Proteins

Imaradenant

Cat. No.: HY-101980 CAS No.: 1321514-06-0 Molecular Formula: C₁₅H₁₁ClFN₅ Molecular Weight: 315.73

Target: Adenosine Receptor Pathway: GPCR/G Protein

Storage: Powder -20°C

3 years 2 years

In solvent -80°C 2 years

> -20°C 1 year

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: ≥83.3 mg/mL (263.83 mM)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.1673 mL	15.8363 mL	31.6726 mL
	5 mM	0.6335 mL	3.1673 mL	6.3345 mL
	10 mM	0.3167 mL	1.5836 mL	3.1673 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (6.59 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (6.59 mM); Clear solution

BIOLOGICAL ACTIVITY

AZD4635 (HTL1071) is a potent, selective and orally active adenosine A2A receptor (A2AR) antagonist. AZD4635 binds to Description human A2AR with a K_i of 1.7 nM and shows >30-fold selectivity over other adenosine receptors^[1]. IC₅₀ & Target Ki: 1.7 nM (A2AR)[1]

In the presence of 0.1, 1 and 10 μ M adenosine, the IC₅₀s of AZD4635 for inhibition of cAMP production are 0.79, 10.0 and 142.9 nM, respectively^[1].

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

In Vitro

CUSTOMER VALIDATION

- Adv Sci (Weinh). 2022 Jan 22;e2104793.
- J Exp Clin Cancer Res. 2022 Oct 14;41(1):302.

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REFERENCES

[1]. Alexandra Borodovsky, et al. Abstract 5580: Preclinical pharmacodynamics and antitumor activity of AZD4635, a novel adenosine 2A receptor inhibitor that reverses adenosine mediated T cell suppression. AACR; Cancer Res 2017;77(13 Suppl):Abstract nr 5580.

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

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