Ulipristal acetate-d₆

Cat. No.:	HY-16508S	
CAS No.:	1621894-64-1	D D_ _D
Molecular Formula:	C ₃₀ H ₃₁ D ₆ NO ₄	D, N, A
Molecular Weight:	481.66	
Target:	Progesterone Receptor; Autophagy; Isotope-Labeled Compounds	
Pathway:	Vitamin D Related/Nuclear Receptor; Autophagy; Others	
Storage:	Please store the product under the recommended conditions in the Certificate of	0~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
	Analysis.	

Description	Ulipristal acetate-d ₆ is deuterium labeled Ulipristal acetate. Ulipristal acetate (CDB-2914) is an orally active, selective progesterone receptor modulator (SPRM). Ulipristal acetate stimulates the autophagic response selectively in leiomyoma cells. Ulipristal acetate has the potential for benign gynecological conditions treatment, such as uterine myoma[1][2].	
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

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[2]. Attardi BJ, et al. In vitro antiprogestational/antiglucocorticoid activity and progestin and glucocorticoid receptor binding of the putative metabolites and synthetic derivatives of CDB-2914, CDB-4124, and mifepristone. J Steroid Biochem Mol Biol.

[3]. Ciarmela P, et al. Ulipristal acetate modulates the expression and functions of activin a in leiomyoma cells. Reprod Sci. 2014 Sep;21(9):1120-5.

[4]. Del Bello B, et al. Autophagy up-regulation by ulipristal acetate as a novel target mechanism in the treatment of uterine leiomyoma: an in vitro study. Fertil Steril. 2019 Dec;112(6):1150-1159.

[5]. Hild SA, et al. CDB-2914: anti-progestational/anti-glucocorticoid profile and post-coital anti-fertility activity in rats and rabbits. Hum Reprod. 2000 Apr; 15(4):822-9.

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[7]. Pohl O, et al. A 39-week oral toxicity study of ulipristal acetate in cynomolgus monkeys. Regul Toxicol Pharmacol. 2013 Jun;66(1):6-12.

[8]. Pohl O, et al. Carcinogenicity and chronic rodent toxicity of the selective progesterone receptor modulator ulipristal acetate. Curr Drug Saf. 2013 Apr;8(2):77-97.

Inhibitors

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Screening Libraries

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Proteins

Product Data Sheet

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

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