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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic



PRODUCT INFORMATION



C18 Ganglioside G_{M3}-d₃ (d18:1/18:0-d₃) (ammonium salt, synthetic) Item No. 39226

CAS Registry No.: 2692623-85-9

Formal Name: N-[(1S,2R,3E)-1-[[[O-(N-acetyl- α -neuraminosyl)-(2→3)-O- β -D-galactopyranosyl-(1→4)- β -D-glucopyranosyl]oxy]methyl]-2-hydroxy-3-heptadecen-1-yl]-octadecanamide-18,18,18-d₃, monoammonium salt

Synonyms: C18 G_{M3}-d₃, N- ω -CD₃-Octadecanoyl monosialoganglioside G_{M3}, Monosialoganglioside G_{M3}-d₃, N-Stearoyl Monosialoganglioside G_{M3}-d₃,

MF: C₅₉H₁₀₄D₃N₂O₂₁ • NH₄

FW: 1,201.6

Chemical Purity: ≥95% (Ganglioside G_{M3})

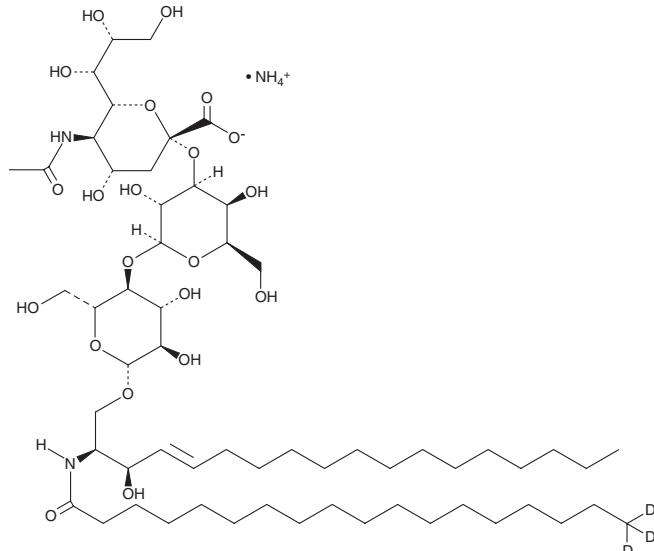
Deuterium

Incorporation: ≥99% deuterated forms (d₁-d₃); ≤1% d₀

Supplied as: A film

Storage: -20°C

Stability: ≥3 years



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

Laboratory Procedures

C18 Ganglioside G_{M3}-d₃ (d18:1/18:0-d₃) (ammonium salt, synthetic) is intended for use as an internal standard for the quantification of ganglioside G_{M3} (Item No. 15587) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

C18 Ganglioside G_{M3}-d₃ (d18:1/18:0-d₃) (ammonium salt, synthetic) is supplied as a solid. A stock solution may be made by dissolving the C18 ganglioside G_{M3}-d₃ (d18:1/18:0-d₃) (ammonium salt, synthetic) in the solvent of choice, which should be purged with an inert gas. C18 Ganglioside G_{M3}-d₃ (d18:1/18:0-d₃) (ammonium salt, synthetic) is soluble in the organic solvent chloroform:methanol (2:1).

Description

Ganglioside G_{M3} is a simple monosialoganglioside that demonstrates both antiproliferative and proapoptotic effects in tumor cells by modulating cell adhesion, proliferation, and differentiation.^{1,2} When used at a concentration of 20 μ M, it suppresses angiogenesis and reduces endothelial cell proliferation and migration by inhibiting VEGFR2 and Akt phosphorylation.¹ Furthermore, ganglioside G_{M3} has been shown to induce the dissociation of the insulin receptor-caveolin-1 complex from lipid microdomains, functioning as an inhibitor of insulin signaling and contributing to insulin resistance.³

References

1. Mukherjee, P., Faber, A.C., Shelton, L.M., et al. *J. Lipid Res.* **49**(5), 929-938 (2008).
2. Seyfried, T.N. and Mukherjee, P. *J. Oncol.* 961243 (2010).
3. Kabayama, K., Sato, T., Saito, K., et al. *Proc. Natl. Acad. Sci. USA* **104**(34), 13678-13683 (2007).

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.

WARRANTY AND LIMITATION OF REMEDY

Buyer agrees to purchase the material subject to Cayman's Terms and Conditions. Complete Terms and Conditions including Warranty and Limitation of Liability information can be found on our website.

CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD
ANN ARBOR, MI 48108 - USA

PHONE: [800] 364-9897
[734] 971-3335

FAX: [734] 971-3640

CUSTSERV@CAYMANCHEM.COM
WWW.CAYMANCHEM.COM