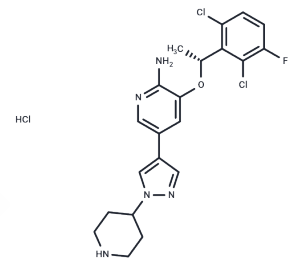


## Crizotinib hydrochloride

## Chemical Properties

CAS No. :	1415560-69-8
Formula:	C <sub>21</sub> H <sub>23</sub> Cl <sub>3</sub> FN <sub>5</sub> O
Molecular Weight:	486.8
Storage:	Store at low temperature Powder: -20°C for 3 years   In solvent: -80°C for 1 year <i>Actual storage temperature shall be subject to the COA.</i>



## Biological Description

Description	Crizotinib hydrochloride (PF-02341066 hydrochloride) is a novel inhibitor of anaplastic lymphoma kinase and c-Met, with IC <sub>50</sub> values of 20 nM and 8 nM, respectively.
Targets(IC <sub>50</sub> )	ALK, Autophagy, c-Met/HGFR, ROS, ROS Kinase
In vitro	Crizotinib hydrochloride (PF-2341066) potently inhibited NPM-ALK phosphorylation in Karpas299 or SU-DHL-1 ALCL cells (mean IC <sub>50</sub> value, 24 nmol/L). In biochemical and cellular screens, PF-2341066 was shown to be selective for c-Met and ALK at pharmacologically relevant concentrations across a panel of >120 diverse kinases. PF-2341066 potently inhibited cell proliferation, which was associated with G(1)-S-phase cell cycle arrest and induction of apoptosis in ALK-positive ALCL cells (IC <sub>50</sub> values, approximately 30 nmol/L) but not ALK-negative lymphoma cells[1].

## Solubility Information

Solubility	DMSO: 10 mg/mL (20.54 mM), Sonication is recommended. H <sub>2</sub> O: 50 mg/mL (102.71 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	2.0542 mL	10.2712 mL	20.5423 mL
5 mM	0.4108 mL	2.0542 mL	4.1085 mL
10 mM	0.2054 mL	1.0271 mL	2.0542 mL
50 mM	0.0411 mL	0.2054 mL	0.4108 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Christensen JG, et al. Cytoreductive antitumor activity of PF-2341066, a novel inhibitor of anaplastic lymphoma kinase and c-Met, in experimental models of anaplastic large-cell lymphoma. *Mol Cancer Ther.* 2007, 6(12 Pt 1), 3314-3322.

Cui JJ, et al. Structure based drug design of crizotinib (PF-02341066), a potent and selective dual inhibitor of mesenchymal-epithelial transition factor (c-MET) kinase and anaplastic lymphoma kinase (ALK). *J Med Chem.* 2011 Sep 22;54(18):6342-63.

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