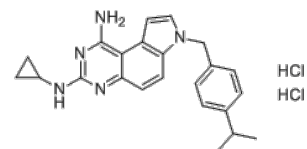


**Product Name** : SCH 79797 dihydrochloride  
**Cat. No.** : PC-61305  
**CAS No.** : 1216720-69-2  
**Molecular Formula** : C<sub>23</sub>H<sub>25</sub>N<sub>5</sub>·2HCl  
**Molecular Weight** : 444.41  
**Target** : Protease-activated Receptor (PAR)  
**Solubility** : 10 mM in DMSO



## Biological Activity

SCH79797 dihydrochloride is a potent, selective competitive and nonpeptide **PAR-1** antagonist with IC<sub>50</sub> of 70 nM. SCH79797 inhibits α-thrombin- and haTRAP-induced aggregation of human platelets, but does not inhibit human platelet aggregation induced by the tethered ligand agonist for PAR-4, γ-thrombin, ADP, or collagen. SCH79797 also blocks angiogenesis associated with endothelial cell growth suppression and induction of apoptosis.

## References

- Ahn HS, et al. *Biochem Pharmacol.* 2000 Nov 15;60(10):1425-34.  
Ahn HS, et al. *Bioorg Med Chem Lett.* 1999 Jul 19;9(14):2073-8.  
Zania P, et al. *J Pharmacol Exp Ther.* 2006 Jul;318(1):246-54.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

E-mail: tech@probechem.com