

# **Tolvaptan** Catalog No: tcsc0572

Available Sizes

Size: 10mg

Size: 50mg

**Specifications** 

**CAS No:** 150683-30-0

### Formula:

 $C_{26}H_{25}CIN_2O_3$ 

**Pathway:** GPCR/G Protein;Autophagy

### **Target:**

Vasopressin Receptor; Autophagy

#### Purity / Grade:

>98%

## Alternative Names:

OPC-41061

**Observed Molecular Weight:** 

448.94

## **Product Description**

Tolvaptan is a selective, competitive arginine vasopressin receptor 2 antagonist with an IC50 of 1.28µM for the inhibition of AVPinduced platelet aggregation.

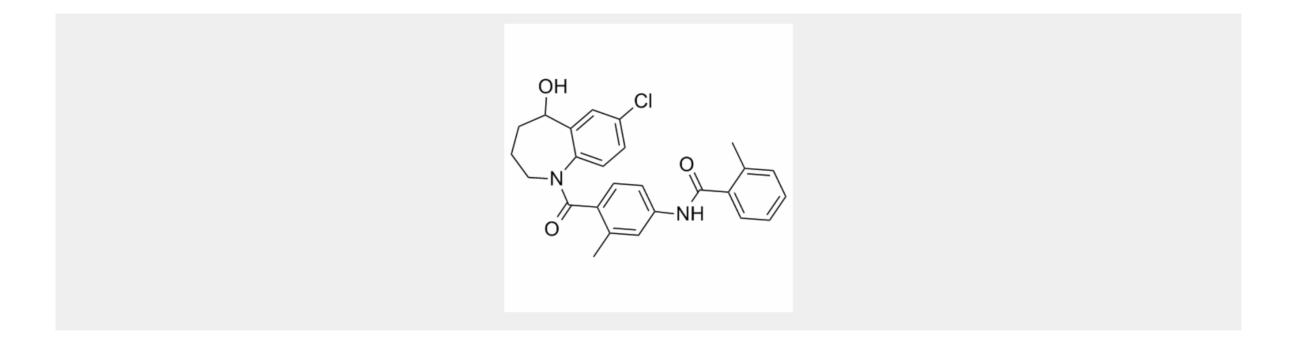
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IC50 value: 1.28 uM (inhibition of AVP-induced platelet aggregation)

#### Target: vasopressin receptor 2

Tolvaptan (OPC-41061) is a selective, competitive arginine vasopressin receptor 2 antagonist with an IC50 of 1.28µM for the inhibition of AVP-induced platelet aggregation. Tolvaptan (OPC-41061) is used to treat hyponatremia (low blood sodium levels) associated with congestive heart failure, cirrhosis, and the syndrome of inappropriate antidiuretic hormone (SIADH). Tolvaptan (OPC-41061) is also in fast-track clinical trials for polycystic kidney disease. Treatment with t tolvaptan (OPC-41061) causes rapid and sustained body weight reductions concurrent with increases in urine output, improves and/or normalizes serum sodium in hyponatremic patients, reduces signs and symptoms of congestion and increases thirst. However, tolvaptan (OPC-41061) has not been shown to decrease heart failure re-hospitalization or mortality. As an adjunct to standard therapy, tolvaptan (OPC-41061) is unique in that it is virtually the only novel agent tested in patients hospitalized for acute heart failure syndrome (AHFS) to reach its primary end point for short-term efficacy without causing deleterious side effects.



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