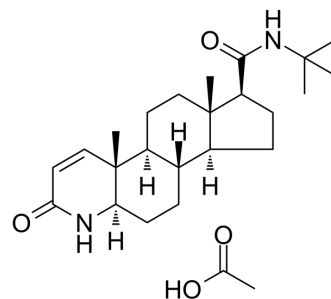


Finasteride acetate

Cat. No.:	HY-13635A
CAS No.:	222989-99-3
Molecular Formula:	C ₂₅ H ₄₀ N ₂ O ₄
Molecular Weight:	432.6
Target:	5 alpha Reductase
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Finasteride (MK-906) acetate is a potent and competitive 5 α -reductase inhibitor, with an IC ₅₀ of 4.2 nM for type II 5 α -reductase. Finasteride acetate has approximately a 100-fold greater affinity for type II 5 α -reductase enzyme than for the type I enzyme. Finasteride acetate can be used for the research of benign prostatic hyperplasia (BPH) and androgenic alopecia ^[1] ^[2] ^[3] .								
IC₅₀ & Target	IC ₅₀ : 4.2 nM (type II 5 α -reductase) ^[1]								
In Vitro	<p>Finasteride (10 μM; 6-24 h) induces the expression of HO-1 and Nrf2 proteins in PC-3 cells^[2].</p> <p>Finasteride decreases the conversion of [³H]testosterone (T) to [³H]dihydrotestosterone (DHT) in <i>P. crustosum</i>^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>PC-3, DU-145, and LNCaP cells</td> </tr> <tr> <td>Concentration:</td> <td>10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>6, 12, 24 h</td> </tr> <tr> <td>Result:</td> <td>Increased the expression of HO-1 protein in a time-dependent manner in PC-3 cells. Induced the expression of Nrf2 protein in DU-145 and PC-3 cells, but not in LNCaP cells.</td> </tr> </table>	Cell Line:	PC-3, DU-145, and LNCaP cells	Concentration:	10 μ M	Incubation Time:	6, 12, 24 h	Result:	Increased the expression of HO-1 protein in a time-dependent manner in PC-3 cells. Induced the expression of Nrf2 protein in DU-145 and PC-3 cells, but not in LNCaP cells.
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In Vivo	<p>Finasteride (0.1-0.5 mg/kg; p.o. once daily for 16 weeks) reduces prostatic size in dogs with BPH without adversely affecting semen quality or serum testosterone concentration^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Male dogs with spontaneous BPH (2.7-11 years old; 10.3-49 kg)^[3]</td> </tr> <tr> <td>Dosage:</td> <td>0.1-0.5 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>P.o. once daily for 16 weeks</td> </tr> <tr> <td>Result:</td> <td>Decreased prostatic diameter (20%), prostatic volume (43%), and serum DHT concentration (58%).</td> </tr> </table>	Animal Model:	Male dogs with spontaneous BPH (2.7-11 years old; 10.3-49 kg) ^[3]	Dosage:	0.1-0.5 mg/kg	Administration:	P.o. once daily for 16 weeks	Result:	Decreased prostatic diameter (20%), prostatic volume (43%), and serum DHT concentration (58%).
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Result:	Decreased prostatic diameter (20%), prostatic volume (43%), and serum DHT concentration (58%).								

Decreased semen volume but did not adversely effect on semen quality or serum testosterone concentration.
No adverse effects on dogs.

CUSTOMER VALIDATION

- J Pain. 2019 May;20(5):577-591.
- Sci Rep. 2019 Dec 23;9(1):19703.

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REFERENCES

- [1]. Flores E, et, al. Steroid 5alpha-reductase inhibitors. Mini Rev Med Chem. 2003 May;3(3):225-37.
- [2]. Yun DK, et, al. Finasteride Increases the Expression of Hemoxygenase-1 (HO-1) and NF-E2-Related Factor-2 (Nrf2) Proteins in PC-3 Cells: Implication of Finasteride-Mediated High-Grade Prostate Tumor Occurrence. Biomol Ther (Seoul). 2013 Jan;21(1):49-53.
- [3]. Sirinarumitr K, et, al. Effects of finasteride on size of the prostate gland and semen quality in dogs with benign prostatic hypertrophy. J Am Vet Med Assoc. 2001 Apr 15;218(8):1275-80.
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Caution: Product has not been fully validated for medical applications. For research use only.

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