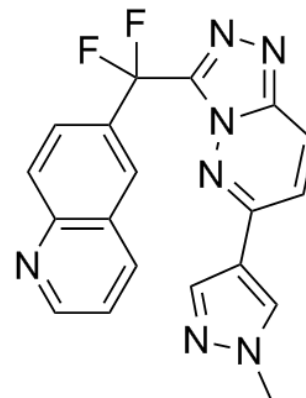


**Product Name** : JNJ-38877605  
**Cat. No.** : PC-45841  
**CAS No.** : 943540-75-8  
**Molecular Formula** : C<sub>19</sub>H<sub>13</sub>F<sub>2</sub>N<sub>7</sub>  
**Molecular Weight** : 377.3502  
**Target** : c-Met (HGFR)  
**Solubility** : DMSO: ≥ 30 mg/mL



## Biological Activity

JNJ-38877605 is a potent, selective, ATP-competitive inhibitor of catalytic activity **c-Met** with IC<sub>50</sub> of 4 nM. JNJ-38877605 exhibits >600-fold selectivity (cFMS IC<sub>50</sub>=2.6 μM, the next potentially inhibited kinase) against a panel of 250 diverse tyrosine and serine-threonine kinases. JNJ-38877605 inhibits Met phosphorylation associated with dose-dependent tumor growth inhibition in tumor xenograft models; orally bioavailable.

## References

- Peter King, et al. DOI: 10.1158/1538-7445.AM10-3628 Published April 2010  
Lolkema MP, et al. *Clin Cancer Res.* 2015 May 15;21(10):2297-2304.  
Galimi F, et al. *Clin Cancer Res.* 2011 May 15;17(10):3146-56.

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

E-mail: tech@probechem.com