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## DATASHEET

JNJ 10191584 maleate

### Product overview

|                   |   |
|-------------------|---|
| Name              | JNJ 10191584 maleate  |
| Cat No            | HB1596  |
| Alternative names | VUF 6002  |
| Biological action | Antagonist  |
| Purity            | >99%  |
| Description       | Potent, selective H <sub>4</sub> receptor silent antagonist |

### Biological Data

|                        |  |
|------------------------|--|
| Biological description | Potent and selective H <sub>4</sub> histamine receptor silent antagonist. Selective for human H <sub>4</sub> over H <sub>3</sub> by >540-fold (K <sub>i</sub> values are 26 nM and 14.1 μM respectively). Inhibits chemotaxis of mast cells and eosinophils in vitro. Shows anti-inflammatory and antinociceptive actions. |
|------------------------|--|

### Solubility & Handling

|                      |   |
|----------------------|---|
| Storage instructions | room temperature (desiccate)  |
| Solubility overview  | Soluble in DMSO (50mM)  |
| Important            | This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use. |

### Chemical Data

|                    |   |
|--------------------|---|
| Chemical name      | 1-[(5-Chloro-1 <i>H</i> -benzimidazol-2-yl)carbonyl]-4-methylpiperazine maleate   |
| Molecular Weight   | 394.81  |
| Chemical structure | The chemical structure shows a 5-chlorobenzimidazole ring system linked via its carbonyl group to the nitrogen of a piperazine ring. The piperazine ring is substituted with a methyl group and a maleate ester side chain. The maleate ester group consists of a four-carbon chain ending in a carboxylic acid group, which is esterified to a maleic anhydride residue. |
| Molecular Formula  | C <sub>13</sub> H <sub>15</sub> ClN <sub>4</sub> O <sub>4</sub>   |
| CAS Number         | 869497-77-6   |
| PubChem identifier | 11718163  |
| SMILES             | O=C(O)/C=C\ C(O)=O.C1C(C=C3)=CC2=C3N=C(N2)C(N1CCN(C)CC1)=O  |
| InChiKey           | KOTJFAYEELTYCZ-BTJKTKUSA-N  |

### References

**Synthesis and structure-activity relationships of indole and benzimidazole piperazines as histamine H(4) receptor antagonists.**

Terzioglu N et al (2004) Bioorg Med Chem Lett 14(21)

PubMedID [15454206](#)

**Inhibitory effects of histamine H4 receptor antagonists on experimental colitis in the rat.**

Varga C *et al* (2005) Eur J Pharmacol 522(1-3)

**PubMedID**

16213481

**Preparation and biological evaluation of indole, benzimidazole, and thienopyrrole piperazine carboxamides: potent human histamine h(4) antagonists.**

Venable JD *et al* (2005) J Med Chem 48(26)

**PubMedID**

16366610

**Antiinflammatory and antinociceptive effects of the selective histamine H4-receptor antagonists JNJ7777120 and VUF6002 in a rat model of carrageenan-induced acute inflammation.**

Coruzzi G *et al* (2007) Eur J Pharmacol 563(1-3)

**PubMedID**

17382315

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