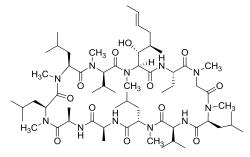


PRODUCT DATA SHEET

Code No.: BIA-C1248

Pack sizes: 1 mg, 5 mg



Synonyms

5-(N-methyl-D-valine)cyclosporine A

Specifications

Cyclosporin H

CAS #	:	83602-39-5
Molecular Formula	:	C ₆₂ H ₁₁₁ N ₁₁ O ₁₂
Molecular Weight	:	1202.6
Source	:	Trichoderma sp.
Appearance	:	White powder
Purity	:	>95% by HPLC
Long Term Storage	:	-20°C
Solubility	:	Soluble in ethanol, methanol, DMF or DMSO. Limited water solubility.

Application Notes

Cyclosporin H is a minor analogue of the cyclosporin family which is immunologically inactive as it does not bind to immunophilin. Cyclosporin H is the most extensively investigated of the minor cyclosporin analogues. It is a potent inhibitor of tumor-promoting phorbol esters on mouse skin in vivo, and of calcium/calmodulin-dependent EF-2 phosphorylation in vitro, a potent and selective antagonist of formyl peptide receptor and inhibitor of formyl peptide-induced superoxide formation.

References

- 1. The weak immunosuppressant cyclosporine D as well as the immunologically inactive cyclosporine H are potent inhibitors in vivo of phorbol ester TPA-induced biological effects in mouse skin and of Ca2+/calmodulin dependent EF-2 phosphorylation in vitro. Gschwendt M. et al., BBRC 1988, 150, 545.
- Differential inhibition of human neutrophil activation by cyclosporins A, D, and H. Cyclosporin H is a potent and effective inhibitor of formyl peptide-induced superoxide formation. Wenzel-Seifert K. et al., J. Immunol. 1991, 147, 1940.
- Cyclosporin H is a potent and selective formyl peptide receptor antagonist. Comparison with N-t-butoxycarbonyl-L-phenylalanyl-L-leucyl-L-phenylalanyl-L-leucyl-L-phenylalanine and cyclosporins A, B, C, D, and E. Wenzel-Seifert K. & Seifert R. J. Immunol. 1993, 150, 4591.
- 4. Cyclosporin H is a potent and selective competitive antagonist of human basophil activation by N-formylmethionyl-leucyl-phenylalanine. de Paulis A. et al., J. Allergy Clin. Immunol. 1996, 98, 152.

Updated: 24 May 2019

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