

## Catalog # 10-4813 Quizartinib 950769-58-1

AC220; 1-(5-(t-Butyl)isoxazol-3-yl)-3-(4-(7-(2-morpholinoethoxy)benzo[d]imidazo[2,1-b]thiazol-2-yl)phenyl)urea Lot #FBS1092

Quizartinib is a potent and selective inhibitor of FLT3 ( $K_d = 1.6$ nM,  $IC_{50} = 0.56$  nM MV4-11 cells). It is in clinical trials for treatment of Acute Myelogenous Leukemia (AML). Quizartinib priming resulted in prevention of myelosuppression in mice treated with 5-FU and Gemcitabine. Quizartinib showed significant reversal of ABCG2-mediated multidrug resistance (@ 3  $\mu$ M) via antagonism of drug efflux function.  $^{5,6}$ 

- 1) Chao et al. (2009) Identification of N-(5-tert-butyl-isoxazol-3-yl)-N'-{4-[7-(2-morpholin-4-yl-ethoxy)imidazo[2,1-b][1,3]benzothiazol-2-yl]phenyl}urea dihydrochloride (AC220), a uniquely potent, selective, and efficacious FMS-like tyrosine kinase-3 (FLT3) inhibitor; J.Med.Chem. **52** 7808
- 2) Zarrinkar et al. (2009); AC220 is a uniquely potent and selective inhibitor of FLT3 for the treatment of acute myeloid leukemia(AML); Blood **114** 2984
- 3) Fathi and Chen (2017); The role of FLT3 inhibitors in the treatment of FLT3-mutated acute myeloid leukemia; Eur.J.Haematol. **98** 330
- 4) Taylor and Langdon (2017); Hitting the snooze button: Inducing quiescence with the FLT3 inhibitor quizartinib protects hematopoietic progenitors from chemotherapy; Mol.Cell Oncol. **4** e1378156
- Li et al. (2017); Quizartinib (AC220) reverses ABCG2-mediated multidrug resistance: In vitro and In vivo studies; Oncotarget 8 93785
- 6) Bhullar et al. (2013) The FLT3 inhibitor quizartinib inhibits ABCG2 at pharmacologically relevant concentrations, with implications for both chemosensitization and adverse drug reactions; PLoS One **8** e71266

## **PHYSICAL DATA**

Molecular Weight: 560.67

NMR: (Conforms)

Solubility: DMSO (>25 mg/ml)
Physical Description: Off-white solid

Storage and Stability: Store as supplied at -20°C for up to 1 year from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 1 month.

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