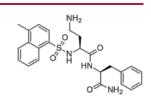


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**Data Sheet** 

Global Supplier of Chemical Probes, Inhibitors & Agonists.

| 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 Ki 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 EC50 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! E-mail: tech@probechem.com