

PF-05089771, Nav1.7 inhibitor (selective)

Catalog	Unit
TBI3990-5MG	5 mg
TBI3990-25MG	25 mg

Product Details

Formal Name: 4-[2-(5-Amino-1H-pyrazol-4-yl)-4-chlorophenoxy]-5-chloro-2-fluoro-N-(1,3-thiazol-4-yl)-4-chlorophenoxy]

yl)benzenesulfonamide **Alternate Names:** PF771

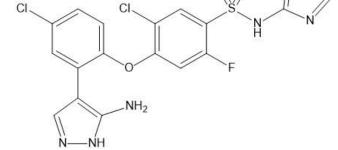
 $\textbf{Molecular Formula:} \ C_{18}H_{12}Cl_2FN_5O_3S_2$

Formula Weight: 500.34 **CAS Number:** 1235403-62-9

Purity: >98% **Formulation:** powder

Solubility: Soluble in DMSO (up to 25 mg/ml)

Storage: -20° C Stability: ≥ 1 year.



Applications

Nav1.7 inhibitor (selective)

Functions

Potent and selective inhibitor of the voltage gated sodium channel 1.7 (Nav1.7). It is a state-dependent inhibitor with IC50 = 11 nM for the half-inactivated channels and IC50 ~ 10 μ M for resting channels. Equipotent for human, monkey, dog, and mouse channels but 15x less potent for rat channels. Interacts equally with fast and slow inactivated Nav1.7 channels.

Application Procedures

First dissolved in DMSO (up to 25 mg/ml), then diluted to aqueous buffer. Solutions in DMSO may be stored at -20° C for up to 3 months.

For research use only.