

Data Sheet

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 Product Name
 :
 Orlistat

 Cat. No.
 :
 PC-20125

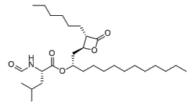
 CAS No.
 :
 96829-58-2

 Molecular Formula
 :
 C₂₉H₅₃NO₅

 Molecular Weight
 :
 495.75

Target : Diacylglycerol Lipase (DAGL)

Solubility : 10 mM in DMSO



Biological Activity

Orlistat ((–)-Tetrahydrolipstatin, Ro-18-0647) is a small molecule inhibitor of pancreatic and gastric **lipase**, inhibits **DAGL** α , DAGL β , ABHD12, ABHD16A, and platelet-activating factor acetylhydrolase (PAF-AH) with IC50 of 0.06, 0.1, 0.08, 0.03, and 0.05 μ M, respectively.

Orlistat does not inhibit fatty acid amide hydrolase (FAAH) with IC50 of >100 uM.

Orlistat (1 uM) decreases ionomycin-induced production of the endocannabinoid 2-arachidonoyl glycerol (2-AG) in N18TG2 murine neuroblastoma cells.

Orlistat also inhibits fatty acid synthase (**FASN**; apparent Ki =0.1 μ M)) and inhibits the proliferation of PC3 prostate cancer cells in a concentration-dependent manner.

Orlistat (10 mg/kg) decreases serum cholesterol levels and total body weight in a mouse model of obesity induced by a high-fat diet.

References

Bisogno, T.et al. *J. Cell Biol.* 163(3), 463-468 (2003).

Hoover, H.S.et al. *Bioorg. Med. Chem. Lett.* 18(22), 5838-5841 (2008).

Bisogno, T., et al. Biochim. Biophys. Acta 1761(2), 205-212 (2006).

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

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