

# BF738735

# Catalog No: tcsc0035356

Available Sizes

Size: 5mg

Size: 10mg

Size: 25mg

Specifications

#### CAS No:

1436383-95-7

#### Formula:

 $\mathrm{C_{21}H_{19}FN_4O_3S}$ 

#### Pathway:

PI3K/Akt/mTOR

#### **Target:**

PI4K

**Purity / Grade:** 

### **Solubility:** 10 mM in DMSO

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

426.46

## **Product Description**

BF738735 is a phosphatidylinositol 4-kinase III beta (**PI4KIII** $\beta$ ) inhibitor with an **IC**<sub>50</sub> of 5.7 nM.

IC50 & Target: IC50: 5.7 nM (PI4KIII $\beta$ ), 1.7  $\mu$ M (PI4KIII $\alpha$ )<sup>[1]</sup>

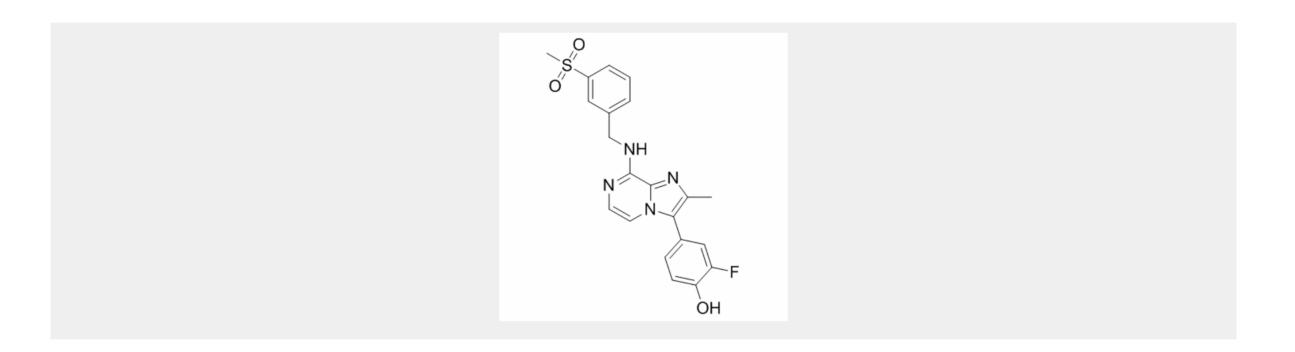
#### In Vitro:

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BF738735 (Compound 1) strongly inhibits PI4KIII $\beta$  activity in vitro, with an IC<sub>50</sub> of 5.7 nM. BF738735 also impairs PI4KIII $\alpha$ , but only at an ~300-fold-higher concentration (IC<sub>50</sub> of 1.7  $\mu$ M). In addition, the activity of BF738735 is analyzed on a set of 150 cellular kinases, including 13 lipid kinases at a concentration of 10  $\mu$ M. For all kinases, the inhibition is less than 10%, indicating that BF738735 specifically inhibits PI4KIII $\beta$  in vitro. BF738735 exhibits a broad spectrum of antiviral activity, as it inhibits all tested species of enteroviruses and rhinoviruses, with 50% effective concentrations ranging between 4 and 71 nM. BF738735 potently inhibits all viruses tested, with EC<sub>50</sub> s ranging between 4 and 71 nM. The cytotoxicity of BF738735, determined in parallel with the EC<sub>50</sub> and using the same culture conditions for 3 to 4 days, is low, with CC<sub>50</sub> values ranging from 11 to 65  $\mu$ M, resulting in high selectivity indices. Low concentrations of BF738735 reduce the amount of luciferase to GuaHCI-treated levels, suggesting that the BF738735 blocks viral RNA replication. The EC<sub>50</sub> of BF738735 in this assay is 77 nM, which is comparable to the inhibition observed in the multicycle assay for coxsackievirus serotype B3 (CVB3)<sup>[1]</sup>.

*In Vivo:* BF738735 is well tolerated by specimens with good plasma levels of the antiviral in circulation and a complete inhibition with 25 mg/kg and some inhibition with 5 mg/kg dose is observed<sup>[2]</sup>.



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