PRODUCT INFORMATION



Tetrahydroepiberberine

Item No. 36177

CAS Registry No.: 38853-67-7

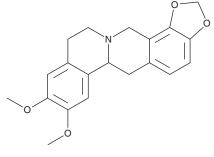
Formal Name: 6,6a,11,14-tetrahydro-8,9-dimethoxy-12H-

benzo[a]-1,3-benzodioxolo[4,5-g]quinolizine

Synonym: MF: $C_{20}H_{21}NO_4$ 339.4 FW: **Purity:** ≥98% Supplied as: A solid Storage: -20°C Stability: ≥4 years

Item Origin: Plant/Sinomenium acutum

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



Laboratory Procedures

Tetrahydroepiberberine is supplied as a solid. A stock solution may be made by dissolving the tetrahydroepiberberine in the solvent of choice, which should be purged with an inert gas. Tetrahydroepiberberine is soluble in chloroform.

Description

Tetrahydroepiberberine is an alkaloid originally isolated from P. rhoeas that has diverse biological activities. 1,2 It is active against the fungus C. albicans (MIC = 8 µg/ml), as well as the bacteria E. coli, P. aeruginosa, P. mirabilis, K. pneumoniae, A. baumannii, S. aureus, and B. subtilis (MICs = 32-64 µg/ml, respectively).² Tetrahydroepiberberine (16 μg/ml) inhibits the cytopathogenic effect of parainfluenza-3 (PI-3) virus in Vero cells. It decreases the viability of HaCaT keratinocytes (IC₅₀ = 150 μ M).

References

- 1. Hijazi, M.A., Aboul-Ela, M., Bouhadir, K., et al. Cytotoxic activity of alkaloids from Papaver rhoeas growing in Lebanon. Rec. Nat. Prod. 11(2), 211-216 (2017).
- 2. Orhana, I., Ozcelik, B., Karaoğlu, T., et al. Antiviral and antimicrobial profiles of selected isoquinoline alkaloids from Fumaria and Corydalis species. Z. Naturforsch. C. J. Biosci. 62(1-2), 19-26 (2007).

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

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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