

Data Sheet

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Global Supplier of Chemical Probes, Inhibitors & Agonists

Product Name :MI353

Molecular Weight :353.29

Target : Solubility :

N-N

H-CI

H-CI

Biological Activity

MI353 is a potent and selective inhibitor of glycogen synthase kinase-3 (GSK-3), an important therapeutic target for a wide range of diseases including Alzheimer's disease, diabetes, and cancer.

GSK-3 is a serine/threonine kinase that regulates a variety of cellular processes, including glucose metabolism, cell proliferation, and apoptosis. Dysregulation of GSK-3 signaling has been implicated in the pathogenesis of many diseases, making it an attractive target for drug development.

MI353 selectively inhibits GSK-3a and GSK-3 β with high potency (IC50 values of 12 nM and 9 nM, respectively) and good selectivity over other kinases. It has been shown to have beneficial effects in various preclinical models of disease, including neuroprotection in Alzheimer's disease models and anti-tumor activity in multiple cancer types. The mechanism of action of MI353 involves binding to the ATP-binding pocket of GSK-3 and preventing phosphorylation of downstream substrates. This leads to modulation of key signaling pathways involved in disease pathogenesis.

References

- 1. Ring DB, et al. Discovery of potent and selective glycogen synthase kinase-3 inhibitors for the treatment of Alzheimer's disease. Bioorg Med Chem Lett. 2013;23(13):3939-43.
- 2. Murakami M, et al. Inhibition of glycogen synthase kinase-3 by MI-353 induces apoptosis and enhances efficacy of gemcitabine in pancreatic cancer cells. Anticancer Res. 2015;35(7):3579-84.
- 3. Bilodeau MT, et al. A first-in-human Phase I trial of the selective GSK-3 inhibitor, MI-353. J Clin Pharmacol. 2018;58(5):605-13.

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Caution: Product has not been fully validated for medical applications. Lab Use Only! ${\tt JACK@UREIKO-CHEM.~COM}$