

Product Data Sheet

GSK-2793660

Cat. No.: HY-112318A CAS No.: 1613458-78-8 Molecular Formula: $C_{20}H_{28}CIN_3O_3$ Molecular Weight: 393.91

Target: Cathepsin

Pathway: Metabolic Enzyme/Protease

Storage: -20°C, sealed storage, away from moisture

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (253.87 mM; Need ultrasonic)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.5387 mL	12.6933 mL	25.3865 mL
	5 mM	0.5077 mL	2.5387 mL	5.0773 mL
	10 mM	0.2539 mL	1.2693 mL	2.5387 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (6.35 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.35 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.35 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	GSK-2793660 is an orally active and irreversible inhibitor of Cathepsin C (CTSC). GSK-2793660 can be used for the research of bronchiectasis $^{[1][2]}$.
IC ₅₀ & Target	Cathepsin C
In Vivo	GSK-2793660 is a cathepsin C (also known as dipeptidyl peptidase I enzyme) inhibitor for the reaearch of cystic fibrosis, non-cystic fibrosis bronchiectasis, anti-neutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis and bronchiectasis ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

[1]. Szcześniak P, et al. The Synthesis of α,α-Disubstituted α-Amino Acids via Ichikawa Rearrangement. J Org Chem. 2016;81(3):1057-1074.				
[2]. Miller BE, et al. Epithelial desquamation observed in a phase I study of an oral cathepsin C inhibitor (GSK2793660). Br J Clin Pharmacol. 2017;83(12):2813-2820.				
(Caution: Product has not been fully validated for medical applications. For research use only.			
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