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Insights on the synthesis of asymmetric curcumin derivatives and their biological activities (2019) *European Journal of Medicinal Chemistry*, 183, art. no. 111704, .

DOI: 10.1016/j.ejmech.2019.111704

Abstract

Curcumin is a small organic molecule with pleiotropic biological activities. However, its multiple structural-pharmacokinetic challenges prevent its development into a clinical drug. Various structural modifications have been made to improve its drug profile. In this review, we focus on the methods adopted in the synthesis of asymmetric curcumin derivatives and their biological activities and forecast the future of this exciting class of compounds in the field of medicine. © 2019 Elsevier Masson SAS

Author Keywords

Asymmetric curcumin; Bioactivity; Diketone; Monoketone; Synthesis

Funding details

Ministry of Higher Education, MalaysiaMOHEGUP-2017-001

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Publisher: Elsevier Masson SAS

ISSN: 02235234 CODEN: EJMCA

Language of Original Document: English Abbreviated Source Title: Eur. J. Med. Chem.

2-s2.0-85072380584

Document Type: Short Survey **Publication Stage:** Final

Source: Scopus



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