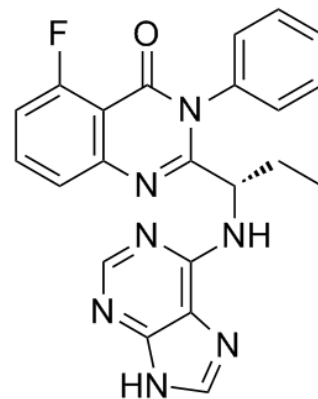


Product Name : Idelalisib
Cat. No. : PC-43336
CAS No. : 870281-82-6
Molecular Formula : C₂₂H₁₈FN₇O
Molecular Weight : 415.423
Target : PI3K
Solubility : DMSO: ≥ 59.7 mg/mL



Biological Activity

Idelalisib (CAL-101, GS-1101) is a potent, selective inhibitor of **PI3K p110 δ** with IC₅₀ of 2.5 nM, 40- to 300-fold selectivity over other class I PI3Ks (p110 α / β γ , IC₅₀=820/565/89nM), and 400- to 4000-fold over C2 β , hVPS34, DNA-PK, and mTOR. Idelalisib (CAL-101, GS-1101) blocks Fc ϵ RI p110 δ -mediated CD63 expression in basophils with an EC₅₀ of 8 nM. Idelalisib (CAL-101, GS-1101) blocks constitutive PI3K signaling, resulting in decreased phosphorylation of Akt and other downstream effectors, an increase in PARP and caspase cleavage and an induction of apoptosis in multiple B-cell malignancies.

References

- Hoellenriegel J, et al. *Blood*. 2011 Sep 29;118(13):3603-12.
Herman SE, et al. *Blood*. 2010 Sep 23;116(12):2078-88.
Lannutti BJ, et al. *Blood*. 2011 Jan 13;117(2):591-4.
Bodo J, et al. *Br J Haematol*. 2013 Oct;163(1):72-80.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

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