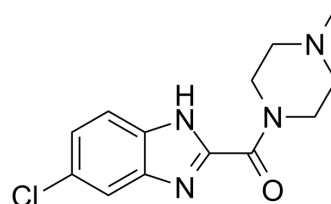


## JNJ10191584

<b>Cat. No.:</b>	HY-123532
<b>CAS No.:</b>	73903-17-0
<b>Molecular Formula:</b>	C <sub>13</sub> H <sub>15</sub> ClN <sub>4</sub> O
<b>Molecular Weight:</b>	278.74
<b>Target:</b>	Histamine Receptor
<b>Pathway:</b>	GPCR/G Protein; Immunology/Inflammation; Neuronal Signaling
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	<p>JNJ10191584 (VUF6002) is an orally active and selective histamine H<sub>4</sub> receptor antagonist with a K<sub>i</sub> value of 26 nM. JNJ10191584 shows 540-fold selectivity to H<sub>4</sub> receptor over H<sub>3</sub> receptor with a K<sub>i</sub> value of 14.1 μM. JNJ10191584 inhibits chemotaxis of eosinophils and mast cells with IC<sub>50</sub> values of 530 nM and 138 nM, respectively<sup>[1][2]</sup>.</p>	
<b>IC<sub>50</sub> &amp; Target</b>	Human H <sub>4</sub> Receptor 26 nM (K <sub>i</sub> )	human H <sub>3</sub> receptor 14.1 μM (K <sub>i</sub> )
<b>In Vitro</b>	<p>JNJ10191584 shows binding affinity of 26 nM and 14.1 μM to H<sub>4</sub> and H<sub>3</sub> receptor, respectively<sup>[1]</sup>. JNJ10191584 (3 h) shows inhibitory effects to chemotaxis of eosinophils and mast cells with IC<sub>50</sub> values of 530 nM and 138 nM, respectively<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>	
<b>In Vivo</b>	<p>JNJ10191584 (10 μg/μL; intra locus coeruleus (LC) administration; once) abolishes VUF-induced anti-allodynic effect in spared nerve injury (SNI) mice<sup>[1]</sup>. JNJ10191584 (10 μg/μL; intra LC administration; once) prevents the anti-allodynic effect of VUF 8430 in SNI mice<sup>[1]</sup>. JNJ10191584 (6 μg/mouse; intrathecal administration; pretreat once) prevents VUF 8430-induced anti-allodynic effect in SNI mice<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>	

### REFERENCES

[1]. Venable JD, et al. Preparation and biological evaluation of indole, benzimidazole, and thienopyrrole piperazine carboxamides: potent human histamine h(4) antagonists. *J Med Chem.* 2005 Dec 29;48(26):8289-98.

[2]. Sanna MD, et al. Histamine H<sub>4</sub> receptor stimulation in the locus coeruleus attenuates neuropathic pain by promoting the coeruleospinal noradrenergic inhibitory pathway. *Eur J Pharmacol.* 2020 Feb 5;868:172859.

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

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