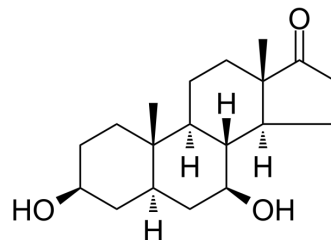


7β-Hydroxy-epi-androsterone

Cat. No.:	HY-107043
CAS No.:	25848-69-5
Molecular Formula:	C ₁₉ H ₃₀ O ₃
Molecular Weight:	306.44
Target:	Estrogen Receptor/ERR
Pathway:	Vitamin D Related/Nuclear Receptor
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	7β-Hydroxy-epi-androsterone (7β-Hydroxy-EpiA) can bind to ERβ and has anti-inflammatory and neuroprotective properties. 7β-Hydroxy-epi-androsterone is an endogenous androgenic derivative of dehydroepiandrosterone ^{[1][2][3]} .								
IC₅₀ & Target	ERβ								
In Vitro	<p>7β-Hydroxy-epi-androsterone (1, 10 and 100 nM, 72 h) inhibits MCF-7 and MDA-MB-231 cell proliferation in the presence of E2 (10 nM)^[2].</p> <p>7β-Hydroxy-epi-androsterone (1, 10 and 100 nM, 48 h) leads to a G0/G1 cell cycle arrest in MCF-7 and MDA-MB-231 cells^[2].</p> <p>7β-Hydroxy-epi-androsterone (1, 10 and 100 nM, 48 h) increases cell apoptosis in the presence of E2 (10 nM) in MCF-7 cells^[2].</p> <p>7β-Hydroxy-epi-androsterone (1, 10 and 100 nM, 24 h) binds to ERβ and decreases in transactivation in MDA-MB-231 cells^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Apoptosis Analysis^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MCF-7 cells</td> </tr> <tr> <td>Concentration:</td> <td>1, 10 and 100 nM (in presence of 10 nM E2)</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Increased cell apoptosis in presence of E2.</td> </tr> </table>	Cell Line:	MCF-7 cells	Concentration:	1, 10 and 100 nM (in presence of 10 nM E2)	Incubation Time:	48 h	Result:	Increased cell apoptosis in presence of E2.
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Concentration:	1, 10 and 100 nM (in presence of 10 nM E2)								
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Result:	Increased cell apoptosis in presence of E2.								
In Vivo	<p>7β-Hydroxy-epi-androsterone (i.p., once a day for 7 days) displays anti-inflammatory effects in rat with DSS-induced colitis^[3].</p> <p>7β-Hydroxy-epi-androsterone (0.1 mg/kg s.c., b.i.d. for 10 days) shows protective effect against inflammatory neurodegeneration and glial cell death in Alzheimer's disease rats^[4].</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>Rat with DSS-induced colitis^[3]</td> </tr> <tr> <td>Dosage:</td> <td>0.01, 0.1 and 1 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intraperitoneal injection (i.p.), once a day for 7 days.</td> </tr> </table>	Animal Model:	Rat with DSS-induced colitis ^[3]	Dosage:	0.01, 0.1 and 1 mg/kg	Administration:	Intraperitoneal injection (i.p.), once a day for 7 days.		
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Dosage:	0.01, 0.1 and 1 mg/kg								
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Result:

Increased in COX-2 and PGE synthase expression.
Increased colonic tissue levels of 15d-PGJ2 levels.

REFERENCES

- [1]. Christophe Ricco, et al. Synthesis of 7 β -hydroxy-epiandrosterone. Steroids. Volume 76, Issues 1-2, January 2011, Pages 28-30.
- [2]. Sandra N, et al. The DHEA metabolite 7 β -hydroxy-epiandrosterone exerts anti-estrogenic effects on breast cancer cell lines. Steroids. 2012 Apr;77(5):542-51.
- [3]. Hennebert O, et al. Anti-inflammatory effects and changes in prostaglandin patterns induced by 7beta-hydroxy-epiandrosterone in rats with colitis. J Steroid Biochem Mol Biol. 2008 Jun;110(3-5):255-62.
- [4]. Dudas B, et al. Protection against inflammatory neurodegeneration and glial cell death by 7beta-hydroxy epiandrosterone, a novel neurosteroid. Neurobiol Dis. 2004 Mar;15(2):262-8.
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Caution: Product has not been fully validated for medical applications. For research use only.

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