyclic Chemistry **Homolytic Aromatic Substitution** James Charles Arnold, 1974 **Homolytic Aromatic Substitution** David Ian Davies, Donald Holroyde Hey, Brian Michael Summers, King's College London. Department of Chemistry, King's College London. Members, 1971 **Homolytic Aromatic Substitution** David Ian Davies, Donald Holroyde Hey, Brian Michael Summers, King's College London. Department of Chemistry, King's College London. Members, 1970 *Modern Arylation Methods* Lutz Ackermann, 2009-04-22 Today arylation methods are belonging to the most important reaction types in organic synthesis Lutz Ackermann a young and ambitious professor has gathered a number of top international authors to present the first comprehensive book on the topic Starting from a historical review the book covers hot topics like Palladium catalyzed arylation of N H and alpha C H acidic Bonds Copper catalyzed arylation of N H and O H Bonds direct arylation reactions carbanion aromatic synthesis arylation reactions of alkenes alkynes and much more This compact source of high quality information is indispensable to synthetic chemists and those working in the pharmaceutical and chemical industry

*Aromatic and Heteroaromatic Chemistry* C W Bird, G W H Cheeseman, 2007-10-31 Specialist Periodical Reports provide systematic and detailed review coverage of progress in the major areas of chemical research. Written by experts in their specialist fields, the series creates a unique service for the active research chemist, supplying regular critical in-depth accounts of progress in particular areas of chemistry. For over 80 years, the Royal Society of Chemistry and its predecessor the Chemical Society have been publishing reports charting developments in chemistry, which originally took the form of Annual Reports. However, by 1967, the whole spectrum of chemistry could no longer be contained within one volume, and the series Specialist Periodical Reports was born. The Annual Reports themselves still existed but were divided into two, and subsequently three volumes covering Inorganic, Organic, and Physical Chemistry. For more general coverage of the highlights in chemistry, they remain a must. Since that time, the SPR series has altered according to the fluctuating degree of activity in various fields of chemistry. Some titles have remained unchanged, while others have altered their emphasis along with their titles; some have been combined under a new name, whereas others have had to be discontinued.

***Aromatic Reactions. -1 Homolytic Aromatic Substitution; -2 Hydrazobenzene Rearrangements*** J. Ray, 1965      Development of New Radical Cascades and Multi-Component Reactions Marie-Helene Larraufie, 2013-12-16

In this dissertation, Marie-Helene Larraufie develops original radical and palladium catalyzed methodologies to enable the synthesis of several classes of bioactive nitrogen-containing heterocycles. New radical cascades employing the N-acylcyanamide moiety offer straightforward routes to quinazolinones and guanidines, as well as new insights into the mechanism of homolytic aromatic substitutions. In parallel, Larraufie expands the scope of visible light photoredox catalysis to the ring opening of epoxides and aziridines, thus providing new sustainable alternatives for the generation of radicals. Furthermore, in a collaborative effort with the Catellani group, the author investigates dual palladium norbornene catalysis. First, she develops a C-amination coupling variant of the Catellani reaction with unprotected amines, which provides an expeditious route to phenanthridines. Then, she examines the influence of the chelating effect on Pd(IV) intermediates' reactivity, with the help of experimental studies and DFT calculations. The work in this thesis has resulted in numerous publications in high-impact journals. The clarity and depth of the experimental section will be useful for students and researchers working in this field.

***Aromatic Reactions. -1 Homolytic Aromatic Substitution; -2 Hydrazobenzene Rearrangements*** Gerald L. Ray, 1965      *Sulfonyl Ynamides as Useful Tools for N-Heterocyclic Chemistry* Tim Wezeman, 2016-12-31

Sulfonyl ynamides are highly versatile and synthetically useful reagents. This thesis details the modular synthesis and use of sulfonyl ynamides in order to access N-heterocyclic scaffolds such as quinolines and pyrazoles. The synthesis of a wide array of sulfonyl ynamides can be realized via copper-catalyzed amidative cross-couplings or by elimination of dichloroamide precursors. Additionally, the use of Sonogashira chemistry to further diversify terminal ynamides and the synthesis of solid-supported ynamides was investigated. Electrophilically activated amides can be reacted with sulfonyl ynamides in order to access highly functionalized 4-aminoquinolines. The straightforward

amide activation procedure with triflic anhydride and 2-chloropyridine was found to tolerate a wide range of substrates which allowed for the development of a library of 4-aminoquinolines with ease. Moreover, 4-aminopyrazoles can be prepared by reacting terminal sulfonyl ynamides with sydnone under copper catalysis. However, as the copper catalysts were also found to promote the degradation of the ynamides, a copper-free strain-promoted alternative was developed. An in situ prepared 3-azacyclohexyne was found to tolerate a wide array of C-4 substituted sydnone, producing a mixture of both the 3,4- and 4,3-fused pyrazoles in good yields. Additional investigations into heterocyclic methodology led to the development of highly sophisticated non-symmetrical and axially chiral dibenzo 1,3-diazepines, oxazepines, and thiazepines from simple commercially available anilines. The anilines were coupled to their corresponding reaction partners via a chloromethyl intermediate and the 7-membered ring was subsequently formed using direct arylation.

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