



## Simeprevir

Catalog No: tcsc0338

Available Si	zes		
Size: 5mg			
Size: 10mg			
Size: 50mg			
Size: 100mg			
Specification	ns		
<b>CAS No:</b> 923604-59-5			
<b>Formula:</b> C <sub>38</sub> H <sub>47</sub> N <sub>5</sub> O <sub>7</sub> S <sub>2</sub>			
<b>Pathway:</b> Metabolic Enzyme/Prof	tease;Anti-infection		
<b>Target:</b> HCV Protease;HCV			
Purity / Grade: >98%			
<b>Solubility:</b> DMSO: 14.29 mg/mL (	(19.05 mM; Need ultrasonic)		
<b>Alternative Names:</b> TMC435			
<b>Observed Molecular</b> 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<sup>[1]</sup>. TMC435350 is a potent inhibitor of HCV NS3/4A protease ( $K_{i}$ =0.36 nM) and viral replication (replicon EC $_{50}$ =7.8 nM)<sup>[2]</sup>.

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!