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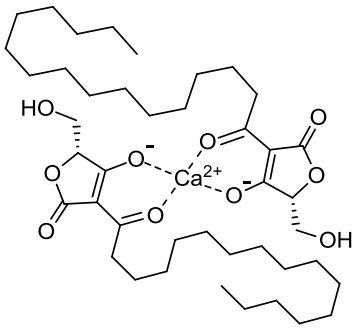
PRODUCT DATA SHEET

RK-682

Code No.: **BIA-R1082**

Pack sizes: **0.5 mg, 2.5 mg**



Synonyms : TAN 1364B, CI 010

Specifications

CAS #	:	**332131-32-5**
Molecular Formula	:	**$C_{42}H_{70}CaO_{10}$**
Molecular Weight	:	**775.1**
Source	:	***Streptomyces* sp.**
Appearance	:	**White Solid**
Purity	:	**>95% by HPLC**
Long Term Storage	:	**-20°C**
Solubility	:	**Soluble in ethanol, methanol, DMF or DMSO. Poor water solubility.**

Application Notes

RK-682 is a dimeric calcium complex of the major analogue of a tetronic acid complex isolated from Streptomyces. Although reported by researchers at RIKEN in 1995, subsequent synthesis in 2001 showed that RK-682 was in fact the calcium complex formed during silica chromatography. Confusion about the structure of RK-682 has lead to the monomeric sub-unit, TAN 1364B (3-hexadecanoyl-5-hydroxymethyltetronic acid) being mis-named as RK-682 by many suppliers. As either the dimer or monomer, RK-682 inhibits protein tyrosine phosphatases, phospoholipase A2, heparinase and HIV-1 protease. However, is unclear whether biological activity is due to the monomer (TAN 1364B) or dimeric complex (RK-682).

References

1. RK-682, a potent inhibitor of tyrosine phosphatase, arrested the mammalian cell cycle progression at G1phase. Hamaguchi T. et al., FEBS Lett. 1995, 372, 54.
2. Structure-based design of a selective heparanase inhibitor as an antimetastatic agent. Ishida K. et al., Mol. Cancer Ther. 2004, 3, 1069.
3. The mechanism of ATP-induced long-term potentiation involves extracellular phosphorylation of membrane proteins in guinea-pig hippocampal CA1 neurons. Fujii S. et al., Neurosci. Lett. 1995, 187, 130.
4. Asymmetric synthesis of a 3-acyltetronic acid derivative, RK-682, and formation of its calcium salt during silica gel column chromatography. Sodeoka M. et al., Chem. Pharm. Bull. 2001, 49, 206

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