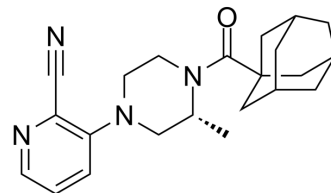


## VU0469650

<b>Cat. No.:</b>	HY-110191
<b>CAS No.:</b>	1443748-47-7
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>28</sub> N <sub>4</sub> O
<b>Molecular Weight:</b>	364.48
<b>Target:</b>	mGluR
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	VU0469650 is a potent, selective and CNS-penetrated negative allosteric modulator of mGlu <sub>1</sub> receptor, with an IC <sub>50</sub> of 99 nM [1].
<b>IC<sub>50</sub> &amp; Target</b>	mGluR 1 99 nM (IC <sub>50</sub> )
<b>In Vivo</b>	VU0469650 (male Sprague-Dawley rats, 0.2 mg/kg, IV, once) exhibits moderate to high clearance, a moderate volume of distribution, and a half-life of approximately 1 hour <sup>[1]</sup> . VU0469650 (male Sprague-Dawley rats, 10 mg/kg, IP, once) shows an excellent selectivity profile and good exposure in both plasma and brain samples <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Lovell KM, Felts AS, Rodriguez AL, et al. N-Acyl-N'-arylpiperazines as negative allosteric modulators of mGlu1: identification of VU0469650, a potent and selective tool compound with CNS exposure in rats. *Bioorg Med Chem Lett.* 2013;23(13):3713-3718.

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

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