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Photoactive formyl benzimidazoles: New molecular scaffolds for optical sensing

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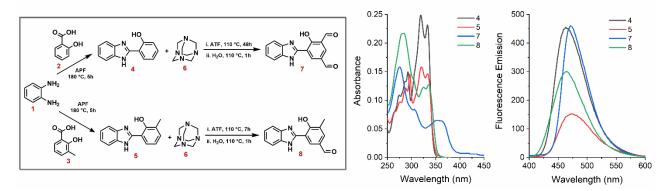
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ABSTRACT

Excited-state intramolecular proton transfer (ESIPT) is a photochemical process that results in the formation of a tautomer with an electronic structure different from that of the initial conformer. ESIPT-responsive fluorophores exhibit remarkable photophysical properties, such as large Stokes shifts and intense fluorescence emission.^{1,2} This work presents the synthesis and photophysical characterization of photoactive compounds via ESIPT, specifically substituted hydroxyphenyl benzazolic heterocycles, obtained through the classical condensation reaction methodology between a functionalized aromatic amine and a salicylic acid derivative.³ From these precursors, the Duff formylation methodology enabled the synthesis of mono- and biformylated benzazolic heterocycle derivatives. This modification preserved important photophysical characteristics, such as absorption in the UV-region and fluorescence emission in the visible region, with a large Stokes shift (~150 nm), while causing a significant increase in the fluorescence of these molecules if compared with their precursors. Notably, the presence of a formyl group in an aromatic system facilitates the design of new fluorescent compounds.⁴



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