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DATASHEET

GV-58

Product overview

 Name
 GV-58

 Cat No
 HB3920

 Purity
 >97%

Description Selective N- and P/Q-type Ca²⁺-channel agonist

Biological Data

Biological description Selective N- and P/Q-type Ca2+-channel agonist. These Ca2+-channels regulate transmitter release in

synapses. Potential lead compound for a variety of disorders that result in neuromuscular weakness.

Solubility & Handling

Storage instructions +4°C

Storage of solutions Prepare and use solutions on the same day if possible. Store solutions at -20 °C for up to one month if

storage is required. Equilibrate to RT and ensure the solution is precipitate free before use.

Shipping Conditions

Important

Stable for ambient temperature shipping. Follow storage instructions on receipt.

This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not

for human or veterinary use.

Chemical Data

Chemical name Molecular Weight Chemical structure (2R)-2-[[6-[[(5-Methyl-2-thienyl)methyl]amino]-9-propyl-9H-purin-2-yl]amino]-1-butanol

H₃C H

Molecular Formula $C_{18}H_{26}N_6OS$ CAS Number1402821-41-3

InChiKey DPTXJOUVBMUSGY-CYBMUJFWSA-N

374.5

Appearance White to off-white solid

References

Potentiation of neuromuscular transmission by a small molecule calcium channel gating modifier improves motor function in a severe spinal muscular atrophy mouse model.

Ojala KS et al (2023) Human molecular genetics 32 **PubMedID** 36757138

Evaluation of a novel calcium channel agonist for therapeutic potential in Lambert-Eaton myasthenic syndrome.

Tarr TB et al (2013) The Journal of neuroscience : the official journal of the Society for Neuroscience 33

PubMedID 23785168

Activation of Voltage-Gated Na(+) Current by GV-58, a Known Activator of Ca(V) Channels.

Cho HY et al (2022) Biomedicines 10

PubMedID 35327523