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Zuschläge

- Mindermengenzuschlag
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SZABO-SCANDIC Handels GmbH

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

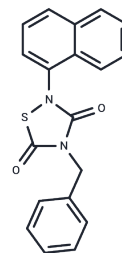
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Tideglusib

Chemical Properties

CAS No. : 865854-05-3
 Formula: C₁₉H₁₄N₂O₂S
 Molecular Weight: 334.39
 Appearance: no data available
 Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



Biological Description

Description	Tideglusib (NP031112), a non-ATP competitive inhibitor of glycogen synthase kinase 3 (GSK-3), is with anti-inflammatory and neuroprotective activities.
Targets(IC50)	GSK-3
Kinase Assay	[35S]Tideglusib (207 Bq/nmol) at 55 μ M is incubated with 5 μ M GSK-3 β for 1 h at 25°C in 315 μ L of 50 mM Tris-HCl, pH 7.5, containing 150 mM NaCl and 0.1 mM EGTA. The incubation is extended for another 30 min after having added 35 μ L of the same buffer with or without 100 mM DTE. Samples are then processed in three different ways. First, an aliquot of 125 μ L of each sample is mixed with 375 μ L of 8 M GdnHCl in Water and heated at 80°C for 5 min. A second aliquot of 125 μ L is diluted up to 500 μ L with Water and left at room temperature for 5 min. In both cases, the free drug is removed afterwards by gel filtration through Sephadex G-25, and the amount of bound drug is determined by liquid scintillation counting on a 1450-MicroBeta TriLux counter. Finally, a third 40 μ L aliquot of each original sample is mixed with 10 μ L of denaturing electrophoresis sample buffer without reducing agents, and 35 μ L of this mixture is loaded onto a 10% polyacrylamide gel and subjected to SDS-PAGE (again in the absence of reducing agents except for the DTE already included in the corresponding sample), followed by fluorography of the dried gel[1].

Solubility Information

Solubility	Ethanol: < 1 mg/mL (insoluble or slightly soluble), DMSO: >15 mg/mL (44.9 mM), < 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.9905 mL	14.9526 mL	29.9052 mL
5 mM	0.5981 mL	2.9905 mL	5.981 mL
10 mM	0.2991 mL	1.4953 mL	2.9905 mL
50 mM	0.0598 mL	0.2991 mL	0.5981 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Dou X, Sun Q, Xu G, et al. Discovery of 2-(furan-2-ylmethylene) hydrazine-1-carbothioamide derivatives as novel inhibitors of SARS-CoV-2 main protease. European Journal of Medicinal Chemistry. 2022: 114508

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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Tel: 781-999-4286 E_mail: info@targetmol.com Address: 36 Washington Street, Wellesley Hills, MA 02481