Photodimerization of Arenediynes

Significance: Cyclic enediynes as well as terminal acyclic enediynes are well known to undergo photochemical C₁–C₆ cycloaromatization. The authors explored the reactivity of naphthalenyl-substituted arenediynes (1 and 2). While the methoxy-substituted derivative 2 undergoes a photo-Bergman cyclization upon irradiation at 300 nm, 1 shows no formation of C₁–C₆ or C₁–C₅ under these conditions. Irradiation of 1 at 350 nm, however, yields a photodimerized product (4).

Comment: The tandem [2+2] photocyclization yields two products in a 4:1 ratio. The structure of the major product (4) was determined by X-ray crystallographic analysis. Based on NMR studies, the authors suggest that the minor product is a diastereomer of 4.

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