PRODUCT INFORMATION



Ginsenoside F₁

Item No. 31137

CAS Registry No.: 53963-43-2

Formal Name: $(3\beta,6\alpha,12\beta)$ -3,6,12-trihydroxydammar-

24-en-20-yl β-D-glucopyranoside

Synonyms: GF₁, Panaxoside A Progenin

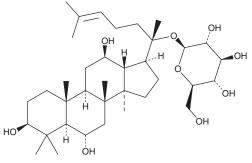
MF: $C_{36}H_{62}O_{9}$ 638.9 FW: **Purity:** ≥98%

Supplied as: A crystalline solid

Storage: -20°C Stability: ≥4 years

Item Origin: Plant/Panax ginseng

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



Laboratory Procedures

Ginsenoside F_1 (GF₁) is supplied as a crystalline solid. A stock solution may be made by dissolving the GF₁ in the solvent of choice, which should be purged with an inert gas. GF_1 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of GF₁ is approximately 10 mg/ml in ethanol and DMF and approximately 30 mg/ml in DMSO.

GF₁ is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, GF₁ should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. GF₁ has a solubility of approximately 0.12 mg/ml in a 1:7 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

GF₁ is a ginsenoside that has been found in P. ginseng and has diverse biological activities.¹⁻⁴ It increases cytotoxic degranulation of natural killer (NK) cells against K562 leukemia cells in vitro when used at a concentration of 10 μM and enhances NK cell-mediated lymphoma clearance in vivo. 1 GF₄ (50 mg/kg) reduces atherosclerotic lesion area, whole body myeloperoxidase (MPO) levels, and aortic root NF-κΒ, TLR4, and LOX-1 levels in ApoE^{-/-} mice.² It increases microvessel density and improves focal cerebral blood perfusion in a rat model of ischemic stroke induced by middle cerebral artery occlusion (MCAO).³ GF₁ also restores spatial working memory, but not context-dependent fear memory, in the APP/PS1 transgenic mouse model of Alzheimer's disease.4

References

- 1. Kwon, H.-J., Lee, H., Choi, G.-E., et al. Ginsenoside F1 promotes cytotoxic activity of NK cells via insulinlike growth factor-1-dependent mechanism. Front. Immunol. 9, 2785 (2018).
- 2. Qin, M., Luo, Y., Lu, S., et al. Ginsenoside F1 ameliorates endothelial cell inflammatory injury and prevents atherosclerosis in mice through A20-mediated suppression of NF-kB signaling. Front. Pharmacol. 8, 953
- 3. Zhang, J., Liu, M., Huang, M., et al. Ginsenoside F1 promotes angiogenesis by activating the IGF-1/IGF1R pathway. Pharmacol. Res. 144, 292-305 (2019).
- 4. Han, J., Oh, J.-P., Yoo, M., et al. Minor ginsenoside F1 improves memory in APP/PS1 mice. Mol. Brain 12(1), 77 (2019).

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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