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[Anti-infection](#) [Fungal](#) [Terbinafine](#)

## Terbinafine (Synonyms: TDT 067)

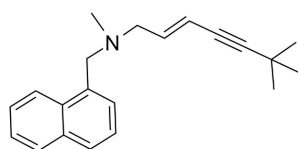
Cat. No.: HY-17395A

Purity: 99.91%

[Data Sheet](#)[SDS](#)[Handling Instructions](#)

Terbinafine (TDT 067) is an orally active and potent **antifungal** agent. Terbinafine is a potent non-competitive inhibitor of **squalene epoxidase** from *Candida*, with a  $K_i$  of 30 nM. Terbinafine also shows antibacterial activity against certain Gram-positive and Gram-negative **bacteria**.

**For research use only. We do not sell to patients.**



Terbinafine Chemical Structure  
CAS No. : 91161-71-6

Size	Price	Stock	Quantity
<b>Free Sample</b> (0.5-1 mg)		<a href="#">Apply Now</a>	
<b>Solution</b>			
10 mM * 1 mL in DMSO	USD 73	In-stock	<input type="text" value="0"/>
<b>Solid + Solvent</b>			
10 mM * 1 mL ready for reconstitution	USD 73	In-stock	<input type="text" value="0"/>
<b>Solid</b>			
100 mg	USD 66	In-stock	<input type="text" value="0"/>
200 mg	USD 117	In-stock	<input type="text" value="0"/>
500 mg		<a href="#">Get quote</a>	
1 g		<a href="#">Get quote</a>	

### Customer Review

Based on 2 publication(s) in Google Scholar

### Other Forms of Terbinafine:

Terbinafine hydrochloride	In-stock
Terbinafine lactate	Get quote
Terbinafine-d3 hydrochloride	Get quote

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### 2 Publications Citing Use of MCE Terbinafine

- *Cell Death Dis.* 2021 May 13;12(5):482.

- *Cancer Commun (Lond).* 2021 Jul 16.



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Cancer Communications | 2021 Jul 16 | PubMed ID: 34268906 | [Read Article](#)

Squalene epoxidase promotes colorectal cancer cell proliferation through accumulating calcitriol and activating CYP24A1-mediated MAPK signaling, et al. Squalene epoxidase promotes colorectal cancer cell proliferation through accumulating calcitriol and activating CYP24A1-mediated MAPK signaling

Luwei He, Huaguang Li, ..., Moubin Lin

**Article Snippet**

"... mice were sacrificed by cervical dislocation before the volume of tumors reached 1000 mm<sup>3</sup>. For **terbinafine** (CatHY17395A; **MedChemExpress**, Shanghai, China) treatment, HT29 and RKO cells (4×10<sup>6</sup> cells in 0.1 mL PBS) were subcutaneously injected into the left flank of nude mice. Once tumor volumes reached 30 mm<sup>3</sup>, these mice were divided into control group (PBS ...)" ([More...](#))

**Figure Legend**

"**Terbinafine** suppressed the growth of CRC organoids and xenograft ..." ([More...](#))

Cancer Science | 2022 Jan 06 | PubMed ID: 34939274 | [Read Article](#)

Squalene synthase predicts poor prognosis in stage I-III colon adenocarcinoma and synergizes squalene epoxidase to promote tumor progression, et al. Squalene synthase predicts poor prognosis in stage I-III colon adenocarcinoma and synergizes squalene epoxidase to promote tumor progression

Huihong Jiang, Erjiang Tang, ..., Luwei He

**Article Snippet**

"2.1 Reagents D-pantethine (abs816989) was purchased from absin. Lapaquistat (HY-14925) and **terbinafine** (HY-17395A) were purchased from **MCE**. N -acetyl-L-cysteine (S0077) was purchased from Beyotime." ([More...](#))

**Figure Legend**

"... HT29 cells were treated with lapaquistat (La) and **terbinafine** (Te) at different concentrations for 48 or 72 ..." ([More...](#))

Cell Death & Disease | 2021 May 13 | PubMed ID: 33986254 | [Read Article](#)

Targeting epigenetic modulation of cholesterol synthesis as a therapeutic strategy for head and neck squamous cell carcinoma

Xing Xu, Jun Chen, ..., Xu Wang

**Article Snippet**

"Chemical compoundsThe following compounds were purchased from commercial vendors: GSK126 (S7061, Selleck, China), GSK343 (S7164, Selleck, China), EPZ6438 (S7168, Selleck, China), NB-598 (HY-16343C, MCE, China), Atorvastatin (HY-17379, MCE, China), Ro48-8071 (HY-18630A, MCE, China), **Terbinafine** (HY-17395, **MCE**, China), Butenafine (HY-17396, MCE, China), Cholesterol (C3045, Sigma, China), Desmosterol (H130206, Aladdin, China), Lathosterol (HY-17395, MCE, China), Squalene (S3626, Sigma, China)." ([More...](#))

**Figure Legend**

"... treatment groups. Cells were treated with vehicle, EPZ6438, **terbinafine** or the combination for 48 h. Bar = ..." ([More...](#))



