

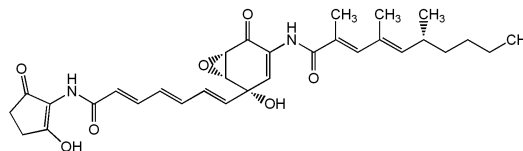
BVT-0091

08-Jun-2011

Manumycin A

BVT-0091-M001	1 mg
BVT-0091-M005	5 mg
BVT-0091-M010	10 mg

Formula	C ₃₁ H ₃₈ N ₂ O ₇
MW	550.6
CAS	52665-74-4



Handling / Storage

Shipping	BLUE ICE
Short Term Storage	+4°C
Long Term Storage	-20°C

Protect from light.

Use / Stability

Stable for at least 1 year after receipt when stored at -20°C. After reconstitution protect from light at -20°C.

MSDS available at www.adipogen.com or upon request.

Product Specifications

Source/Host	Isolated from <i>Streptomyces parvulus</i> .
Purity	≥96% (HPLC)
Identity	Determined by ¹ H-NMR.
Appearance	Yellow to brown powder.
Solubility	Soluble in DMSO or methanol; insoluble in water.

Product Description

- Antibiotic.
- Potent, selective and competitive cell permeable rasfarnesyltransferase inhibitor (IC₅₀ = 30 nM).
- Does not affect geranylgeranyltransferase (IC₅₀ = 180 μM).
- Inhibition is competitive with respect to farnesyl pyrophosphate and non-competitive with respect to Ras.
- Neutral sphingomyelinase inhibitor.
- Blocks insulin-induced MAP kinase activation in rat cardiac myocytes (19 μM).

WARNING: Intended for research use only. This product is not intended or approved for human, diagnostics, therapeutic or veterinary use. Use of this product for human or animal testing is extremely hazardous and may result in disease, severe injury, or death. **MATERIAL SAFETY DATA:** Review the complete Material Safety Data Sheet before use.

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Product Specific References

1. Identification of Rasfarnesyltransferase inhibitors by microbial screening: M. Hara et al.; PNAS **90**, 2281 (1993)
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9. A yeast-based genomic strategy highlights the cell protein networks altered by FTase inhibitor peptidomimetics: G. Porcu, et al.; Mol. Cancer **9**:197 (2010)
10. Targeting farnesyl-transferase as a novel therapeutic strategy for mevalonate kinase deficiency: In vitro and in vivo approaches: L. De Leo, et al.; Pharmacol. Res. **61**, 506 (2010)

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