# PRODUCT INFORMATION



## Ranaconitine

Item No. 34774

**CAS Registry No.:** 1360-76-5

Formal Name: 20-ethyl- $1\alpha$ , $14\alpha$ , $16\beta$ -trimethoxy-aconitane-

4,7,8,9-tetrol 4-[2-(acetylamino)benzoate]

Synonym: 7-hydroxy Lappaconitine

MF:  $C_{32}H_{44}N_2O_9$ 600.7 FW:

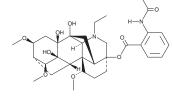
≥98% **Purity:** 

UV/Vis.:  $\lambda_{max}$ : 223, 252 nm

Supplied as: A solid Storage: -20°C Stability: ≥4 years

Item Origin: Plant/Aconitum carmichaelii

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



## **Laboratory Procedures**

Ranaconitine is supplied as a solid. A stock solution may be made by dissolving the ranaconitine in the solvent of choice, which should be purged with an inert gas. Ranaconitine is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of ranaconitine in these solvents is approximately 1, 10, and 30 mg/ml, respectively.

Ranaconitine is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, ranaconitine should first be dissolved in DMF and then diluted with the aqueous buffer of choice. Ranaconitine has a solubility of approximately 0.2 mg/ml in a 1:4 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

## Description

Ranaconitine is an alkaloid that has been found in Aconitum and has analgesic and local anesthetic activities in mice (ED<sub>50</sub>s = 6.99 and 0.167  $\mu$ mol/kg, respectively).<sup>1</sup>

## Reference

1. Bello-Ramírez, A.M. and Nava-Ocampo, A.A. A QSAR analysis of toxicity of Aconitum alkaloids. Fundam. Clin. Pharmacol. 18(6), 699-704 (2004).

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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