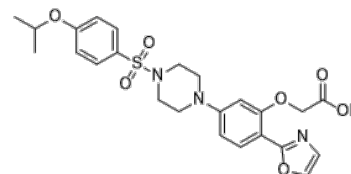


**Product Name** : Asapiprant  
**Cat. No.** : PC-22638  
**CAS No.** : 932372-01-5  
**Molecular Formula** : C<sub>24</sub>H<sub>27</sub>N<sub>3</sub>O<sub>7</sub>S  
**Molecular Weight** : 501.55  
**Target** : Prostaglandin Receptor  
**Solubility** : 10 mM in DMSO



CAS: 932372-01-5

## Biological Activity

Asapiprant (S-555739) is a potent and selective prostaglandin DP1 receptor antagonist with K<sub>i</sub> of 0.44 nM, >300-fold selective over EP2 and other prostanoid receptors.

Asapiprant (S-555739) strongly inhibits the cAMP elevation elicited by PGD<sub>2</sub> in human platelets with IC<sub>50</sub> of 16 nM, 50-fold than another DP1 antagonist, S-5751 (840 nM).

Asapiprant (S-555739) inhibits cAMP elevation induced by PGD<sub>2</sub> in guinea pigs, rats, and sheep with IC<sub>50</sub> of 61, 74, and 15, respectively.

Asapiprant (S-555739) (3, 5 mg/kg) inhibits PGD<sub>2</sub>-induced nasal resistance (sRaw) in guinea pigs.

Asapiprant (S-555739) suppresses antigen-induced asthmatic responses, airway hyper-responsiveness, and cell infiltration and mucin production in lung.

## References

Takahashi G, et al. Eur J Pharmacol. 2015 Oct 15;765:15-23.

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

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