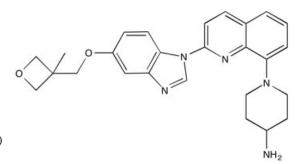
# **Tribioscience**

## Crenolanib, PDGFR and FLT3 inhibitor

Catalog	Unit
TBI4782-5MG	5 mg
TBI4782-25MG	25 mg

## **Product Details**

Formal Name: 1-[2-[5-[(3-Methyloxetan-3-yl)methoxy]benzimidazole-1yl]quinoline-8-yl]piperidin-4-amine Alternate Names: CP-868,596 Molecular Formula:  $C_{26}H_{29}N_5O_2$ Formula Weight: 443.55 CAS Number: 670220-88-9 Purity: >97% Formulation: powder Solubility: Soluble in DMSO (up to 15 mg/ml) or in Ethanol (up to 10 mg/ml) Storage: -20°C Stability:  $\geq$  1 year.



### **Applications**

PDGFR and FLT3 inhibitor

#### **Functions**

Crenolanib is a potent inhibitor of PDGFR (Kd for  $\alpha = 2.1$  nM;  $\beta = 3.2$  nM) and FLT3 (Kd = 0.74 nM). Crenolanib is a type I inhibitor binding only to the active kinase conformation. It showed potent activity against imatinib-resistant PDGFR $\alpha$  mutations D842I, D842V, D842Y, D1842-843M, and deletion 1843 as well as FLT3/ITD and FLT3/D835 mutants. Crenolanib acted synergistically with FLT3-CAR T-cells in a FLT3-ITD+ AML murine xenograft model.

#### **Application Procedures**

First dissolved in DMSO (up to 15 mg/ml) or in Ethanol (up to 10 mg/ml), then diluted to aqueous buffer. Solutions in DMSO or ethanol may be stored at  $-20^{\circ}$  for up to 1 month.

For research use only.