

Product Data Sheet

Product Name: ST1060
Cat. No.: GC79557

Chemical Properties

Cas. No. 1402702-76-4

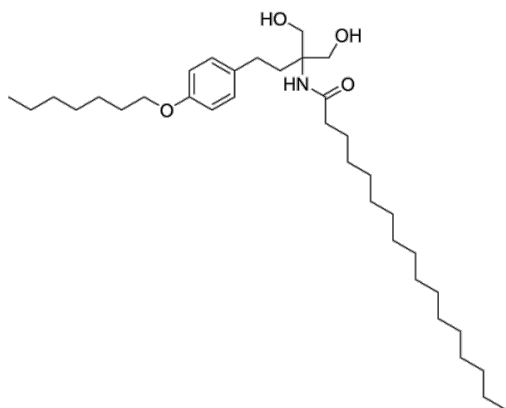
Formula $C_{35}H_{63}NO_4$ M.Wt 561.88

Solubility DMSO: 100 mg/mL (177.97 mM; Need ultrasonic) Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution: ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure



Background

Caution: Product has not been fully validated for medical applications. For research use only.

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ST1060 is a Ceramide synthase 2 (CerS2) inhibitor. ST1060 reduces the levels of long-chain dihydroceramide (dhCer) produced by CerS2 catalysis, with IC₅₀ values of 63.6 μ M and 61.3 μ M for C24:0 and C24:1, respectively. ST1060 is applicable to the research of ceramide metabolism disorders and cervical cancer [1].

In Vitro, ST1060 (preincubated for 10 min, reacted for 30 min) inhibits the synthesis of long-chain dihydroceramides (C24:0-dhCer, C24:1-dhCer) by targeting CerS2 in the microsomal fraction of HCT-116 cells, with IC₅₀ values of 63.6 μ M and 61.3 μ M, respectively [1]. ST1060 (25 μ M; pre-incubated for 10 minutes, reacted for 30 minutes) specifically inhibits CerS2 activity in lysates of HCT-116 cells overexpressing CerS2, as evidenced by a significant reduction in the production of C24:0 and C24:1-dhCer [1]. ST1060 (30 μ M) significantly reduces the levels of long-chain dihydroceramides (C24:0-dhCer, C24:1-dhCer) in HeLa cervical cancer cells, confirming its targeted action on CerS2 [1]. ST1060 (10-70 μ M; 48 h) inhibits C14:0-ceramide synthesis in HeLa cervical cancer cells in a concentration-dependent manner, with an IC₅₀ of 45.1 μ M after 48 h of treatment, while increasing the level of long-chain ceramides [1]. ST1060 (0-80 μ M; 48 h) increases the cell viability of HCT-116 colon cancer cells and induces their proliferation in a concentration-dependent manner following 48 h of treatment [1].

References:

[1]. Schiffmann S, et al. Inhibitors of specific ceramide synthases. Biochimie. 2012 Feb;94(2):558-65.

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