# SANTA CRUZ BIOTECHNOLOGY, INC.

# Tra-2β (D-4): sc-166769



## BACKGROUND

Human transformer-2  $\alpha$  (Tra-2 $\alpha$ ) and Tra-2 $\beta$  are nuclear proteins that associate with distinct pre-mRNA splicing enhancer elements. Tra-2 $\alpha$  is the functional homolog of the *Drosophila* TRA-2 protein, which regulates the female specific pre-mRNA splicing pattern of the doublesex (dsx) gene in *Drosophila*. Human Tra-2 proteins are able to actively splice *Drosophila* sex factors; however, human Tra-2 $\alpha$  has not been shown to induce sexual differentiation, which suggests that human Tra-2 $\alpha$  proteins may regulate splicing patterns involving alternative mechanisms. Tra-2 $\alpha$  and Tra-2 $\beta$  proteins contain a single RNP-type RNA-binding domain and selectively bind to purine-rich sequences to facilitate mRNA splicing. Expression of Tra-2 $\beta$  is upregulated during the reoxygenation of hypoxic astrocytes, and both Tra-2 $\alpha$  and Tra-2 $\beta$  interact with the serine/arginine-rich (SR) family of splicing factors to form Tra-2/SR complexes that then regulate tissue-specific alternative splicing patterns of many pre-mRNAs.

## REFERENCES

- 1. Amrein, H., et al. 1994. The role of specific protein-RNA and proteinprotein interactions in positive and negative control of pre-mRNA splicing by transformer 2. Cell 76: 735-746.
- 2. Matsuo, N., et al. 1995. Cloning of a novel RNA binding polypeptide (RA301) induced by hypoxia/reoxygenation. J. Biol. Chem. 270: 28216-28222.
- Dauwalder, B., et al. 1996. A human homologue of the *Drosophila* sex determination factor transformer-2 has conserved splicing regulatory functions. Proc. Natl. Acad. Sci. USA 93: 9004-9009.
- Segade, F., et al. 1996. Molecular cloning of a mouse homologue for the Drosophila splicing regulator Tra-2. FEBS Lett. 387: 152-156.

### **CHROMOSOMAL LOCATION**

Genetic locus: TRA2B (human) mapping to 3q27.2; Tra2b (mouse) mapping to 16 B1.

#### SOURCE

Tra-2 $\beta$  (D-4) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 10-50 near the N-terminus of Tra-2 $\beta$  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-166769 X, 200  $\mu$ g/0.1 ml.

Blocking peptide available for competition studies, sc-166769 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

#### **STORAGE**

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

#### **APPLICATIONS**

Tra-2 $\beta$  (D-4) is recommended for detection of Tra-2 $\beta$  of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg 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 Tra-2 $\beta$  siRNA (h): sc-38566, Tra-2 $\beta$  siRNA (m): sc-38567, Tra-2 $\beta$  shRNA Plasmid (h): sc-38566-SH, Tra-2 $\beta$  shRNA Plasmid (m): sc-38567-SH, Tra-2 $\beta$  shRNA (h) Lentiviral Particles: sc-38566-V and Tra-2 $\beta$  shRNA (m) Lentiviral Particles: sc-38567-V.

 $Tra-2\beta$  (D-4) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight (predicted) of Tra-2 β isoforms: 33/22/4 kDa.

Molecular Weight (observed) of Tra-2ß isoforms: 40 kDa.

Positive Controls: IMR-32 nuclear extract: sc-2148, HeLa nuclear extract: sc-2120 or SK-N-MC nuclear extract: sc-2154.

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> 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). 3) Immunofluorescence: use m-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz<sup>®</sup> Mounting Medium: sc-24941 or UltraCruz<sup>®</sup> Hard-set Mounting Medium: sc-359850.

#### DATA





Tra-2  $\beta$  (D-4): sc-166769. Western blot analysis of Tra-2  $\beta$  expression in IMR-32 nuclear extract.

Tra-2  $\beta$  (D-4): sc-166769. Western blot analysis of Tra-2  $\beta$  expression in IMR-32 nuclear extract.

#### **RESEARCH USE**

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

## PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.