

# **Data Sheet**

Product Name:	Azacyclonol	$\land$
Cat. No.:	CS-2754	
CAS No.:	115-46-8	
Molecular Formula:	C <sub>18</sub> H <sub>21</sub> NO	~ Ѷ∩н
Molecular Weight:	267.37	
Target:	Histamine Receptor	$\langle \rangle$
Pathway:	GPCR/G Protein; Immunology/Inflammation; Neuronal Signaling	
Solubility:	DMSO : 100 mg/mL (374.01 mM; Need ultrasonic)	N H

# **BIOLOGICAL ACTIVITY:**

Azacyclonol (y-pipradol), a metabolite of Terfenadine, is a central depressant agent. Azacyclonol is a ganglion-blocking agent. Azacyclonol can be used to diminish psychoses-induced hallucinations<sup>[1][2][3]</sup>. In Vitro: Azacyclonol is formed from Terfenadine in rat liver<sup>[2]</sup>. In Vivo: Azacyclonol causes depressed activity in mice and rats<sup>[3]</sup>.

Azacyclonol antagonizes increased coordination activity in mice induced by pipradrol, amphetamine, morphine and cocaine and prolongs Hexobarbital hypnosis<sup>[3]</sup>.

## **References:**

[1]. Brown DA, et, al. The effects of some centrally acting drugs on ganglionic transmission in the cat.

[2]. Jurima-Romet M, et, al. Induction of CYP3A and associated terfenadine N-dealkylation in rat hepatocytes cocultured with 3T3 cells. Cell Biol Toxicol. 1995 Dec;11(6):313-27.

[3]. BRAUN DL, et, al. The pharmacologic activity of alpha-(4-piperidyl)-benzhydrol hydrochloride (azacyclonol hydrochloride); an ataractive agent. J Pharmacol Exp Ther. 1956 Oct;118(2):153-61.

# CAIndexNames:

4-Piperidinemethanol, α,α-diphenyl-

## SMILES:

OC(C1=CC=CC=C1)(C2CCNCC2)C3=CC=CC=C3

## Caution: Product has not been fully validated for medical applications. For research use only.

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