



Simeprevir

Catalog No: tcsc0338

Available Si	zes		
Size: 5mg			
Size: 10mg			
Size: 50mg			
Size: 100mg			
Specification	ns		
CAS No: 923604-59-5			
Formula: C ₃₈ H ₄₇ N ₅ O ₇ S ₂			
Pathway: Metabolic Enzyme/Prof	tease;Anti-infection		
Target: HCV Protease;HCV			
Purity / Grade: >98%			
Solubility: DMSO: 14.29 mg/mL ((19.05 mM; Need ultrasonic)		
Alternative Names: TMC435			
Observed Molecular 749.94	Weight:		



Product Description

Simeprevir is a potent **HCV NS3/4A protease** inhibitor, and inhibits HCV replication with EC_{50} of 8 nM.

IC50 & Target: EC50: 8 nM

In Vitro: In Huh7-Luc cells, antiviral activity of simeprevir (TMC435350) is dose dependent, and the EC $_{50}$ and EC $_{90}$ values determined for TMC435350 are 8 nM and 24 nM, respectively. Inhibition of TMC435350 on NS3/4A protease is time dependent, and the overall K $_{i}$ s are estimated to be 0.5 nM for genotype 1a and 0.4 nM for genotype 1b, respectively^[1]. TMC435350 is a potent inhibitor of HCV NS3/4A protease (K_{i} =0.36 nM) and viral replication (replicon EC $_{50}$ =7.8 nM)^[2].

In Vivo: In rats, TMC435350 (40 mg/kg, p.o.) is extensively distributed to the liver and intestinal tract (tissue/plasma area under the concentration-time curve ratios of >35), and the absolute bioavailability is $44\%^{[1]}$.

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