

Product Data Sheet

A3AR modulator 1

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-151899} \\ \textbf{CAS No.:} & 2991693-62-8 \\ \textbf{Molecular Formula:} & \textbf{C}_{23}\textbf{H}_{25}\textbf{IN}_{4} \\ \textbf{Molecular Weight:} & 484.38 \\ \end{array}$

Target: Adenosine Receptor
Pathway: GPCR/G Protein

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description	A3AR modulator 1 (MRS8054) is an orally active A3 adenosine receptor (A3AR) (Adenosine Receptor) positive allosteric modulator (PAM). A3AR modulator 1 greatly enhances Cl-IB-MECA-stimulated [35 S]GTP $_{\gamma}$ S binding E $_{max}$ [1].
IC ₅₀ & Target	Adenosine A ₃ receptor
In Vitro	A3AR modulator 1 (compound 39) shows a few weak off-target interactions, with K_i values of 0.123 μ M, 0.891 μ M, and 2.6 μ M for translocator protein (TSPO), opioid receptor σ 2, and 5HT2B receptor, respectively ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	In fasted Wistar rats, A3AR modulator 1 (compound 39) has considerably longer in vivo half-life and improved oral bioavailability (3.44 h, 64.0%F at 3 mg/kg; 3.84 h, 61.5%F at 10 mg/kg), indicating substantial oral bioavailability ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Lucas B Fallot, e al. Structure-Activity Studies of 1 H-Imidazo[4,5- c]quinolin-4-amine Derivatives as A3 Adenosine Receptor Positive Allosteric Modulators. J Med Chem. 2022 Nov 24;65(22):15238-15262.

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

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