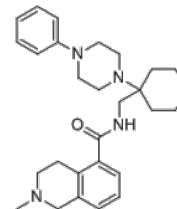


**Product Name** : JNJ-42253432  
**Cat. No.** : PC-60767  
**CAS No.** : 1428327-35-8  
**Molecular Formula** : C<sub>28</sub>H<sub>38</sub>N<sub>4</sub>O  
**Molecular Weight** : 446.639  
**Target** : P2X Receptor  
**Solubility** : 10 mM in DMSO



## Biological Activity

JNJ-42253432 is a potent, selective, CNS-penetrant **P2X7** antagonist with pK<sub>i</sub> of 9.1 and 7.9 for rat and human P2X7 channel, respectively.

JNJ-42253432 does not block calcium flux via human P2X1, P2X2, P2X3, P2X2/P2X3, and P2X4 at 10 μM.

JNJ-42253432 attenuates both ATP- and Bz-ATP-induced currents from hP2X7-1321N1 cells with similar potencies (pEC<sub>50</sub>=7.0).

JNJ-42253432 blocks the release of IL-1β induced by Bz-ATP in freely moving rat brain, also increases serotonin levels.

JNJ-42253432 attenuates amphetamine-induced hyperactivity in vivo.

## References

Lord B, et al. *J Pharmacol Exp Ther*. 2014 Dec;351(3):628-41.

Amhaoul H, et al. *Neuropharmacology*. 2016 Jun;105:175-185.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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