

Catalog	Unit
TBI4726-5MG	5 mg
TBI4726-25MG	25 mg

### Product Details

**Formal Name:** 6-Fluoro-N-[(5-fluoro-2-methoxypyridin-3-yl)methyl]-5-[(5-methyl-1H-pyrrolo[2,3-b]pyridine-3-yl)methyl]pyridine-2-amine

**Molecular Formula:** C<sub>21</sub>H<sub>19</sub>F<sub>2</sub>N<sub>5</sub>O

**Formula Weight:** 395.41

**CAS Number:** 1303420-67-8

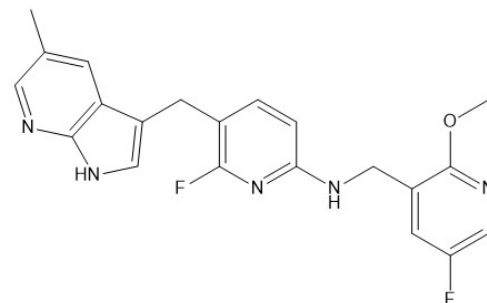
**Purity:** >98%

**Formulation:** powder

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

**Storage:** -20°C

**Stability:** ≥ 2 years.



### Applications

CSF1R inhibitor (highly selective)

### Functions

Highly selective (>20 fold over KIT and FLT3, >60 fold over 200 other kinases) and brain-penetrant inhibitor of colony-stimulating factor 1 receptor (CSF1R; IC<sub>50</sub> = 16 nM). It prevented plaque formation in 5xFAD and 3xTg mouse models of Alzheimer's disease via elimination of microglia in a CSF1R-dependent manner. PLX5622 showed efficacy in a mouse neuropathic pain model via reduction of CD86+ macrophages resulting in reduced expression of pro-inflammatory cytokines. It also was able to ameliorate peripheral neuropathy in aging mice. PLX5622 displayed neuroprotective effects during the chronic phase of a traumatic brain injury mouse model. PLX5622 has also been shown to affect myeloid and lymphoid compartments, indicating that its effects are not limited to microglia and include peripheral immune cells.

### Application Procedures

First dissolved in DMSO (up to at least 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.**