

<b>Product Name:</b>	Clotrimazole
<b>Product Number:</b>	C037
<b>CAS Number:</b>	23593-75-1
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>17</sub> ClN <sub>2</sub>
<b>Molecular Weight:</b>	344.84
<b>Form:</b>	Powder
<b>Appearance:</b>	White crystalline powder
<b>Solubility:</b>	freely soluble in water (30 mg/mL)
<b>Source:</b>	Synthetic
<b>Melting Point:</b>	141- 145°C
<b>Storage Conditions:</b>	2-8°C
<b>Description:</b>	Clotrimazole is a broad-spectrum antifungal. It falls into the imidazole subclass of azole compounds, which interfere with the biosynthesis of ergosterol, a major membrane component of the fungal cytoplasmic membrane. It was discovered in 1969 and was developed by Schering Plough. It inhibits Ca <sup>2+</sup> -activated potassium channels. The compound has promising anti-cancer effects. It is an CYP450 enzyme inhibitor. It is freely soluble in water.
<b>Mechanism of Action:</b>	Clotrimazole increases fungal cell permeability by inhibiting ergosterol synthesis, a major cell membrane component found exclusively in fungi, thus is fungistatic and inhibits fungal growth. Specifically, it inhibits the microsomal cytochrome P450-dependent 14α-demethylase, which is critical to ergosterol biosynthesis.
<b>Spectrum:</b>	Clotrimazole is broad-spectrum, targeting a broad range of fungi including <i>Candida</i> and <i>Aspergillus</i> species.
<b>Cancer Applications</b>	Clotrimazole has promising anti-cancer effects, interfering with glycolytic enzymes, specifically their cellular distribution and their activity. Cell lines from human breast tissue (MCF10A, MCF-7 and MDA-MB-231) were used, and Clotrimazole induced a dose-dependent decrease in glucose uptake in all three cell lines, affecting the metabolism, growth, and migration of human breast cancer cell lines. It was non-toxic to non-tumor human breast cell lines (Furtado et al, 2012).

**References:**

- Crowley PD and Gallagher HC (2014) Clotrimazole as a pharmaceutical: Past, present, and future. *J. Appl. Microbiol.* 117(3):611-617
- Furtado CM, Marcondes MC, Sola-Penna M, de Souza MLS, and Zancan P (2012) Clotrimazole preferentially inhibits human breast cancer cell proliferation, viability and glycolysis. *PLoS ONE* 7(2): e30462
- Jensen BS, Strøbaek D, Olesen SP, Christophersen P (2001) The  $\text{Ca}^{2+}$ -activated  $\text{K}^{+}$  channel of intermediate conductance: a molecular target for novel treatments? *Curr Drug Targets.* 2(4):401-422 PMID 11732639
- Rice LB and Ghannoum MA (1999) Antifungal agents: Mode of action, mechanisms of resistance, and correlation of these mechanisms with bacterial resistance. *Clin. Microbiol. Rev.* 12(4):501-517 PMID 10515900
- Yan Z, Rafferty B, Caldwell GW and MASucci JA (2002) Rapidly distinguishing reversible and irreversible CYP450 inhibitors by using fluorometric kinetic analyses. *E. J. Drug Metab. And Pharmacokin.* 27(4):281-287

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