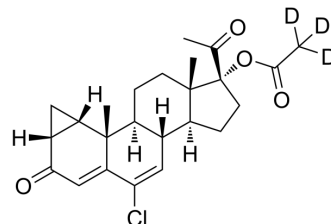


## Cyproterone acetate-d<sub>3</sub>

<b>Cat. No.:</b>	HY-13604S
<b>CAS No.:</b>	2376035-90-2
<b>Molecular Formula:</b>	C <sub>24</sub> H <sub>26</sub> D <sub>3</sub> ClO <sub>4</sub>
<b>Molecular Weight:</b>	419.96
<b>Target:</b>	Androgen Receptor; Isotope-Labeled Compounds
<b>Pathway:</b>	Vitamin D Related/Nuclear Receptor; Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Cyproterone acetate-d <sub>3</sub> is deuterium labeled Cyproterone acetate. Cyproterone acetate is an anti-androgen (IC <sub>50</sub> =7.1 nM) and progestogen synthetic steroid. Cyproterone acetate has affinity with progesteron and with glucocorticoidal receptors[1][2].
<b>In Vitro</b>	Stable heavy isotopes of hydrogen, carbon, and other elements have been incorporated into drug molecules, largely as tracers for quantitation during the drug development process. Deuteration has gained attention because of its potential to affect the pharmacokinetic and metabolic profiles of drugs <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

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- [2]. Chen L, et al. Cyproterone acetate enhances TRAIL-induced androgen-independent prostate cancer cell apoptosis via up-regulation of death receptor 5. *BMC Cancer.* 2017;17(1):179. Published 2017 Mar 7.
- [3]. Migally N. Effect of cyproterone acetate on the structure of the adrenal cortex. *Arch Androl.* 1979;2(2):109-115.
- [4]. Sonneveld, E., et al., Development of androgen- and estrogen-responsive bioassays, members of a panel of human cell line-based highly selective steroid-responsive bioassays. *Toxicol Sci.* 2005. 83(1): p. 136-48.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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