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Product Data Sheet

Rodatristat

Cat. No.: HY-120083 CAS No.: 1673568-73-4 Molecular Formula: $C_{27}H_{27}ClF_{3}N_{5}O_{3}$

Molecular Weight: 561.98

Target: 5-HT Receptor; Tryptophan Hydroxylase

Pathway: GPCR/G Protein; Neuronal Signaling; Metabolic Enzyme/Protease

Storage: Powder -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

BIOLOGICAL ACTIVITY

Rodatristat (KAR5417) is a potent tryptophan hydroxylase 1 (TPH1) and TPH2 inhibitor with IC $_{50}$ s value of 33 nM and 7 nM, Description respectively, and shows robust reduction of intestinal serotonin (5-HT) levels in $mice^{[1]}$.

IC₅₀ & Target TPH1 TPH2 serotonin 33 nM (IC₅₀) 7 nM (IC₅₀)

In Vivo Rodatristat (10-50 mg/kg; oral administration; mice) treatment decreases intestinal 5-HT concentrations at 50 mg/kg, their efficacy drop off significantly at the lower 10 mg/kg dose^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

| Animal Model: | $Mice^{[1]}$ |
|-----------------|--|
| Dosage: | 10 mg/kg and 50 mg/kg |
| Administration: | Oral administration |
| Result: | Decreased intestinal 5-HT concentrations at 50 mg/kg, their efficacy dropped off significantly at the lower 10 mg/kg dose. |

REFERENCES

[1]. Goldberg DR, et al. Optimization of spirocyclic proline tryptophan hydroxylase-1 inhibitors. Bioorg Med Chem Lett. 2017 Feb 1;27(3):413-419.

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

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