#### Crizotinib

# REAGENTS DIRECT

## 5 mg

## For research purposes only

Crizotinib is a potent ATP competitive inhibitor of c-MET and anaplastic lymphoma kinase (ALK). It has been shown to display antitumor efficacy in multiple tumor models and has been shown to inhibit c-MET dependent proliferation, migration and invasion of tumor cells *in vitro*. Crizotinib is selective for c-MET and ALK against over 120 different kinases.

#### **TECHNICAL INFORMATION**

Other Names: PF-02341066, PF-1066, PF-2341066

Chemical Formula: C<sub>21</sub>H<sub>22</sub>Cl<sub>2</sub>FN<sub>5</sub>O

**CAS Number:** 877399-52-5

Molecular Weight: 450.34

**Purity: >98%** 

Appearance: a crystalline solid

**Solubility: DMSO** 

#### STORAGE AND HANDLING

**Storage:** Store at 4°C and protected from light. Following reconstitution, store aliquots at -20°C.

**Stability:** Stock solutions stable at -20°C for up to 2 years.

**Shipping Conditions:** Shipped at room temperature.

### **PRODUCT USE**

Soluble in DMSO at 25 mg/ml. If precipitate is observed, vortex for 5 minutes. For most cells, the maximum tolerance to DMSO is less than 0.5%.

#### REFERENCES

- Cui et al. (2011) 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. 54(18):6342-63.
- Rodig et al. (2010) Crizotinib, a small-molecule dual inhibitor of the c-Met and ALK receptor tyrosine kinases. Curr Opin Investig Drugs. 11 (12):1477-90.
- 3. Gerber et al. (2010) ALK inhibition for non-small cell lung cancer: from discovery to therapy in record time. Cancer Cell. 18(6):548-51.

