## PF-06737007

Cat. No.: HY-112437

CAS No.: 1863905-38-7 Molecular Formula:  $C_{25}H_{28}F_4N_2O_6$ 

Molecular Weight: 528.49

Target: Trk Receptor

Pathway: Neuronal Signaling; Protein Tyrosine Kinase/RTK

Storage: Powder -20°C 3 years

4°C 2 years

-80°C In solvent 6 months

> -20°C 1 month

## **SOLVENT & SOLUBILITY**

DMSO : ≥ 100 mg/mL (189.22 mM) In Vitro

\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.8922 mL	9.4609 mL	18.9218 mL
	5 mM	0.3784 mL	1.8922 mL	3.7844 mL
	10 mM	0.1892 mL	0.9461 mL	1.8922 mL

Please refer to the solubility information to select the appropriate solvent.

## **BIOLOGICAL ACTIVITY**

Description  $PF-06737007\ is\ a\ potent\ pan-Trk\ inhibitor\ in\ cell-based\ assays\ with\ IC_{50}s\ of\ 7.7\ nM,\ 15\ nM\ and\ 3.9\ nM\ for\ TrkA,\ TrkB\ and\ TrkC,$ respectively<sup>[1]</sup>. Anti-hyperalgesic effect<sup>[1]</sup>.

TrkA TrkB TrkC IC<sub>50</sub> & Target

> 3.9 nM (IC<sub>50</sub>, in cell-based assays) 7.7 nM (IC<sub>50</sub>, in cell-based 15 nM (IC<sub>50</sub>, in cell-based

assays) assays)

In Vitro PF-06737007 (Compound 13b) exhibits superb Trk selectivity with >95% inhibition of TrkA with no other kinase being

inhibited by >40% at 1  $\mu$ M<sup>[1]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

1]. Bagal SK, et al. Discovery 8800.	of Potent, Selective, and Peripl	nerally Restricted Pan-Trk Kinas	se Inhibitors for the Treatment of F	Pain. J Med Chem. 2018 Aug 9;6	1(15):6779-
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