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Biological Activity	Protocol	Purity & Documentation	References	Customer Review																													
<b>Description</b>	Terbinafine (TDT 067) is an orally active and potent <b>antifungal</b> agent. Terbinafine is a potent non-competitive inhibitor of <b>squalene epoxidase</b> from <i>Candida</i> , with a $K_i$ of 30 nM. Terbinafine also shows antibacterial activity against certain Gram-positive and Gram-negative <b>bacteria</b> <sup>[1] [2] [3]</sup> .																																
<b>IC<sub>50</sub> &amp; Target</b>	K <sub>i</sub> : 30 nM (squalene epoxidase) <sup>[1]</sup>																																
<b>In Vitro</b>	Terbinafine has a primary fungicidal action <i>in vitro</i> against most fungal pathogens, including dermatophytes, and dimorphic and filamentous fungi. Terbinafine specifically inhibits fungal ergosterol biosynthesis at the point of squalene epoxidation. The treated fungal cells rapidly accumulate tlic intermediate squalene and become deficient in the end-product of the pathway, ergosterol <sup>[1]</sup> . <b>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</b>																																
<b>In Vivo</b>	Terbinafine is not only active after topical application but is very effective in experimental dermatophytoses following oral administration. In fungi infected guinea-pigs, the skin temperature dropps dramatically after the fourth treatment of terbinafine <sup>[2]</sup> . <b>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</b>																																
<b>Clinical Trial</b>	<b>NCT Number</b>	<b>Sponsor</b>	<b>Condition</b>	<b>Start Date</b>	<b>Phase</b>																												
	<a href="#">NCT01433107</a>	Novartis	Tinea Pedis	August 2011	Phase 3																												
	<a href="#">NCT03171584</a>	Assiut University	Antifungal Drugs in Onychomycosis	July 1, 2017	Phase 3																												
	<a href="#">NCT00443898</a>	Novartis Pharmaceuticals Novartis	Onychomycosis	December 2006	Phase 3																												
	<a href="#">View More</a> ▾																																
<b>Molecular Weight</b>	291.43																																
<b>Formula</b>	C <sub>21</sub> H <sub>25</sub> N																																
<b>CAS No.</b>	91161-71-6																																
<b>SMILES</b>	CN(C/C=C/C#CC(C)(C)C)CC1=C2C=CC=CC2=CC=C1																																
<b>Shipping</b>	Room temperature in continental US; may vary elsewhere.																																
<b>Storage</b>	Powder	-20°C	3 years																														
		4°C	2 years																														
	In solvent	-80°C	6 months																														
		-20°C	1 month																														
<b>Solvent &amp; Solubility</b>	<p><b>In Vitro:</b></p> <p><b>DMSO : ≥ 100 mg/mL (343.14 mM)</b></p> <p>* "≥" means soluble, but saturation unknown.</p> <table border="1"> <thead> <tr> <th rowspan="2">Preparing Stock Solutions</th> <th>Solvent</th> <th>Mass</th> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> <tr> <th colspan="5">Concentration</th> </tr> </thead> <tbody> <tr> <td></td> <td>1 mM</td> <td></td> <td>3.4314 mL</td> <td>17.1568 mL</td> <td>34.3136 mL</td> </tr> <tr> <td></td> <td>5 mM</td> <td></td> <td>0.6863 mL</td> <td>3.4314 mL</td> <td>6.8627 mL</td> </tr> <tr> <td></td> <td>10 mM</td> <td></td> <td>0.3431 mL</td> <td>1.7157 mL</td> <td>3.4314 mL</td> </tr> </tbody> </table> <p>* Please refer to the solubility information to select the appropriate solvent.</p> <p><b>In Vivo:</b></p>				Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg	Concentration						1 mM		3.4314 mL	17.1568 mL	34.3136 mL		5 mM		0.6863 mL	3.4314 mL	6.8627 mL		10 mM		0.3431 mL	1.7157 mL	3.4314 mL
Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg		10 mg																											
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3. Add each solvent one by one: **10% DMSO** » **90% (20% SBE-β-CD in saline)**  
Solubility: 2.5 mg/mL (8.58 mM); Suspended solution; Need ultrasonic
4. Add each solvent one by one: **10% DMSO** » **90% corn oil**  
Solubility: 2.5 mg/mL (8.58 mM); Clear solution; Need ultrasonic
- \* All of the co-solvents are available by MCE.

**Purity & Documentation**

Purity: 99.91%

Select Batch: [Data Sheet \(276 KB\)](#)[SDS \(393 KB\)](#)[COA \(248 KB\)](#)[LCMS \(267 KB\)](#)[Handling Instructions \(2659 KB\)](#)**References**

- [1]. Ryder NS, et al. Terbinafine: mode of action and properties of the squalene epoxidase inhibition. Br J Dermatol. 1992 Feb;126 Suppl 39:2-7.
- [2]. Mieth H, et al. Preclinical evaluation of terbinafine in vivo. Clin Exp Dermatol. 1989 Mar;14(2):104-7.
- [3]. Ciftci E, et al. Mupirocin vs terbinafine in impetigo. Indian J Pediatr. 2002 Aug;69(8):679-82.

[Molarity Calculator](#)[Dilution Calculator](#)**The molarity calculator equation**

$$\text{Mass (g)} = \text{Concentration (mol/L)} \times \text{Volume (L)} \times \text{Molecular Weight (g/mol)}$$

Mass

mg

Concentration

mM

Volume

mL

Molecular Weight \*

Calculate

Reset

**Keywords:** Terbinafine 91161-71-6 TDT 067 TDT067 TDT-067 Fungal Bacterial Antibiotic infections bacteria  
Candida squalene epoxidase antibacterial activity Inhibitor inhibitor inhibit

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Specific Active Calculator

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