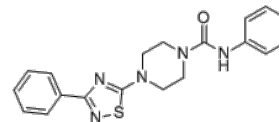


Product Name : JNJ-1661010
Cat. No. : PC-21982
CAS No. : 681136-29-8
Molecular Formula : C₁₉H₁₉N₅OS
Molecular Weight : 365.46
Target : FAAH
Solubility : 10 mM in DMSO



Biological Activity

JNJ-1661010 (Takeda-25) is a potent and selective fatty acid amide hydrolase (FAAH) inhibitor with IC₅₀ of 34 and 33 nM for rat FAAH and human FAAH, respectively.

JNJ-1661010 is >100-fold preferentially selective for FAAH-1 when compared to FAAH-2.

JNJ-1661010 dose-dependently increases arachidonoyl ethanolamide, oleoyl ethanolamide, and palmitoyl ethanolamide in the rat brain.

JNJ-1661010 attenuates tactile allodynia in the rat mild thermal injury model of acute tissue damage and in the rat spinal nerve ligation (Chung) model of neuropathic pain.

JNJ-1661010 also diminishes thermal hyperalgesia in the inflammatory rat carrageenan paw model.

References

Keith JM, et al. Bioorg Med Chem Lett. 2008 Sep 1;18(17):4838-43.

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

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