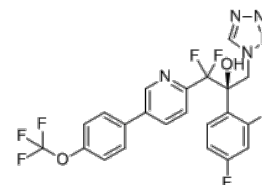


**Product Name** : Quilseconazole  
**Cat. No.** : PC-60051  
**CAS No.** : 1340593-70-5  
**Molecular Formula** : C<sub>22</sub>H<sub>14</sub>F<sub>7</sub>N<sub>5</sub>O<sub>2</sub>  
**Molecular Weight** : 513.376  
**Target** : Fungal  
**Solubility** : 10 mM in DMSO



## Biological Activity

Quilseconazole (VT-1129) is a selective, orally available **fungal CYP51** (lanosterol 14- $\alpha$ -demethylase) inhibitor. Quilseconazole displays greater selectivity for fungal enzyme than for mammalian CYP450 compared to the approved azole class of antifungal drugs. Quilseconazole inhibits *C. albicans* and *T. rubrum* with MIC of <0.001  $\mu$ g/mL, also demonstrated potent activities against *Cryptococcus* species as demonstrated by low MIC<sub>50</sub> and MIC<sub>90</sub> values.

## References

- Hoekstra WJ, et al. *Bioorg Med Chem Lett*. 2014 Aug 1;24(15):3455-8.
- Lockhart SR, et al. *Antimicrob Agents Chemother*. 2016 Mar 25;60(4):2528-31.
- Warrilow AG, et al. *Antimicrob Agents Chemother*. 2016 Jul 22;60(8):4530-8.
- Nielsen K, et al. *Med Mycol*. 2017 Jun 1;55(4):453-456.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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