SANTA CRUZ BIOTECHNOLOGY, INC.

TGFβ2 (H-6): sc-374659



BACKGROUND

Transforming growth factor β s (TGF β s) were originally discovered due to their ability to promote anchorage-independent growth of rat NRK fibroblasts in the presence of TGF α . It is now realized that TGF β s mediate many cell-cell interactions that occur during embryonic development. Three TGF β s have been identified in mammals. TGF β 1, TGF β 2 and TGF β 3 are each synthesized as precursor proteins that are very similar in that each is cleaved to yield a 112 amino acid polypeptide that remains associated with the latent portion of the molecules. Biologically active TGF β requires dimerization of the monomers (usually homodimers) and release of the latent peptide portion. Overall, the mature region of the TGF β 1 and TGF β 2. However, the NH₂ terminals or precursor regions of their molecules share only 27% sequence identity.

REFERENCES

- Todaro, G.J., et al. 1980. Transforming growth factors produced by certain human tumor cells: polypeptides that interact with epidermal growth factor receptors. Proc. Natl. Acad. Sci. USA 77: 5258-5262.
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- Derynck, R., et al. 1985. Human transforming growth factor-β cDNA sequence and expression in tumor cell lines. Nature 316: 701-705.
- deMartin, R., et al. 1987. Complementary DNA for human glioblastomaderived factor-β family. EMBO J. 6: 3673-3677.
- 5. ten Dijke, P., et al. 1988. Identification of a new member of the transforming growth factor type β gene family. Proc. Natl. Acad. Sci. USA 85: 4715-4719.

CHROMOSOMAL LOCATION

Genetic locus: TGFB2 (human) mapping to 1q41; Tgfb2 (mouse) mapping to 1 H5.

SOURCE

TGF β 2 (H-6) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 343-377 at the C-terminus of TGF β 2 of human origin.

PRODUCT

Each vial contains 200 μg lgG_3 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-374659 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein).

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.

APPLICATIONS

TGF β 2 (H-6) is recommended for detection of mature and precursor forms of TGF β 2 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).

TGF β 2 (H-6) is also recommended for detection of mature and precursor forms of TGF β 2 in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for TGF β 2 siRNA (h): sc-39802, TGF β 2 siRNA (m): sc-39803, TGF β 2 shRNA Plasmid (h): sc-39802-SH, TGF β 2 shRNA Plasmid (m): sc-39803-SH, TGF β 2 shRNA (h) Lentiviral Particles: sc-39802-V and TGF β 2 shRNA (m) Lentiviral Particles: sc-39803-V.

Molecular Weight of TGF_β2 monomer: 13 kDa.

Molecular Weight of TGF_β2 dimer: 25 kDa.

Positive Controls: human placenta extract: sc-363772.

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[®] 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-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850.

DATA



TGF β 2 (H-6): sc-374659. Western blot analysis of human recombinant TGF β 2 fusion protein.

SELECT PRODUCT CITATIONS

 Zepeda-Morales, A.S., et al. 2016. Liver fibrosis in bile duct-ligated rats correlates with increased hepatic IL-17 and TGFβ2 expression. Ann. Hepatol. 15: 418-426.

PROTOCOLS

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