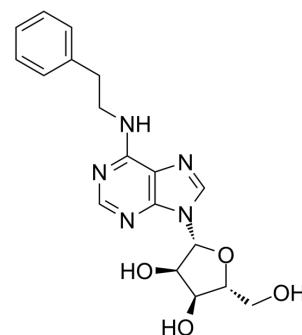


N6-(2-Phenylethyl)adenosine

| | | | |
|---------------------------|---|-------|---------|
| Cat. No.: | HY-101854 | | |
| CAS No.: | 20125-39-7 | | |
| Molecular Formula: | C ₁₈ H ₂₁ N ₅ O ₄ | | |
| Molecular Weight: | 371.39 | | |
| Target: | Adenosine Receptor | | |
| Pathway: | GPCR/G Protein | | |
| Storage: | Powder | -20°C | 3 years |
| | | 4°C | 2 years |
| | In solvent | -80°C | 2 years |
| | | -20°C | 1 year |



SOLVENT & SOLUBILITY

In Vitro

DMSO : 100 mg/mL (269.26 mM; Need ultrasonic)
 H₂O : < 0.1 mg/mL (insoluble)

| Preparing Stock Solutions | Solvent | | Mass | | |
|---------------------------|---------------|--|-----------|------------|------------|
| | Concentration | | 1 mg | 5 mg | 10 mg |
| | 1 mM | | 2.6926 mL | 13.4629 mL | 26.9259 mL |
| | 5 mM | | 0.5385 mL | 2.6926 mL | 5.3852 mL |
| | 10 mM | | 0.2693 mL | 1.3463 mL | 2.6926 mL |

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.5 mg/mL (6.73 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.5 mg/mL (6.73 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.5 mg/mL (6.73 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

N6-(2-Phenylethyl)adenosine (N6-Phenethyladenosine), an adenosine derivative, is a potent adenosine receptors (AR) agonist with K_i values of 11.8 nM, 30.1 nM, 0.63 nM for rat A₁AR, human A₁AR and hA₃AR, respectively^[1].

IC₅₀ & Target

Ki: 11.8 nM (rA₁AR), 30.1 nM (hA₁AR) and 0.63 nM (hA₃AR)^[1]

In Vitro

N6-(2-Phenylethyl)adenosine (N6-Phenethyladenosine) inhibits rA₂AR (IC₅₀=560 nM), hA₂AR (IC₅₀=2250 nM) in CHO cells^[1].

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

REFERENCES

[1]. Tchilibon S, et al. Exploring distal regions of the A3 adenosine receptor binding site: sterically constrained N6-(2-phenylethyl)adenosine derivatives as potent ligands. Bioorg Med Chem. 2004 May 1;12(9):2021-34.

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

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