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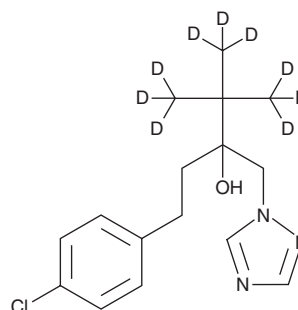
PRODUCT INFORMATION



Tebuconazole-d₉

Item No. 33287

CAS Registry No.: 1246818-83-6
Formal Name: α-[2-(4-chlorophenyl)ethyl]-α-(1,1-dimethylethyl-d₉)-1H-1,2,4-triazole-1-ethanol
MF: C₁₆H₁₃ClD₉N₃O
FW: 316.9
Chemical Purity: ≥98% (Tebuconazole)
Deuterium Incorporation: ≥99% deuterated forms (d₁-d₉); ≤1% d₀
Supplied as: A solid
Storage: -20°C
Stability: ≥2 years



Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Tebuconazole-d₉ is intended for use as an internal standard for the quantification of tebuconazole (Item No. 24052) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

Tebuconazole-d₉ is supplied as a solid. A stock solution may be made by dissolving the tebuconazole-d₉ in the solvent of choice, which should be purged with an inert gas. Tebuconazole-d₉ is soluble in methanol, DMSO, and acetonitrile.

Description

Tebuconazole is a triazole fungicide that is active against both seed and foliar fungi.¹ It inhibits 14α-demethylase isolated from *U. maydis* and *S. bicolor* with IC₅₀ values of 0.05 and 0.16 nM, respectively.² It inhibits the androgenic effect of the androgen receptor agonist DHT (IC₂₀ = 2.89 μM) and is cytotoxic (EC₂₀ = 38.9 μM) in an MDA-kb2 assay.³ Tebuconazole (50 and 100 mg/kg per day) administered during gestation reduces testosterone levels and increases testicular levels of progesterone and 17α-hydroxyprogesterone in male rat fetuses.⁴ It has a feminizing effect on male pups and a virilizing effect on female pups. When administered to rats gestationally through postnatal day 42, tebuconazole (20 and 60 mg/kg per day) leads to cell death of pyramidal cells in the CA3-4 region of the hippocampus and layer V of the cortex concomitant with impairment in learning the platform location in the Morris water maze.⁵

References

1. Reinecke, P., Kaspers, H., Scheinplug, H., et al. BAY HWG 1608, a new fungicide for foliar spray and seed-treatment use against a wide spectrum of fungal pathogens. *Br. Crop Prot. Conf.-Pests Dis., Proc.* **1**, 41-46 (1986).
2. Zarn, J.A., Brüschweiler, B.J., and Schlatter, J.R. Azole fungicides affect mammalian steroidogenesis by inhibiting sterol 14 α-demethylase and aromatase. *Environ. Health Perspect.* **111**(3), 255-261 (2003).
3. Orton, F., Rosivatz, E., Scholze, M., et al. Widely used pesticides with previously unknown endocrine activity revealed as *in vitro* antiandrogens. *Environ. Health Perspect.* **119**(6), 794-800 (2011).
4. Taxvig, C., Hass, U., Axelstad, M., et al. Endocrine-disrupting activities *in vivo* of the fungicides tebuconazole and epoxiconazole. *Toxicol. Sci.* **100**(2), 464-473 (2007).
5. Moser, V.C., Barone, S., Jr., Smialowicz, R.J., et al. The effects of perinatal tebuconazole exposure on adult neurological, immunological, and reproductive function in rats. *Toxicol. Sci.* **62**(2), 339-352 (2001).

WARNING

THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the [complete](#) Safety Data Sheet, which has been sent via email to your institution.

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