



Data Sheet

 Product Name:
 HJC0350

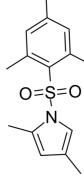
 Cat. No.:
 CS-6510

 CAS No.:
 885434-70-8

 Molecular Formula:
 C15H19NO2S

Molecular Weight: 277.38
Target: Others
Pathway: Others

Solubility: DMSO: 33.33 mg/mL (120.16 mM; Need ultrasonic)



BIOLOGICAL ACTIVITY:

HJC0350 is a potent and specific **EPAC2** antagonist with an **IC**₅₀ of 0.3 μ M. IC50 & Target: IC50: 0.3 μ M (EPAC2)^[1] **In Vitro**: HJC0350 has an apparent IC₅₀ value of 0.3 μ M for competing with 8-NBD-cAMP binding of EPAC2, and is about 133-fold more potent than cAMP. HJC0350 is found not to inhibit EPAC1-mediated Rap1-GDP exchange activity at 25 μ M in the presence of equal concentration of cAMP, indicating that it is EPAC2-specific antagonists. Pretreatment of HEK293/EPAC2-FL cells with 10 μ M HJC0350 fully blocks the 007-AM induced decrease of FRET^[1].

References:

[1]. Chen H, et al. Identification and characterization of small molecules as potent and specific EPAC2 antagonists. J Med Chem. 2013 Feb 14;56(3):952-62.

CAIndexNames:

1H-Pyrrole, 2,4-dimethyl-1-[(2,4,6-trimethylphenyl)sulfonyl]-

SMILES:

O=S(N1C(C)=CC(C)=C1)(C2=C(C)C=C(C)C=C2C)=O

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

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