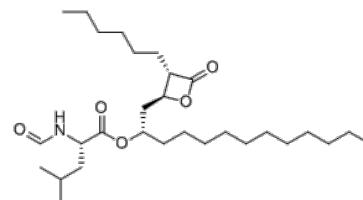


Product Name : Orlistat
Cat. No. : PC-20125
CAS No. : 96829-58-2
Molecular Formula : C₂₉H₅₃NO₅
Molecular Weight : 495.75
Target : Phospholipase
Solubility : 10 mM in DMSO



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 μ M.

Orlistat (1 μ M) 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 K_i = 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

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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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