EP2 receptor antagonist-2

| Cat. No.: | HY-153129 |
|--------------------|---|
| CAS No.: | 615273-95-5 |
| Molecular Formula: | C ₁₅ H ₁₄ F ₃ N ₃ O |
| Molecular Weight: | 309.29 |
| Target: | Prostaglandin Receptor |
| Pathway: | GPCR/G Protein |
| Storage: | 4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light) |

SOLVENT & SOLUBILITY

| | | | Solvent Concentration | Preparing Stock Solutions |
|---|------------|---------------------|-------------------------------------|------------------------------|
| 32.3321 mL | 16.1661 mL | 3.2332 mL | 1 mM | |
| 6.4664 mL | 3.2332 mL | 0.6466 mL | 5 mM | |
| 3.2332 mL | 1.6166 mL | 0.3233 mL | 10 mM | |
| | i i | ppropriate solvent. | ubility information to select the a | Please refer to the solu |
| 10 mM0.3233 mL1.6166 mL3.2332 mLPlease refer to the solubility information to select the appropriate solvent.1. Add each solvent one by one: 10% DMSO >> 90% corn oil | | | | |

| BIOLOGICAL ACTIV | |
|---------------------------|--|
| Description | EP2 receptor antagonist-2 (CID891729) is an antagonist of EP2 receptor. EP2 receptor antagonist-2 inhibits the EP2 receptor activation induced by PGE2. EP2 receptor antagonist-2 also suppresses lactate dehydrogenase (LDH) release induced by N-methyl-D-aspartate (NMDA) ^[1] . |
| IC ₅₀ & Target | EP |
| In Vitro | EP2 receptor antagonist-2 shows cytotoxicity against C6G cells with an CC ₅₀ value of 575 μM ^[1] . EP2 receptor antagonist-2 (20 μM; 30 min) decreases NMDA (30 μM)-induced LDH release in rat primary hippocampal neurons (DIV14) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

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[1]. Jiang J, et al. Discovery of 2-Piperidinyl Phenyl Benzamides and Trisubstituted Pyrimidines as Positive Allosteric Modulators of the Prostaglandin Receptor EP2. ACS Chem Neurosci. 2018 Apr 18;9(4):699-707.

Caution: Product has not been fully validated for medical applications. For research use only.

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