

Catalog Number: CM04959

## 产品信息

**Catalog Number:**  
CM04959

**CAS号:**  
1446817-84-0

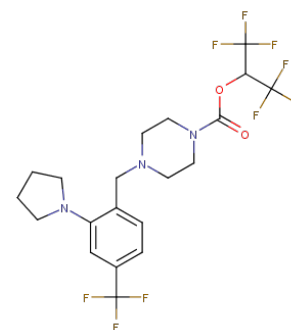
**分子式:**  
 $C_{20}H_{22}F_9N_3O_2$

**主要靶点:**  
Lipase

**主要通路:**  
代谢

**分子量:**  
507.39

**溶解度:**  
Ethanol:10 mg/mL (19.71 mM); H<sub>2</sub>O:Insoluble; DMSO:55 mg/mL (108.4 mM)



## 靶点活性

MGLL (mouse):27 nM|MGLL (human):14 nM

## 体外活性

ABX-1431是一种强效的人类MGLL抑制剂(平均IC<sub>50</sub>值为0.014 μM),对ABHD6具有超过100倍的选择性,对PLA2G7具有超过200倍的选择性。在对完整的人类PC3细胞进行处理时,经过30分钟的抑制剂孵育之后,ABX-1431对MGLL活性产生了浓度依赖性的抑制,其IC<sub>50</sub>值为0.0022 μM,约为体外观察到的效力的6倍。在细胞基础实验中,维持了对MGLL超过100倍的选择性,相比之下,ABHD6的IC<sub>50</sub>值为0.253 μM,PLA2G7的IC<sub>50</sub>值为494 μM。

## 体内活性

在体内,ABX-1431抑制啮齿动物脑内的MGLL活性(ED<sub>50</sub>=0.5-1.4 mg/kg),提高脑内2-AG浓度,并在大鼠甲醛疼痛模型中抑制疼痛行为。

## 动物实验

All animals were 6-8 weeks old at the time of study and weighed between 197-217 g. Animals (n = 3 per dosing route) were dosed with ABX-1431 at 1 mg/kg iv (in 70% polyethylene glycol (PEG) 400 in hydroxypropyl-β-cyclodextran (HPBCD) in saline) or 5 mg/kg po (in 70% PEG 400 in 0.5% carboxymethylcellulose (CMC, w/v) in saline). Animals were fasted overnight, and food withheld until 4 h post dose. Approximately 100 μL of blood were collected via a jugular vein catheter at 0.033, 0.083, 0.25, 0.5, 1, 2, 4, 6, 8, and 24 h after intravenous and 0.25, 0.5, 1, 2, 4, 6, 8, and 24 h after oral administration. All blood samples were collected into tubes containing 400 μL acetonitrile to immediately inactivate blood esterase activity and frozen at -80 °C. Samples were thawed and centrifuged (14,000 rpm for 5 min at 4 °C) and the supernatant transferred for LCMS analysis.

## 细胞实验

Human prostate cancer PC3 cells were grown in F-12K medium supplemented with 10% fetal bovine serum at 37 °C with 5% CO<sub>2</sub> to ~80% confluency in 100 mm dishes. ABX-1431 was added to cells to give final concentration of 0.1-10 μM compound in serum free media. Cells were incubated with compound for 30 min at 37 °C with 5% CO<sub>2</sub>. Cells were washed, harvested, and lysed by probe sonication. Cell lysates (2mg/mL) were treated with JW912 (1 μM) and analyzed by SDS-PAGE and in-gel fluorescence scanning.

## 储存

Powder: -20 °C for 3 years | In solvent: -80 °C for 1 year | Shipping with blue ice.