

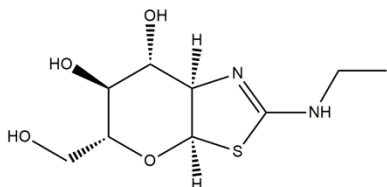
# Thiamet G

12/19

**ALTERNATE NAMES:** 2-ethylimino-5-(hydroxymethyl)-1,3a,5,6,7,7a-hexahydropyrano[3,2-d][1,3]thiazole-6,7-diol; 2-(ethylamino)-3aR,6S,7R,7aR-tetrahydro-5R-(hydroxymethyl)-5H-pyrano[3,2-d]thiazole-6,7-diol; O-GlcNAcase Inhibitor

**CATALOG #:** B2959-5 5 mg  
B2959-25 25 mg

**STRUCTURE:**



**MOLECULAR FORMULA:** C<sub>9</sub>H<sub>16</sub>N<sub>2</sub>O<sub>4</sub>S

**MOLECULAR WEIGHT:** 248.3

**CAS NUMBER:** 1009816-48-1

**APPEARANCE:** Solid

**PURITY:** 98%

**SOLUBILITY:** ~0.5 mg/ml in Ethanol  
~20 mg/ml in DMSO and DMF

**DESCRIPTION:** Thiamet G is a potent and selective inhibitor of O-GlcNAcase with a K<sub>i</sub> value of 21 nM. It increases cellular O-GlcNAc-modified protein levels with an EC<sub>50</sub> of 30 nM and blocks phosphorylation of tau protein both in cultured PC-12 cells and in rats (200 mg/kg/day). It is the first highly potent O-GlcNAcase inhibitor known to be orally bioavailable and can effectively cross the blood brain barrier. It may potentially block pathological hyperphosphorylation of tau in Alzheimer's disease.

**STORAGE TEMPERATURE:** -20°C. Store in the dark. Product is light sensitive. Protect from air. Store under desiccating conditions.

**HANDLING:** Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.

**REFERENCE:** Yuzwa, S.A., Macauley, M.S., Heinonen, J.E., et al. A potent mechanism-inspired O-GlcNAcase inhibitor that blocks phosphorylation of tau in vivo. *Nature Chemical Biology* 4(8), 483-490 (2008).

**RELATED PRODUCTS:**

Congo Red (Cat. No. 2588)  
 Beta-Secretase Inhibitor II (Cat. No. 7502)  
 CHIR-98014 (Cat. No. 2595)  
 10Z-Hymenialdisine (Cat. No. 2212)  
 GSK-3039294 (Cat. No. B2374)

**DISCLAIMER:** *FOR RESEARCH USE ONLY! Not to be used on humans.*