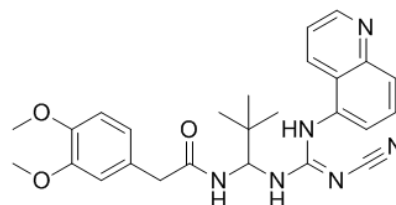


**Product Name** : A-740003  
**Cat. No.** : PC-45846  
**CAS No.** : 861393-28-4  
**Molecular Formula** : C<sub>26</sub>H<sub>30</sub>N<sub>6</sub>O<sub>3</sub>  
**Molecular Weight** : 474.5548  
**Target** : P2X Receptor  
**Solubility** : 10 mM in DMSO



## Biological Activity

A potent, selective, competitive P2X7 receptor antagonist with IC<sub>50</sub> of 40 nM and 18 nM for hP2X7 and rP2X7, respectively; showed weak or no activity for other P2 receptors (IC<sub>50</sub>>10 μM); potently blocks agonist-evoked IL-1β release (IC<sub>50</sub>=156 nM) and pore formation (IC<sub>50</sub>=92 nM) in differentiated human THP-1 cells; produces dose-dependent antinociception in a spinal nerve ligation model (ED<sub>50</sub>=19 mg/kg i.p.) in the rat.

Pain

Discontinued

## References

Honore P, et al. J Pharmacol Exp Ther. 2006 Dec;319(3):1376-85.

Morytko MJ, et al. Bioorg Med Chem Lett. 2008 Mar 15;18(6):2093-6.

King BF. Br J Pharmacol. 2007 Jul;151(5):565-7.

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

E-mail: tech@probechem.com