PRODUCT INFORMATION



Ginsenoside Rh₁

Item No. 27318

CAS Registry No.:	63223-86-9	
Formal Name:	(3β,6α,12β)-3,12,20-trihydroxydammar-24-en-6-yl,	F
	β-D-glucopyranoside	ОН У ОН
Synonyms:	20(S)-Ginsenoside Rh ₁ , Prosapogenin A ₂ ,	
	Sanchinoside Rh ₁	
MF:	C ₃₆ H ₆₂ O ₉	
FW:	638.9	HO*
Purity:	≥98%	OH
Supplied as:	A solid	HO.,
Storage:	-20°C	но
Stability:	≥4 years	
Item Origin:	Plant/Panax ginseng	но

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

Laboratory Procedures

Ginsenoside Rh₁ is supplied as a solid. A stock solution may be made by dissolving the ginsenoside Rh₁ in the solvent of choice, which should be purged with an inert gas. Ginsenoside Rh₁ is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of ginsenoside Rh1 in ethanol is approximately 10 mg/ml and approximately 20 mg/ml in DMSO and DMF.

Ginsenoside Rh₁ is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, ginsenoside Rh₁ should first be dissolved in DMF and then diluted with the aqueous buffer of choice. Ginsenoside Rh_1^{-} 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

Ginsenoside Rh1 is a steroid glycoside and saponin that has been found in red ginseng and has diverse biological activities.^{1,2} It inhibits expression of matrix metalloproteinase-1 (MMP-1), MMP-3, and MMP-9 induced by phorbol 12-myristate 13-acetate (PMA; Item No. 10008014) in U87MG and U373MG human astroglioma cells in a concentration-dependent manner.² Ginsenoside Rh₁ (300 μ M) also inhibits PMA-induced invasion and migration of these cell lines in Matrigel™ invasion and wound healing assays, respectively. It inhibits LPS-induced increases in nitrite accumulation, prostaglandin E_2 (PGE,; Item No. 14010) synthesis, and inducible nitric oxide synthase (iNOS) protein levels in RAW 264. $\overline{7}$ cells and inhibits histamine release induced by compound 48/80 (Item No. 22173) in rat peritoneal mast cells (IC₅₀ = 37 μ M).³ Ginsenoside Rh₁ (25 mg/kg, i.p.) inhibits IgE-induced passive cutaneous anaphylaxis reactions in mice by 87%. Ginsenoside Rh1 is also a metabolite of ginsenoside Re (Item No. 15330) and ginsenoside Rg₁ (Item No. 15315) formed by intestinal microflora.^{2,4}

References

- 1. Tam, D.N.H., Truong, D.H., Nguyen, T.T.H., et al. Planta Med. 84(3), 139-152 (2018).
- 2. Jung, J.S., Ahn, J.H., Le, T.K., et al. Neurochem. Int. 63(2), 80-86 (2013).
- 3. Park, E.K., Choo, M.K., Han, M.J., et al. Int. Arch. Allergy. Immunol. 133(2), 113-120 (2004).
- 4. Peng, D., Wang, H., Qu, C., et al. Chin. Med. 7, 2 (2012).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

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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