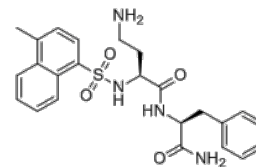


**Product Name** : J-2156  
**Cat. No.** : PC-49080  
**CAS No.** : 848647-56-3  
**Molecular Formula** : C<sub>24</sub>H<sub>28</sub>N<sub>4</sub>O<sub>4</sub>S  
**Molecular Weight** : 468.572  
**Target** : Somatostatin Receptor  
**Solubility** : 10 mM in DMSO



## Biological Activity

J-2156 is a highly potent, selective somatostatin **SST4 receptor** agonist with binding K<sub>i</sub> value of 1.2 nM (hSST4R), >400-fold selectivity over other human somatostatin receptor subtypes.

J-2156 acts as a apparent full agonist by completely inhibiting the forskolin-stimulated adenylyl cyclase activity with EC<sub>50</sub> of 70 nM.

J-2156 (1-100 microg/kg i.p.) inhibited nocifensive behaviour of mice in the second phase of the formalin test.

J-2156 inhibited sciatic nerve ligation-induced neuropathic mechanical hyperalgesia.

## References

Engström M, et al. *J Pharmacol Exp Ther*. 2005 Jan;312(1):332-8.

Engström M, et al. *J Pharmacol Exp Ther*. 2006 Mar;316(3):1262-8.

Sándor K, et al. *Eur J Pharmacol*. 2006 Jun 6;539(1-2):71-5.

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

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