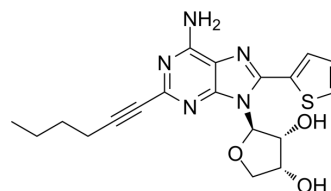


## LJ-4517

Cat. No.:	HY-151139
CAS No.:	2988849-20-1
Molecular Formula:	C <sub>19</sub> H <sub>21</sub> N <sub>5</sub> O <sub>3</sub> S
Molecular Weight:	399.47
Target:	Adenosine Receptor
Pathway:	GPCR/G Protein
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

Description	LJ-4517 is a potent A <sub>2A</sub> AR antagonist, with a K <sub>i</sub> of 18.3 nM. LJ-4517 is potent in displacing the binding of [ <sup>3</sup> H]ZM241385 (HY-19532) at WT A <sub>2A</sub> AR <sup>[1]</sup> . LJ-4517 is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAC) with molecules containing Azide groups.
IC <sub>50</sub> & Target	A2AR 18.3 nM (K <sub>i</sub> )
In Vitro	LJ-4517 (10 μM) induces cAMP accumulation <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Shiriaeva A, et al. GPCR Agonist-to-Antagonist Conversion: Enabling the Design of Nucleoside Functional Switches for the A<sub>2A</sub> Adenosine Receptor. J Med Chem. 2022 Aug 17.

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

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