

# GDC-0879

**Catalog No: tcsc0158** 

Available Sizes

Size: 2mg

Size: 5mg

Size: 25mg

Size: 50mg

Size: 50mg

CAS No:

905281-76-7

Formula:

 $C_{19}H_{18}N_4O_2$ 

**Pathway:** MAPK/ERK Pathway

#### Target: Raf

## Purity / Grade:

>98%

#### Solubility:

10 mM in DMSO

#### **Observed Molecular Weight:**

334.37

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### **Product Description**

GDC-0879 is a potent and selective **B-Raf** inhibitor with an **IC<sub>50</sub>** of 0.13 nM.

IC50 & Target: IC50: 0.13 nM (B-Raf)<sup>[1]</sup>

*In Vitro:* GDC-0879 also inhibits pERK with an  $IC_{50}$  of 63 nM<sup>[1]</sup>. GDC-0879 represents a novel potent and selective B-Raf inhibitor that is being evaluated as a potential antitumor agent. GDC-0879 exhibits potent inhibition of Raf/MEK/ERK signaling pathway in V600E B-Raf mutant cell lines with low cellular pMEK1 inhibition  $IC_{50}$  estimates of 59 and 29 nM in A375 melanoma and Colo205 colorectal carcinoma cells, respectively<sup>[2]</sup>.

In Vivo: The pharmacokinetic parameters of GDC-0879 after oral administration of 15, 25, 50, 100, and 200 mg/kg in MCT in mice are estimated as follows:  $k_a = 8.20 h^{-1}$ ,  $k_e = 0.59 h^{-1}$ , and apparent volume of distribution=6.19 L/kg<sup>[2]</sup>.



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