

Data Sheet

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 Product Name
 : PF-04457845

 Cat. No.
 : PC-20049

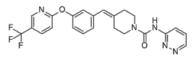
 CAS No.
 : 1020315-31-4

 Molecular Formula
 : C23H20F3N5O2

Molecular Weight : 455.44

Target : FAAH

Solubility : 10 mM in DMSO



Biological Activity

PF-04457845 (Redafamdastat) is a highly potent, selective and covalent fatty acid amide hydrolase (**FAAH**) inhibitor with IC50 of 7.2 nM (human FAAH).

PF-04457845 displays no activity against other FP-reactive serine hydrolases at 100 uM, as well as a broad panel of 68 targets including receptors, enzymes, ion channels, and transporters.

PF-04457845 is orally bioavailable, and PF-04457845 (10 mg/kg, p.o.) displays antihyperalgesic activity with long duration of action in the CFA model of inflammatory pain in rats.

PF-04457845 does not elicit effect in motility, catalepsy, and body temperature.

References

Sébastien Dilly, et al. Antioxidants (Basel). 2023 Feb 10;12(2):440.

Li Yet al. *Proc. Natl. Acad. Sci. USA*. 2012;109:12526–12531.

Ahn K, et al. *J Pharmacol Exp Ther.* 2011 Jul;338(1):114-24.

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

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