# SANTA CRUZ BIOTECHNOLOGY, INC.

# TRβ1 (P-16): sc-10823



# BACKGROUND

Thyroid hormone nuclear receptors (TRs) are ligand-dependent transcription factors which regulate growth, differentiation and development and represent members of the steroid/retinoic acid superfamily. The two genes encoding TRs identified to date, TR $\alpha$  and TR $\beta$ , have been mapped to human chromosomes 17 and 3, respectively. TRs bind to thyroid hormone response elements (TREs) with half-site binding motifs in the orientation of palindromes, direct repeats or inverted palindromes. The affinities of binding are both variable and influenced differentially by 3,5,3'-triiodo-L-thyronine (T3). Transcriptional regulation by TRs is also modulated by heterodimerization with TR nuclear accessory proteins, the most extensively characterized of which are the retinoid X receptors (RXR $\alpha$ , RXR $\beta$  and RXR $\gamma$ ). The TR $\beta$  isoform TR $\beta$ 1 forms a complex with the Pl 3-kinase p85 $\alpha$  subunit and plays an important role in the T3-induced activation of Akt in pancreatic  $\beta$  cells.

#### REFERENCES

- Näär, A., et al. 1991. The orientation and spacing of core DNA-binding motifs dictate selective transcriptional responses to three nuclear receptors. Cell 65: 1267-1271.
- Lazar, M.A. 1993. Thyroid hormone receptors: multiple forms, multiple possibilities. Endocrinol. Rev. 14: 184-193.
- Meier, C.A., et al. 1993. Interaction of human TRβ1 and its mutants with DNA and RXRβ. T3 response element-dependent dominant negative potency. J. Clin. Invest. 92: 1986-1993.
- 4. Zhang, X.K., et al. 1993. Hetero- and homodimeric receptors in thyroid hormone and vitamin A action. Receptor 3: 183-191.
- Bhat, M.K., et al. 1994. Phosphorylation enhances the target gene sequence-dependent dimerization of thyroid hormone receptor with retinoid X receptor. Proc. Natl. Acad. Sci. USA 91: 7927-7931.
- Sugawara, A., et al. 1994. Phosphorylation selectively increases triiodothyronine receptor homodimer binding to DNA. J. Biol. Chem. 269: 433-437.
- 7. Verga Falzacappa, C., et al. 2007. Thyroid hormone receptor TR $\beta$ 1 mediates Akt activation by T3 in pancreatic  $\beta$  cells. J. Mol. Endocrinol. 38: 221-233.

## CHROMOSOMAL LOCATION

Genetic locus: THRB (human) mapping to 3p24.2; Thrb (mouse) mapping to 14 A2.

#### SOURCE

TR $\beta$ 1 (P-16) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of TR $\beta$ 1 of human origin.

#### STORAGE

Store at 4° C, \*\*DO 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.

## PRODUCT

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

Blocking peptide available for competition studies, sc-10823 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

Available as TransCruz reagent for Gel Supershift and ChIP applications, sc-10823 X, 200  $\mu$ g/0.1 ml.

# **APPLICATIONS**

TRβ1 (P-16) is recommended for detection of TRβ1 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), 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).

 $TR\beta1$  (P-16) is also recommended for detection of  $TR\beta1$  in additional species, including equine and canine.

Suitable for use as control antibody for TR $\beta$ 1 siRNA (h): sc-38890, TR $\beta$ 1 siRNA (m): sc-38891, TR $\beta$ 1 shRNA Plasmid (h): sc-38890-SH, TR $\beta$ 1 shRNA Plasmid (m): sc-38891-SH, TR $\beta$ 1 shRNA (h) Lentiviral Particles: sc-38890-V and TR $\beta$ 1 shRNA (m) Lentiviral Particles: sc-38891-V.

 $TR\beta1$  (P-16) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of TR $\beta$ 1: 55 kDa.

Positive Controls: C32 whole cell lysate: sc-2205.

## **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) Immunofluo-rescence: 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

 Fava, G., et al. 2006. Thyroid hormone inhibits biliary growth in bile ductligated rats by PLC/IP<sub>3</sub>/Ca<sup>2+</sup>-dependent downregulation of SRC/ERK1/2. Am. J. Physiol. Cell Physiol. 292: C1467-C1475.



Try **TRβ1 (J51): sc-737** or **TRβ1 (J52): sc-738**, our highly recommended monoclonal alternatives to TRβ1 (P-16). Also, for AC, HRP, FITC, PE, Alexa Fluor<sup>®</sup> 488 and Alexa Fluor<sup>®</sup> 647 conjugates, see **TRβ1 (J51): sc-737**.