

# **GDC-0834**

**Catalog No: tcsc3123** 

Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

Es

Specifications

#### CAS No:

1133432-49-1

#### Formula:

 $C_{33}H_{36}N_6O_3S$ 

**Pathway:** Protein Tyrosine Kinase/RTK

### **Target:**

Btk

**Purity / Grade:** 

Solubility: DMSO :  $\geq$  32 mg/mL (53.62 mM)

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

596.74

## **Product Description**

GDC-0834 is a potent and selective **BTK** inhibitor. GDC-0834 inhibits BTK with an in vitro **IC**<sub>50</sub> of 5.9 and 6.4 nM in biochemical and cellular assays, respectively, and in vivo **IC**<sub>50</sub> of 1.1 and 5.6  $\mu$ M in mouse and rat, respectively.

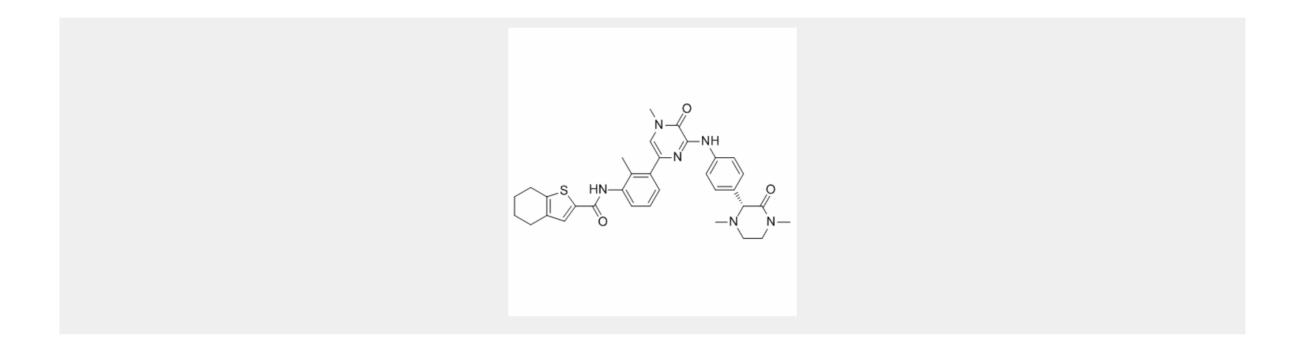
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IC50 & Target: IC50: 5.9 nM (BTK)<sup>[1]</sup>

In Vitro: GDC-0834 suppresses BTK kinase activity with an IC<sub>50</sub> value of  $5.9\pm1.1$  nM with Hill slope value of  $-0.84\pm0.07$  (mean $\pm$ S.E.)<sup>[1]</sup>. GDC-0834 is shown to be a potent reversible inhibitor of six known aldehyde oxidase (AO) substrates with IC<sub>50</sub> values ranging from 0.86 to 1.87  $\mu$ M<sup>[2]</sup>.

In Vivo: The treatment of BALB/c mice with GDC-0834 results in dose-dependent inhibition of pBTK-Tyr223. Animals dosed with 150 or 100 mg/kg GDC-0834 for 2 h show complete inhibition of pBTK-Tyr223 levels in blood, with a mean inhibition of 97 and 96%, respectively. In the rat CIA study, GDC-0834 inhibits pBTK-Tyr223 in rat blood in a dose-dependent manner. The IC<sub>50</sub> estimate of pBTK-Tyr223 inhibition in rats is determined to be  $5.6 \pm 1.6 \mu$ M with m of  $0.51 \pm 0.087$  (mean $\pm$ S.E.)<sup>[1]</sup>.



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