

## Radical cascade sulfonylation/cyclization of benzimidazole derivatives with sulfinates induced by visible light

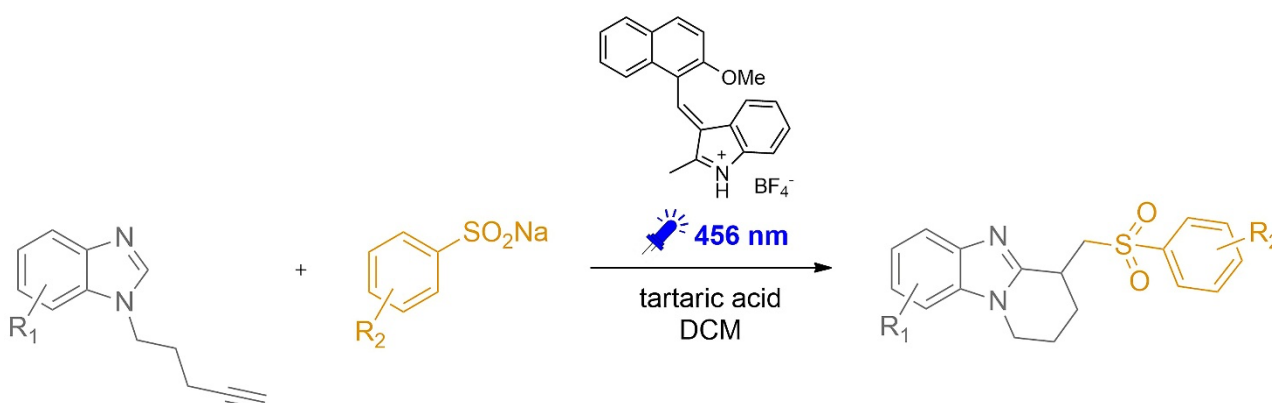
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Organosulfur moieties are essential features in numerous natural products<sup>1</sup> and functional materials.<sup>2</sup> They hold particular significance in the pharmaceutical industry,<sup>3</sup> with over 150 sulfur-containing drugs currently available on the market.<sup>4</sup>

As a result, the development of milder and more environmentally friendly methodologies for C-S bond formation has been the object of intensive work for organic chemists. Photocatalysis has emerged as a powerful approach, enabling unique reactivity pathways that are challenging to achieve through conventional thermal transformations.

In recent years, our research group has focused on the photoinduced generation of key sulfur-centered radicals, such as sulfonyl and thiyl radicals,<sup>5</sup> aiming to leverage them for the functionalization of unsaturated compounds. In this context, benzimidazole derivatives with appended alkyne functionality and sodium sulfinates have been employed as the starting materials in a radical cascade sulfonylation/cyclization mediated by bench-stable diarylmethyl cation tetrafluoroborate organophotocatalysts under visible light irradiation.



**Figure 1.** Visible light sulfonylation-cyclization of benzimidazoles with sulfinates.

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