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Product Information

Product ID V3355 CAS No. 2182-14-1

Chemical Name (28,38,48,5α,128,19α)-4-(Acetyloxy)-6,7-didehydro- 3-hydroxy-16-

methoxy-1-methyl-aspidospermidine-3-carboxylic acid methyl ester

Synonym Vindolin, NSC 91994

Formula $C_{25}H_{32}N_2O_6$ Formula Wt. 456.53 Melting Point 164-165°C Purity ≥98%

Solubility Soluble in chloroform or ethanol.

Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
V3355	25 mg	\$87.60
V3355	100 mg	\$243.10
V3355	500 mg	\$738.90

Store Temp -20°C Ship Temp Ambient

Description Vindoline is a semi-synthetic vinca alkaloid originally found in *Catharanthus* that exhibits anti-diabetic and antacid activities; it is an intermediate in the synthesis of vinblastine. Unlike other vinca alkaloids, vindoline is only weakly cytotoxic, binding poorly

to tubulin. Vindoline increases glucose-stimulated insulin release and inhibits Kv2.1 K+ channels, decreasing outward K+ current and lowering blood glucose, Hb1Ac, and triglyceride levels in animal models of diabetes. Vindoline also inhibits H+/K+ ATPases,

potentially decreasing gastric acid secretion.

References Yao XG, Chen F, Li P, et al. Natural product vindoline stimulates insulin secretion and efficiently ameliorates glucose homeostasis in diabetic murine models. J Ethnopharmacol. 2013 Oct 28;150(1):285-97. PMID: 24012527.

> Freitas CS, Baggio CH, Mayer B, et al. Inhibition of gastric H+, K(+)-ATPase activity by compounds from medicinal plants. Nat Prod Commun. 2011 Sep;6(9):1253-4. PMID: 21941891.

Sertel S, Fu Y, Zu Y, et al. Molecular docking and pharmacogenomics of vinca alkaloids and their monomeric precursors, vindoline and catharanthine. Biochem Pharmacol. 2011 Mar 15;81(6):723-35. PMID: 21219884.

Caution: This product is intended for laboratory and research use only. It is not for human or drug use.