For Research Use Only

GP130 receptor agonist-1



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Catalog Number: CM05850

产品信息

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CAS号: 339303-87-6

分子式: C₁₅H₁₁FN₂S

主要靶点:

Interleukin **主要通路:** 免疫与炎症

分子量: 270.32 溶解度:

DMSO:55 mg/mL (203.46 mM)

Compound 2 (N-(4-Fluorophenyl)-4-phenyl-2-thiazolami) treatment showed a 2-fold increase in phosphorylation of STAT3 within 10 min at its regulatory Tyr705 site in SH-SY5Y cells.. Compound 2 treatment increases phosphorylation of AKT at its regulatory Thr308 site and phosphorylation of ERK1/2 at its regulatory Thr202/Tyr204 site in the serum free media condition in SH-SY5Y cells, and in primary cortical neurons.

For Compound 2 (N-(4-Fluorophenyl)-4-phenyl-2-thiazolami), mice are dosed orally at 10 or 30 mg/kg, or injected subcutaneously (SQ) at 10 mg/kg, and euthanized after 1, 2, 4, 6, and 8 h post dose. At 2 h after SQ delivery at 10 mg/kg the brain Cmax is 161 ng/g while dosing at 30 mg/kg orally, results in the brain Cmax of 156 ng/g (0.57 $\,\mu$ M). The brain to plasma ratio for 2 is $\sim\!4$:1 for oral 30 mg/kg and $\sim\!7.5$:1 for 10 mg/kg SQ injection.

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