

# Ziprasidone

Catalog No: tcsc1071

Available Sizes

Size: 10mg

Size: 50mg

**Specifications** 

**CAS No:** 146939-27-7

Formula:

C<sub>21</sub>H<sub>21</sub>CIN<sub>4</sub>OS

**Pathway:** GPCR/G Protein;Neuronal Signaling;Neuronal Signaling;GPCR/G Protein

#### **Target:**

Dopamine Receptor; Dopamine Receptor; 5-HT Receptor; 5-HT Receptor

#### Purity / Grade:

>98%

### Solubility: DMSO : 13.5 mg/mL (32.69 mM; Need ultrasonic)

# Alternative Names:

CP-88059

**Observed Molecular Weight:** 

412.94

## **Product Description**

Ziprasidone(CP88059) is a combined 5-HT (serotonin) and dopamine receptor antagonist which exhibits potent effects of antipsychotic activity.

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IC50 value:

Target: 5-HT receptor; Dopamine receptor

Ziprasidone possesses an in vitro 5-HT2A/dopamine D2 receptor affinity ratio higher than any clinically available antipsychotic agent. In vivo, ziprasidone antagonizes 5-HT2A receptor-induced head twitch with 6-fold higher potency than for blockade of damphetamine-induced hyperactivity, a measure of central dopamine D2 receptor antagonism. Ziprasidone also has high affinity for the 5-HT1A, 5-HT1D and 5-HT2C receptor subtypes, which may further enhance its therapeutic potential [1]. Ziprasidone sulfoxide and sulfone were the major metabolites in human serum. The affinities of the sulfoxide and sulfone metabolites for 5-HT2 and D2 receptors are low with respect to ziprasidone, and are thus unlikely to contribute to its antipsychotic effects [2]. Ziprasidone was associated with significant differential adverse effects relative to placebo in BPM, BPD, and schizophrenia with no significant difference in weight gain in all 3 groups. Self-reported somnolence was increased across the 3 conditions. Subjects with BPM were more vulnerable to EPS than those with BPD or schizophrenia [3].

Clinical indications: Bipolar I disorder; Bipolar disorder; Mania; Schizophrenia

FDA Approved Date: February 2001



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