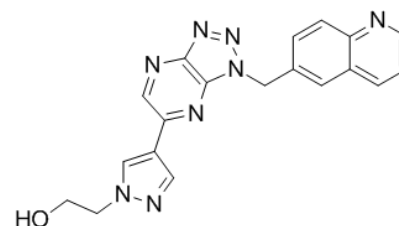


**Product Name** : PF-04217903  
**Cat. No.** : PC-42832  
**CAS No.** : 956905-27-4  
**Molecular Formula** : C<sub>19</sub>H<sub>16</sub>N<sub>8</sub>O  
**Molecular Weight** : 372.3833  
**Target** : c-Met (HGFR)  
**Solubility** : 10 mM in DMSO



## Biological Activity

PF-04217903 is a potent, selective, ATP-competitive **c-Met** kinase inhibitor with  $K_i$  of 4.8 nM, inhibits HGF-stimulated tyrosine phosphorylation of wild type c-Met with  $IC_{50}$  of 7.3 nM in cell-based assays. PF-04217903 shows similar potency for inhibition of c-Met phosphorylation in mIMCD3 mouse epithelial cells ( $IC_{50}$ =6.9 nM) and MDCK cell ( $IC_{50}$ =9.5 nM). PF-04217903 also exhibits inhibitory activities against c-Met mutations including V1092I, H1094R, M1250T, R988C, and T11010I ( $IC_{50}$ =530 nM), but has little to no activity against mutants Y1230C and Y1235D. PF-04217903 inhibits tumor cell proliferation, survival, migration/invasion both in vitro and in vivo.

## References

- Zou HY, et al. *Mol Cancer Ther.* 2012 Apr;11(4):1036-47.  
Sennino B, et al. *Cancer Res.* 2013 Jun 15;73(12):3692-703.  
Cui JJ, et al. *J Med Chem.* 2012 Sep 27;55(18):8091-109.  
Yeh I, et al. *Nat Commun.* 2015 May 27;6:7174.

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

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