## SANTA CRUZ BIOTECHNOLOGY, INC.

# TRα1/β1 (C3): sc-32754



## BACKGROUND

Thyroid hormone nuclear receptors (TRs) are ligand-dependent transcription factors which regulate and control many metabolic and developmental processes. There are two genes encoding TRs identified to date, TR $\alpha$  and TR $\beta$ . 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 $\alpha$  isoform, TR $\alpha$ 1, can display both a nuclear and undefined cytoplasmic location, and is the only TR that is imported into the mitochondrial matrix. The TR $\beta$  isoform TR $\beta$ 1 forms a complex with the PI 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. Endocr. 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.

## CHROMOSOMAL LOCATION

Genetic locus: THRA (human) mapping to 17q21.1, THRB (human) mapping to 3p24.2.

## SOURCE

TR $\alpha$ 1/ $\beta$ 1 (C3) is a mouse monoclonal antibody raised against the ligand binding domain of the thyroid hormone receptor  $\alpha$ 1 of human origin.

## PRODUCT

Each vial contains 200  $\mu$ g lgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin. Also available as TransCruz reagent for Gel Supershift and ChIP applications, sc-32754 X, 200  $\mu$ g/0.1 ml.

## STORAGE

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

## APPLICATIONS

TR $\alpha$ 1/ $\beta$ 1 (C3) is recommended for detection of TR $\alpha$ 1 and TR $\beta$ 1 of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)].

 $TR\alpha 1/\beta 1$  (C3) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of TR $\alpha$ 1: 47 kDa.

Molecular Weight of TR<sub>β1</sub>: 58 kDa.

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

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml).

#### DATA



 $TR\alpha1/\beta1$  (C3): sc-32754. Western blot analysis of  $\mathit{in}$   $\mathit{vito}$  synthesized human TRβ1 (A-C) and TR\alpha1 (D-F) prepared from TNT T7 programmed lysates. 1  $\mu$ I (A,D), 3  $\mu$ I (B,E) and 6  $\mu$ I (C,F).

## SELECT PRODUCT CITATIONS

- Huang, W., et al. 2010. qHTS for inhibitors of the interaction of thyroid hormone receptor and steroid receptor coregulator 2. Molecular Libraries.
- Hwang, J.Y., et al. 2011. Methylsulfonylnitrobenzoates, a new class of irreversible inhibitors of the interaction of the thyroid hormone receptor and its obligate coactivators that functionally antagonizes thyroid hormone. J. Biol. Chem. 286: 11895-11908.

#### **RESEARCH USE**

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



See **TRβ1 (J51): sc-737** for TRβ1 antibody conjugates, including AC, HRP, FITC, PE, and Alexa Fluor<sup>®</sup> 488, 546, 594, 647, 680 and 790.