

Convenient and Efficient Suzuki Miyaura Coupling Reactions of *Meso*-Halogenated BODIPYs

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Abstract – *In this study, we synthesized new conjugated BODIPY compounds with potential fluorescence by treated the BODIPY compound containing bromine in the meso position with different boronic acids. BODIPY cores derived from meso position are used in interesting applications such as pH probe and redox sensors.*

Keywords – BODIPY, halogenated BODIPY, Suzuki-Miyaura coupling reactions, boronic acids, photophysical and electrochemical properties.

Introduction

Boradiaza-s-indacene (boron dipyrin or boron dipyrromethene, BODIPY) dyes have gained outstanding place for use in fluorescent materials, labels, and probes. Derivatization at the 8-position, or *meso*-position, is a preferred method for the construction of complex BODIPY fluorophores. The nature of the new substituent has strong influence on the spectral properties of the dyes [1]. Spectroscopic features can be easily accommodated by different synthetic strategies on BODIPY nucleus [2]. Halogenated BODIPYs are important precursors to synthesize a variety of substituted BODIPYs and potential sensitizers for photodynamic therapy (PDT). Halogenated systems can be derivatized by transition metal catalyzed cross-coupling reactions [3].

The palladium-catalyzed Suzuki cross-coupling reaction is one of the most efficient methods for the construction of C_{aryl}-C_{aryl} bonds and has found widespread use in organic synthesis. Carbon-carbon bond formation reactions are some of the most important processes in chemistry, that provide key steps in the building of more complex molecules from simple precursors [4].

Conclusion

In this study, 4,4-difluoro-8-(4-bromophenyl)-4-bora-3a,4a-diaza-s-indacene (BODIPY) was synthesized from reaction of pyrrole with 4-bromobenzaldehyde and complexation with BF₃.OEt₂. The synthesized BODIPY nucleus was treated with different boronic acids and new BODIPY compounds derived from the 8-position were obtained.

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References

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