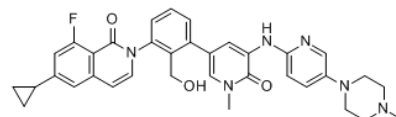


**Product Name** : RN-486  
**Cat. No.** : PC-45403  
**CAS No.** : 1242156-23-5  
**Molecular Formula** : C<sub>35</sub>H<sub>35</sub>FN<sub>6</sub>O<sub>3</sub>  
**Molecular Weight** : 606.6892  
**Target** : BTK  
**Solubility** : DMSO: 24 mg/mL



## Biological Activity

RN-486 is a potent and selective **BTK** inhibitor with IC<sub>50</sub> of 4 nM in enzymatic assays.

RN-486 shows high selectivity against a panel of 369 kinases.

RN-486 blocks Fcε receptor cross-linking-induced degranulation in mast cells (IC<sub>50</sub>=2.9 nM) and Fcγ receptor engagement-mediated TNFα production in monocytes (IC<sub>50</sub>=7.0 nM).

RN-486 shows robust anti-inflammatory and bone-protective effects in mouse CIA and rat adjuvant-induced arthritis (AIA) models.

## References

Xu D, et al. *J Pharmacol Exp Ther*. 2012 Apr;341(1):90-103.

Mina-Osorio P, et al. *Arthritis Rheum*. 2013 Sep;65(9):2380-91.

Lou Y, et al. *J Med Chem*. 2015 Jan 8;58(1):512-6.

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

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