## SANTA CRUZ BIOTECHNOLOGY, INC.

# CYP11A1 (C-16): sc-18043



### BACKGROUND

CYP11A1, also known as cytochrome P450C11A1, cytochrome P450<sup>scc</sup> and cytochrome P450, subfamily XIA, is an enzyme that catalyzes the first step of steroid biosynthesis under the modulation of cAMP signal. CYP11A1 in steroidogenic cells converts cholesterol to pregnenolone, which is determined by hormonal control of cholesterol availability. Expression of the CYP11A1 gene is controlled by the transcription factor SF-1, and the upstream SF-1 binding site in the CYP11A1 gene is required for hormonal stimulation. c-Jun and SF-1 may act synergistically to activate CYP11A1 gene expression. Both Forskolin and 8-Br-cAMPS elevate CYP11A1 mRNA levels in the interstitial cell monolayer, which has a fully functional adenylate cyclase. The CYP11A1 protein is coexpressed with 3 $\beta$ -HSD2 in the rat hippocampus, dentate dyrus, cerebellar granular layer and Purkinje cells, indicating that neurosteriods are synthesized in a region-specific manner in the brain. CYP11A1 interacts with its physiological partner, adrenodoxin, by electrostatic interaction.

## CHROMOSOMAL LOCATION

Genetic locus: CYP11A1 (human) mapping to 15q24.1; Cyp11a1 (mouse) mapping to 9 B.

### SOURCE

CYP11A1 (C-16) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of CYP11A1 of human origin.

#### PRODUCT

Each vial contains 200  $\mu g$  IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-18043 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

#### **STORAGE**

Store at 4° C, \*\*D0 NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

#### **APPLICATIONS**

CYP11A1 (C-16) is recommended for detection of CYP11A1 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for CYP11A1 siRNA (h): sc-41496, CYP11A1 siRNA (m): sc-41497, CYP11A1 shRNA Plasmid (h): sc-41496-SH, CYP11A1 shRNA Plasmid (m): sc-41497-SH, CYP11A1 shRNA (h) Lentiviral Particles: sc-41496-V and CYP11A1 shRNA (m) Lentiviral Particles: sc-41497-V.

Molecular Weight of CYP11A1: 60 kDa.

#### **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

## SELECT PRODUCT CITATIONS

- Niakan, K.K., et al. 2006. Novel role for the orphan nuclear receptor DAX-1 in embryogenesis, different from steroidogenesis. Mol. Genet. Metab. 88: 261-271.
- Armenti, A.E., et al. 2008. Developmental methoxychlor exposure affects multiple reproductive parameters and ovarian folliculogenesis and gene expression in adult rats. Toxicol. Appl. Pharmacol. 233: 286-296.
- Xu, B., et al. 2009. DAX-1 and steroid receptor RNA activator (SRA) function as transcriptional coactivators for steroidogenic factor 1 in steroidogenesis. Mol. Cell. Biol. 29: 1719-1734.
- Huang, Y.H., et al. 2009. Pluripotency of mouse spermatogonial stem cells maintained by IGF-1- dependent pathway. FASEB J. 23: 2076-2087.
- Carmona, F.D., et al. 2009. SOX9 is not required for the cellular events of testicular organogenesis in XX mole ovotestes. J. Exp. Zool. B Mol. Dev. Evol. 312: 734-748.
- 7. Han, D.Y., et al. 2010. Polychlorinated biphenyls have inhibitory effect on testicular steroidogenesis by downregulation of P450<sup>17 $\alpha$ </sup> and P450<sup>scc</sup>. Toxicol. Ind. Health 26: 287-296.
- Sherrill, J.D., et al. 2010. Developmental exposures of male rats to soy isoflavones impact Leydig cell differentiation. Biol. Reprod. 83: 488-501.
- Tinfo, N.S., et al. 2011. Understanding the effects of atrazine on steroidogenesis in rat granulosa and H295R adrenal cortical carcinoma cells. Reprod. Toxicol. 31: 184-193.
- Nanjappa, M.K., et al. 2012. The industrial chemical bisphenol A (BPA) interferes with proliferative activity and development of steroidogenic capacity in rat Leydig cells. Biol. Reprod. 86: 135, 1-12.
- Stojkov, N.J., et al. 2012. Repeated immobilization stress disturbed steroidogenic machinery and stimulated the expression of cAMP signaling elements and adrenergic receptors in Leydig cells. Am. J. Physiol. Endocrinol. Metab. 302: E1239-E1251.
- Ren, X.M., et al. 2012. The protection of selenium on cadmium-induced inhibition of spermatogenesis via activating testosterone synthesis in mice. Food Chem. Toxicol. 50: 3521-3529.
- Zhao, Y., et al. 2012. Perfluorooctanoic acid effects on ovaries mediate its inhibition of peripubertal mammary gland development in Balb/c and C57BI/6 mice. Reprod. Toxicol. 33: 563-576.