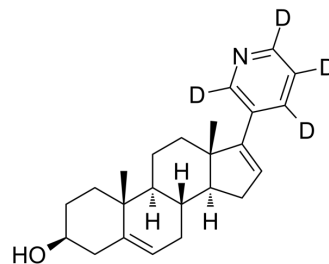


Abiraterone-D4

Cat. No.:	HY-70013S
CAS No.:	2122245-62-7
Molecular Formula:	C ₂₄ H ₂₇ D ₄ NO
Molecular Weight:	353.53
Target:	Cytochrome P450
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Abiraterone-D4 (CB-7598-D4) is the deuterium labeled Abiraterone. Abiraterone is a potent and irreversible CYP17A1 inhibitor with antiandrogen activity, which inhibits both the 17 α -hydroxylase and 17,20-lyase activity of the cytochrome p450 enzyme CYP17 with IC ₅₀ s of 2.5 nM and 15 nM, respectively.
In Vitro	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 ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

- [1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. *Ann Pharmacother.* 2019;53(2):211-216.
- [2]. Attard G, et al. Phase I clinical trial of a selective inhibitor of CYP17, abiraterone acetate, confirms that castration-resistant prostate cancer commonly remains hormone driven. *J Clin Oncol.* 2008 Oct 1;26(28):4563-71.; Richards J, et al. Interactions of abiraterone, eplerenone, and prednisolone with wild-type and mutant androgen receptor: a rationale for increasing abiraterone exposure or combining with MDV3100. *Cancer Res.* 2012 May 1;72(9):2176-82.; Stein MN, et al. Androgen synthesis inhibitors in the treatment of castration-resistant prostate cancer. *Asian J Androl.* 2014 May-Jun;16(3):387-400.; Li R, et al. Abiraterone inhibits 3 β -hydroxysteroid dehydrogenase: a rationale for increasing drug exposure in castration-resistant prostate cancer. *Clin Cancer Res.* 2012 Jul 1;18(13):3571-9.; Kumar SV, et al. Validated RP-HPLC/UV method for the quantitation of abiraterone in rat plasma and its application to a pharmacokinetic study in rats. *Biomed Chromatogr.* 2013 Feb;27(2):203-7.; Stein MN, et al. Androgen synthesis inhibitors in the treatment of castration-resistant prostate cancer. *Asian J Androl.* 2014 May-Jun;16(3):387-400.

Caution: Product has not been fully validated for medical applications. For research use only.

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