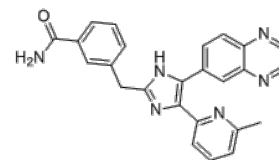


**Product Name** : IN-1130  
**Cat. No.** : PC-70054  
**CAS No.** : 868612-83-3  
**Molecular Formula** : C<sub>25</sub>H<sub>20</sub>N<sub>6</sub>O  
**Molecular Weight** : 420.47  
**Target** : TGF beta Receptor (TGFBR)  
**Solubility** : 10 mM in DMSO



## Biological Activity

IN-1130 is a potent, highly selective **ALK5** inhibitor, inhibits the purified kinase domain of ALK5-mediated Smad3 phosphorylation with an IC<sub>50</sub> of 5.3 nM.

IN-1130 is highly selective in a panel of 27 serine/threonine and tyrosine kinases including p38 $\alpha$ .

IN-1130 decreases levels of TGF-beta1 mRNA, type I collagen mRNA, and pSmad2, compared to UUO control rats.

IN-1130 inhibits renal fibrosis in obstructive nephropathy in vivo models.

## References

Moon JA, et al. *Kidney Int.* 2006 Oct;70(7):1234-43.

Ryu JK, et al. *J Sex Med.* 2009 May;6(5):1284-96.

Park CY, et al. *Cancer Lett.* 2014 Aug 28;351(1):72-80.

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

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