

Veratridine

Product Number **V5754**

Storage Temperature -0 °C

Product Description

Molecular Formula: C₃₆H₅₁NO₁₁

Molecular Weight: 673.8

CAS Number: 71-62-5

Melting Point: 180 °C

λ_{\max} : 262 nm (ethanol)

Specific Rotation: +4.9° (25 mg/ml, ethanol, 25 °C)¹

Veratridine is one of several alkaloids isolated from the seeds of *Schoenocaulon officinale* and from the rhizome of *Veratrum album*. The crude extract is called veratrine or sabadilla, and contains cevadine, veratridine, cevadilline, sabadine, and cevine. It has been used as an insecticide, acting as a paralytic agent with higher toxicity to insects than to mammals.² However, purified veratridine is highly toxic to all animals tested. Sabadilla has an LD₅₀ of 5000 mg/kg, but veratridine has an LD₅₀ of 1350 µg/kg.

In cells, veratridine acts to open voltage-dependent Na⁺ channels and prevents their inactivation. This, in turn, opens voltage-activated calcium channels, increasing intracellular calcium content and inducing neurotransmitter release.^{3,4} It has been used to study Na⁺ channel blockers such as vincamine and vincanol by inducing Na⁺ channels in the presence and absence of the drugs being tested.⁵

Precautions and Disclaimer

For Laboratory Use Only. Not for drug, household or other uses.

Preparation Instructions

Veratridine is soluble in ethanol or DMSO (50 mg/m), and is freely soluble in chloroform. The solubility in water is dependent on pH. The free base form is very slightly soluble in water, but dissolves easily at 50 mg/ml in 1 M HCl.

Storage/Stability

Solutions of veratridine in chloroform are stable for at least 6 months stored at -20 °C.

References

1. McKinney, L. C. et al., Purification, solubility and pKa of veratridine. *Anal. Biochem.*, **153(1)**, 33-38 (1986).
2. Bloomquist, J. R. Ion channels as targets for insecticides. *Ann. Rev. Entomol.*, **41**, 163-190 (1996).
3. Maroto, R., et al., Effects of Ca²⁺ channel antagonists on chromaffin cell death and cytosolic Ca²⁺ oscillations induced by veratridine. *Eur. J. Pharmacol.*, **270(4)**, 331-339 (1994).
4. Dobrev, D., et al., Voltage-activated calcium channels involved in veratridine-evoked [³H]dopamine release in rat striatal slices. *Neuropharmacology*, **37(8)**, 973-982 (1998).
5. Erdo, S. A., et al., Vincamine and vincanol are potent blockers of voltage-gated Na⁺ channels. *Eur. J. Pharmacol.*, **14(1-2)**, 69-73 (1996).

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