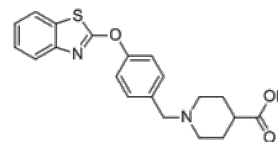


**Product Name** : JNJ-26993135  
**Cat. No.** : PC-63409  
**CAS No.** : 841202-16-2  
**Molecular Formula** : C<sub>20</sub>H<sub>20</sub>N<sub>2</sub>O<sub>3</sub>S  
**Molecular Weight** : 368.451  
**Target** : Aminopeptidase  
**Solubility** : 10 mM in DMSO



### Biological Activity

JNJ-26993135 is a potent, selective leukotriene A4 hydrolase (**LTA4H**) inhibitor, inhibits both the epoxide hydrolase and aminopeptidase activities of recombinant human LTA4H with IC<sub>50</sub> of 10 nM.

JNJ-26993135 has no significant effects on LTC<sub>4</sub>, lipoxin A<sub>4</sub>, or PGE<sub>2</sub> production.

JNJ-26993135 dose-dependently inhibits ex vivo LTB<sub>4</sub> production in blood (IC<sub>50</sub>=339 nM), in parallel with dose-dependent inhibition of neutrophil influx (ED<sub>50</sub>, 1-3 mg/kg) and ear edema in murine model of arachidonic acid-induced ear inflammation.

JNJ-26993135 selectively inhibited LTB<sub>4</sub> production, without affecting cysteinyl leukotriene production.

### References

Rao NL, et al. *J Pharmacol Exp Ther*. 2007 Jun;321(3):1154-60.

Whittle BJ, et al. *Br J Pharmacol*. 2008 Mar;153(5):983-91.

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

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