

Product Data Sheet

Product Name: Nav1.2-IN-2

Cat. No.: GC79520

Chemical Properties

Cas. No. 7469-49-0

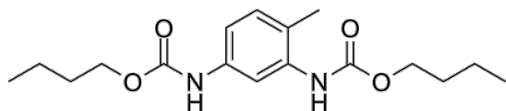
Formula $C_{17}H_{26}N_2O_4$ M.Wt 322.4

Solubility DMSO: 4.17 mg/mL (12.93 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



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

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Background

Nav1.2-IN-2 is a Nav1.2 inhibitor with a human IC₅₀ of 0.18 μ M. Nav1.2-IN-2 preferentially binds to the inactivated state of Nav1.2, reduces window current, suppresses neuronal depolarization and action potential generation. Nav1.2-IN-2 suppresses Veratridine -induced Ca²⁺ influx. Nav1.2-IN-2 can be used for the research of epilepsy [1].

In Vitro, Nav1.2-IN-2 (Compound 1) (0.1-10 μ M; 15 min) potently suppresses spontaneous synchronized Ca²⁺ oscillation amplitude and frequency in primary cultured neocortical neurons with IC₅₀ values of 0.38 μ M and 0.88 μ M, respectively [1]. Nav1.2-IN-2 (0.1-10 μ M; 15 min) potently inhibits Veratridine-induced Ca²⁺ influx in primary cultured neocortical neurons with an IC₅₀ of 1.89 μ M [1]. Nav1.2-IN-2 exhibits state-dependent inhibition of Nav1.2 in transiently transfected HEK-293 cells, shifting steady-state inactivation hyperpolarization by 10.99 mV and inhibiting the inactivated state with an IC₅₀ of 0.18 μ M, while having no effect on steady-state activation [1].

References:

[1]. Xu J, et al. Aniline analogues as synchronized Ca²⁺ oscillation modulators: structure-activity relationship and mechanism. Eur J Med Chem. 2026;301:118267.

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