

# IPI-3063

### Catalog No: tcsc0042285

 Available Sizes

 Size: 2mg

 Size: 5mg

 Size: 10mg

 Size: 25mg

  $\boxed{)}$  Specifications

 CAS No: 1425043-73-7

 Formula:  $c_{25}H_{25}N_{7}O_{2}$  

 Pathway: PI3K/Akt/mTOR

Target:

PI3K

Purity / Grade:

>98%

**Solubility:** DMSO : ≥ 83.33 mg/mL (182.94 mM)

## **Observed Molecular Weight:**

455.51

### **Product Description**

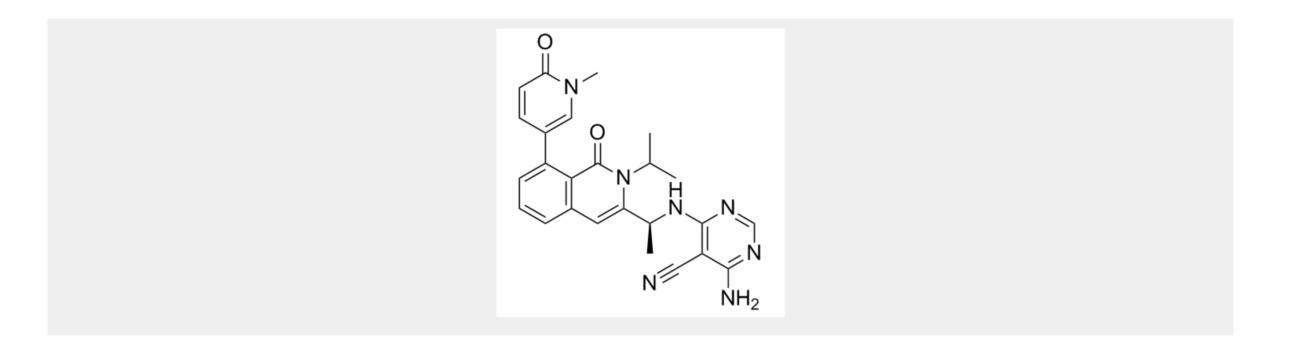
IPI-3063 is a potent and selective **PI3K p1106** inhibitor with an  $IC_{50}$  of 2.5 ± 1.2 nM.

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### IC50 & Target: IC50: 2.5±1.2 nM (p110δ), 1171±533 nM (p110α), 1508±624 nM (p110β), 2187±1529 nM (p110γ)<sup>[1]</sup>

*In Vitro:* IPI-3063 inhibits p110α, p110β, and p110γ with IC<sub>50</sub>s of 1171±533 nM, 1508±624 nM, and 2187±1529 nM, respectively. IPI-3063 potently reduces mouse B cell proliferation, survival, and plasmablast differentiation while increasing antibody class switching to IgG1. IPI-3063 is a p110δ selective compound with an IC<sub>50</sub>=0.1 nM in p110δ-specific cell-based assays and cellular IC<sub>50</sub> values for the other class I PI3K isoforms are at least 1,000-fold higher (IC<sub>50</sub>=1901±1318 nM for p110α, IC<sub>50</sub>=102.8±35.7 nM for p110β, IC<sub>50</sub>=418.8±117.2 nM for p110γ). IPI-3063 is very potent in reducing p-AKT (significant effect at 1 nM). IPI-3063 also reduces p-ERK1/2 with a significant effect at 10 nM. IPI-3063 is very potent, achieving a significant decrease in B cell survival when present at 10 nM<sup>[1]</sup>.



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