



MK-8617

Catalog No: tcsc0020731

Observed Molecular Weight:

443.45

Available Sizes	
Size: 5mg	
Size: 10mg	
Size: 25mg	
Size: 50mg	
Size: 100mg	
Size: 200mg	
Specifications	
CAS No: 1187990-87-9	
Formula: C ₂₄ H ₂₁ N ₅ O ₄	
Pathway: Metabolic Enzyme/Protease	
Target: HIF/HIF Prolyl-Hydroxylase	
Purity / Grade: >98%	
Solubility: DMSO : 6 mg/mL (13.53 mM; Need ultrasonic)	



Product Description

MK-8617 is an orally active pan-inhibitor of hypoxia-inducible factor prolyl hydroxylase 1-3 (**HIF PHD1-3**) with an IC_{50} of 1 nM for **PHD2**.

IC50 & Target: IC50: 1 nM (PHD2)[1]

In Vitro: MK-8617 is an orally active pan-inhibitor of hypoxia-inducible factor prolyl hydroxylase 1-3 (HIF PHD1-3) with an IC $_{50}$ of 1 nM for PHD2. MK-8617 is not a significant inhibitor of the cytochrome p450 enzymes in vitro (IC $_{50}$), CYP1A2, 3A4, 2B6, 2C9, 2C19, or 2D6, >60 μ M, and is a moderate reversible inhibitor of CYP2C8 at 1.6 μ M in vitro. The IC $_{50}$ of MK-8617 is determined for factor inhibiting HIF (FIH) to be 18 μ M $^{[1]}$.

In Vivo: Tritiated MK-8617 exhibits minimal metabolic turnover in liver microsomes from rat, dog, and monkey ([1].

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!