For Research Use Only Esonarimod



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Catalog Number: CM04738

产品信息

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CAS号: 101973-77-7

分子式: C₁₄H₁₆O₄S 主要靶点: IL Receptor

主要通路: 免疫与炎症 DMS0:60 mg/mL (214.03 mM)

280.34

溶解度:

H₂C CH

体外活性

The IC50 of Esonarimod is 117.5 $\,\mu$ g/mL In RAW264.7 cells, Esonarimod (KE-298) (10 to 300 $\,\mu$ g/mL) suppresses the production of NO in a dose-dependent manner.

体内活性

After repeated oral administration of Esonarimod (14C-KE-298), the radioactivity decreases rapidly and no tendency towards accumulation is found.

动物实验

Seven-week-old male Wistar rats is administered Esonarimod (5 mg/kg once daily) orally by gastric intubation.

细胞实验

For in vitro experiment, Esonarimod (KE-298) is dissolved in ethanol and diluted with culture medium or distilled water. RAW264.7 cells are used in this study. For NO production, RAW264.7 cells [2×105/0.2 mL of RPMI-1640 supplemented by 10% heat inactivated fetal bovine serum (FBS), penicillin G (100 U/mL), and streptomycin (100 μ g/mL)] are stimulated with 100 ng/mL of Escherichia coli 026:B6 lipopolysaccharide in the presence of Esonarimod (KE-298) (0, 10, 30, 100, 200, 300 μ g/mL) in 96 well plates and incubated 24 h at 37°C in an atmosphere of 5% CO2 in air. After incubation, the supernatants are collected and assayed for nitrite (NO2-) instead of NO[1].

描述

Esonarimod is an antirheumatic drug.

储存

Powder: -20°C for 3 years | In solvent: -80°C for 2 years