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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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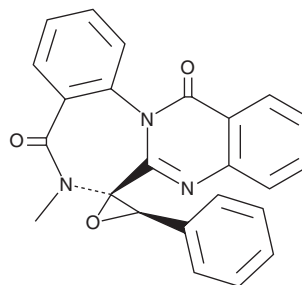
PRODUCT INFORMATION



Benzomalvin C

Item No. 22064

CAS Registry No.: 157047-98-8
Formal Name: (2R,3S)-rel-(+)-6'-methyl-3-phenyl-spiro[oxirane-2,7'(13'H)-quinazolino[3,2-a][1,4]benzodiazepine]-5',13'(6'H)-dione
MF: C₂₄H₁₇N₃O₃
FW: 395.4
Purity: ≥95%
Supplied as: A solid
Storage: -20°C
Stability: ≥2 years



Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Benzomalvin C is supplied as a solid. A stock solution may be made by dissolving the benzomalvin C in the solvent of choice. Benzomalvin C is soluble in organic solvents such as ethanol, methanol, DMSO, and dimethyl formamide, which should be purged with an inert gas.

Description

Benzomalvin C is a weak antagonist of the neurokinin-1 (NK₁) receptor inhibiting binding of substance P by 46% when used at 100 µg/ml *in vitro*.¹ It is also a weak inhibitor of indoleamine 2,3-dioxygenase (IDO) with an IC₅₀ value of 130 µM for recombinant IDO.² It was isolated from *Penicillium* and contains an epoxide group at C-19 and C-20, which is not present in benzomalvins A, B, or E.

References

1. Sun, H.H., Barrow, C.J., Sedlock, D.M., *et al.* Benzomalvins, new substance P inhibitors from a *Penicillium* sp. *J. Antibiot. (Tokyo)* **47(5)**, 515-522 (1994).
2. Jang, J.-P., Jang, J.-H., Soung, N.-K., *et al.* Benzomalvin E, an indoleamine 2,3-dioxygenase inhibitor isolated from *Penicillium* sp. FN070315. *J. Antibiot. (Tokyo)* **65(4)**, 215-217 (2012).

WARNING

THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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