



## Haloperidol

Catalog No: tcsc1971



## **Available Sizes**

Size: 100mg

Size: 500mg



## **Specifications**

**CAS No:** 

52-86-8

Formula:

 $C_{21}H_{23}CIFNO_2$ 

**Pathway:** 

GPCR/G Protein; Neuronal Signaling

**Target:** 

Dopamine Receptor; Dopamine Receptor

**Purity / Grade:** 

>98%

**Solubility:** 

DMSO : ≥ 31 mg/mL (82.48 mM)

**Observed Molecular Weight:** 

375.86

## **Product Description**

Haloperidol is a potent dopamine D2 receptor antagonist, widely used as an antipsychotic.

In Vivo: Haloperidol (1 mg) intra-arterially attenuates the dopamine-induced pancreatic secretion. Haloperidol (3 mg) completely inhibits the action of 10  $\mu$ g of dopamine in the pancreas of the dogs<sup>[1]</sup>. Haloperidol (10 mg/kg) as well as chlorpromazine (CPZ, 15 mg/kg) blocks mescaline-induced altered behavior within 7 to 10 minutes when injected into the mice 45 minutes after 50 mg/kg (2  $\mu$ c) of mescaline. Haloperidol has no effect on mescaline disappearance<sup>[2]</sup>.





All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!