## For Research Use Only 2,5-dimethyl Celecoxib



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Catalog Number: CM13665

产品信息

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CAS号:

457639-26-8

分子式: C<sub>18</sub>H<sub>16</sub>F<sub>3</sub>N<sub>3</sub>O<sub>2</sub>S

主要靶点:

Prostaglandin Receptor|Wnt/beta-catenin|Apoptosis

G蛋白偶联受体|干细胞|凋亡|免疫 与炎症|细胞骨架

分子量: 395.4 溶解度:

Ethanol:3 mg/mL,DMSO:5 mg/mL,DMF:5 mg/mL,DMSO:PBS (pH 7.2) (1:3):0.25 mg/mL

体外活性

2,5-dimethyl Celecoxib(1–100?  $\mu$  M) decreased the viability of GBM cell lines in a dose-dependent manner. 2,5-dimethyl Celecoxib downregulated  $\beta$ -catenin target genes expression in A-172, T98G and U-138 MG cell lines. Apoptosis was induced, and cell cycle distribution was altered after the treatment with 2,5-dimethyl Celecoxib in T98G cell line.2,5-dimethyl Celecoxib downregulated β-catenin target genes expression in patient-derived primary GBM cell lines P1 and P6[1]

体内活性

2,5-dimethyl Celecoxib prevented cardiac remodeling and markedly reduced urinary albumin excretion without altering blood pressure in mice. 2,5-dimethyl Celecoxib prevented podocyte injury, glomerulosclerosis, and interstitial fibrosis in the kidney of mice loaded with angiotensin II and high-salt load. 2,5-dimethyl Celecoxib reduced the phosphorylation level of Akt and activated glycogen synthase kinase-3, which led to the suppression of the Wnt/  $\beta$ -catenin signal in the heart and kidney. 2,5-dimethyl Celecoxib also reduced the expression level of snail, a key transcription factor for the epithelial–mesenchymal transition and of which gene is a target of the Wnt/  $\beta$ -catenin signal[2].

描述

2,5-dimethyl Celecoxib is a celecoxib derivative and a targeted inhibitor of microsomal prostaglandin E synthase-1 (mPGES-1), a key enzyme in the PGE2 synthesis pathway of inflammatory mediators.

储存

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