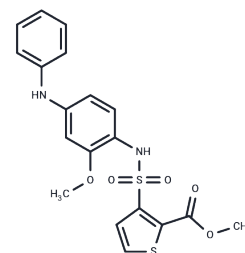


GSK0660

## Chemical Properties

CAS No. : 1014691-61-2  
 Formula: C<sub>19</sub>H<sub>18</sub>N<sub>2</sub>O<sub>5</sub>S<sub>2</sub>  
 Molecular Weight: 418.49  
 Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year  
 Actual storage temperature shall be subject to the COA.



## Biological Description

Description	GSK0660 is an antagonist and inverse agonist of PPAR $\beta/\delta$ .
Targets(IC50)	PPAR
Kinase Assay	Human astrocytoma cells, 1321N1, are grown to stably express rat P2X7, human P2X4, P2X2a, P2X2/3, P2X1, P2Y1 and P2Y2 recombinant receptors. Agonist, BzATP, 2,3-O-(4-benzoylbenzoyl)-ATP or ATP-induced changes in intracellular Ca <sup>2+</sup> concentrations are assessed in all of the cell lines using the Ca <sup>2+</sup> chelating dye, Fluo-4, in conjunction with a Fluorometric Imaging Plate Reader. The cells are plated out the day before the experiment onto poly-D-lysine-coated black 96 well plates. After the agonist addition, changes in intracellular Ca <sup>2+</sup> concentrations are recorded, per second, for 3 min. Ligands are tested at 11 half-log concentrations from 10 <sup>-10</sup> to 10 <sup>-4</sup> M. BzATP or ATP concentrations corresponds to the EC <sub>70</sub> values for each receptor to enable comparison of antagonist potencies across the multiple P2 receptor subtypes. A 438079 is added to the cell plate and fluorescence data are collected for 3 min before the addition of agonist, subsequently, data are then collected for another 2 min. The pEC <sub>50</sub> or pIC <sub>50</sub> values are derived from a single curve fit.

## Solubility Information

Solubility	Ethanol: 4.19 mg/mL (10.01 mM), Sonication is recommended. DMSO: 117.5 mg/mL (280.77 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
In vivo Formulation	10% DMSO+90% (20% SBE- $\beta$ -CD in Saline): < 10 mg/mL (23.9 mM), Lower concentrations may be soluble, but exact solubility limit is unknown. 10% DMSO+40% PEG300+5% Tween 80+45% Saline: < 10 mg/mL (23.9 mM), Lower concentrations may be soluble, but exact solubility limit is unknown. 10% DMSO+90% Saline: < 10 mg/mL (23.9 mM), Lower concentrations may be soluble, but exact solubility limit is unknown. 10% DMSO+90% Corn oil: < 10 mg/mL (23.9 mM), Lower concentrations may be soluble, but exact solubility limit is unknown. Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and

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In vivo Formulation	<i>used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i>
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### Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.3895 mL	11.9477 mL	23.8954 mL
5 mM	0.4779 mL	2.3895 mL	4.7791 mL
10 mM	0.239 mL	1.1948 mL	2.3895 mL
50 mM	0.0478 mL	0.239 mL	0.4779 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Savage SR, et al. Mol Vis. 2015 May 20;21:568-76.

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