roduct Da

A-841720

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-103550} \\ \textbf{CAS No.:} & 869802\text{-}58\text{-}4 \\ \textbf{Molecular Formula:} & \textbf{C}_{17}\textbf{H}_{21}\textbf{N}_{5}\textbf{OS} \\ \textbf{Molecular Weight:} & 343.45 \\ \end{array}$

Target: mGluR

Pathway: GPCR/G Protein; Neuronal Signaling

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

Analysis.

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BIOLOGICAL ACTIVITY

Description	A-841720 is a potent, non-competitive and selective mGlu1 receptor antagonist with an IC $_{50}$ of 10 nM for human mGlu1 receptor. A-841720 displays 34-fold selectivity over mGlu5 (IC $_{50}$ of 342 nM), and no significant activity at a range of other neurotransmitter receptors, ion channels, and transporters. A-841720 has the potential for chronic pain research ^{[1][2]} .	
IC ₅₀ & Target	Human mGluR1 10 nM (IC ₅₀)	human mGluR5 342 nM (IC ₅₀)

REFERENCES

[1]. O El-Kouhen, et al. Blockade of mGluR1 receptor results in analgesia and disruption of motor and cognitive performances: effects of A-841720, a novel non-competitive mGluR1 receptor antagonist. Br J Pharmacol. 2006 Nov;149(6):761-74.

[2]. Lorenzo Morè, et al. Comparison of the mGluR1 antagonist A-841720 in rat models of pain and cognition. Behav Pharmacol. 2007 Jul;18(4):273-81.

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

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