## TOCR a biotechne

## **Certificate of Analysis**

## www.tocris.com

Print Date: Mar 10th 2022

#### Product Name: Mirtazapine

CAS Number: 85650-52-8

IUPAC Name:

## 1. PHYSICAL AND CHEMICAL PROPERTIES

**Batch Molecular Formula: Batch Molecular Weight:** Physical Appearance: Solubility:

 $C_{17}H_{19}N_3.1/_4H_2O$ 269.86 White solid ethanol to 50 mM DMSO to 20 mM Store at RT

Storage: **Batch Molecular Structure:** 

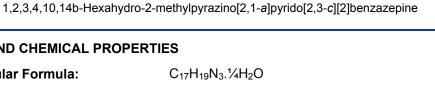
## 2. ANALYTICAL DATA

TLC: HPLC: <sup>1</sup>H NMR: Microanalysis: R<sub>f</sub> = 0.24 (Dichloromethane:Methanol:Acetic acid [9:1:0.1]) Shows 100% purity Consistent with structure

	Carbon Hydrogen Nitrogen				
Theoretical	75.66	7.28	15.57		
Found	75.36	7.35	15.56		

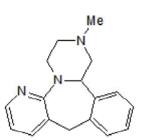
Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

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Catalog No.: 2018

EC Number: 288-060-6



Batch No.: 2

# TOCRIS a biotechne brand

## **Product Information**

## www.tocris.com

Batch No.: 2

### Product Name: Mirtazapine

CAS Number: 85650-52-8

IUPAC Name:

1,2,3,4,10,14b-Hexahydro-2-methylpyrazino[2,1-a]pyrido[2,3-c][2]benzazepine

### **Description:**

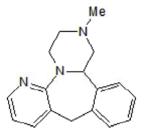
Mirtazapine is an antidepressant agent; potent 5-HT<sub>2</sub>, 5-HT<sub>3</sub> and histamine H<sub>1</sub> receptor antagonist and moderately potent  $\alpha_2$ -adrenoceptor antagonist (pK<sub>i</sub> values are 8.05, ~ 8.1, 9.3 and 6.95 respectively). Enhances noradrenaline (NA) release in rat brain via inhibition of  $\alpha_2$ -adrenergic autoreceptors and displays only weak affinity for monoamine transporters (pK<sub>i</sub> values are 5.6, < 5 and < 5.1 for inhibition of NA, dopamine and 5-HT uptake respectively). Increases hippocampal NA and 5-HT levels in rats following systemic administration in vivo.

### **Physical and Chemical Properties:**

Batch Molecular Formula: C<sub>17</sub>H<sub>19</sub>N<sub>3</sub>.¼H<sub>2</sub>O Batch Molecular Weight: 269.86 Physical Appearance: White solid

Minimum Purity: ≥99%

#### **Batch Molecular Structure:**



#### **References:**

**de Boer** *et al* (1996) Differences in modulation of noradrenergic and serotonergic transmission by the alpha-2 adrenoceptor antagonists, mirtazepine, mianserin and idazoxan. J.Pharmacol.Exp.Ther. **277** 852. PMID: 8627567.

Kooyman et al (1994) Interaction between enantiomers of mianserin and ORG3770 at 5-HT3 receptors in cultured mouse neuroblastoma cells. Neuropharmacology 33 501. PMID: 7984289.

de Boer et al (1988) Neurochemical and autonomic pharmacological profiles of the 6-aza-analogue of mianserin, ORG 3770 and its enantiomers. Neuropharmacology 27 399. PMID: 3419539.

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### Storage: Store at RT

**Solubility & Usage Info:** ethanol to 50 mM

DMSO to 20 mM

#### **Stability and Solubility Advice:**

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a  $45-60^{\circ}C$  water bath).

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Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. Our standard recommendations are:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.