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### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

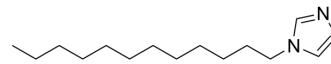
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## 1-Dodecylimidazole

Cat. No.:	HY-138540		
CAS No.:	4303-67-7		
Molecular Formula:	$C_{15}H_{28}N_2$		
Molecular Weight:	236.4		
Target:	Fungal		
Pathway:	Anti-infection		
Storage:	Pure form	-20°C	3 years
		4°C	2 years
In solvent	-80°C	6 months	
	-20°C	1 month	



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 100 mg/mL (423.01 mM; Need ultrasonic)

Preparing Stock Solutions	Concentration	Solvent Mass		
		1 mg	5 mg	10 mg
	1 mM	4.2301 mL	21.1506 mL	42.3012 mL
	5 mM	0.8460 mL	4.2301 mL	8.4602 mL
	10 mM	0.4230 mL	2.1151 mL	4.2301 mL

Please refer to the solubility information to select the appropriate solvent.

#### In Vivo

1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline  
Solubility:  $\geq 2.5 \text{ mg/mL}$  (10.58 mM); Clear solution
2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- $\beta$ -CD in saline)  
Solubility:  $\geq 2.5 \text{ mg/mL}$  (10.58 mM); Clear solution
3. Add each solvent one by one: 10% DMSO >> 90% corn oil  
Solubility:  $\geq 2.5 \text{ mg/mL}$  (10.58 mM); Clear solution

### BIOLOGICAL ACTIVITY

Description	1-Dodecylimidazole (N-Dodecylimidazole) is a lysosomotropic detergent and a cytotoxic agent. 1-Dodecylimidazole causes cell death by its acid-dependent accumulation in lysosomes, disruption of the lysosomal membrane, and release of cysteine proteases into the cytoplasm. 1-Dodecylimidazole has hypocholesterolaemic activity and broad-spectrum antifungal activity <sup>[1][2][3]</sup> .
In Vitro	N-dodecylimidazole, an acid activated detergent with a pKa of 6.3, has been shown to be cytotoxic to cells in culture. N-dodecylimidazole displayed extracellular pH (pHe)-dependent cytotoxicity against EMT-6 and MGH U1 cells. cell killing was

dose dependent and was 100-fold greater at pH 6.0 than pH 7.0<sup>[4]</sup>.  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### In Vivo

The hypocholesterolaemic activity of 1-dodecylimidazole results in part from the inhibition of cholesterol biosynthesis at the level of 2,3-oxidosqualene sterol cyclase<sup>[2]</sup>.

1-dodecylimidazole (150 mg/kg body wt; by stomach tube; daily for 10 days) has lower serum cholesterol concentrations than control rats<sup>[2]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male rats <sup>[2]</sup>
Dosage:	150 mg/kg body wt
Administration:	By stomach tube; daily for 10 days
Result:	Had significantly lower serum cholesterol concentrations than untreated animals.

#### REFERENCES

- [1]. Wilson PD, et al. A relationship between multidrug resistance and growth-state dependent cytotoxicity of the lysosomotropic detergent N-dodecylimidazole. Biochem Biophys Res Commun. 1991;176(3):1377-1382.
- [2]. Atkin SD, et al. The isolation of 2,3-oxidosqualene from the liver of rats treated with 1-dodecylimidazole, a novel hypocholesterolaemic agent. Biochem J. 1972;130(1):153-157.
- [3]. Firestone RA, et al. Lysosomotropic agents. 7. Broad-spectrum antifungal activity of lysosomotropic detergents. J Med Chem. 1987;30(8):1519-1521.
- [4]. Boyer MJ, et al. pH dependent cytotoxicity of N-dodecylimidazole: a compound that acquires detergent properties under acidic conditions. Br J Cancer. 1993;67(1):81-87.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA