### SANTA CRUZ BIOTECHNOLOGY, INC.

# Smad1/5 (T-20): sc-6201



BACKGROUND

Smad proteins, the mammalian homologs of the *Drosophila* mothers against dpp (Mad) have been implicated as downstream effectors of TGF $\beta$ /BMP signaling. Smad1 (also designated Madr1 or JV4-1), Smad5 and mammalian Smad8 (also designated Smad9 or MadH6) are effectors of BMP2 and BMP4 function while Smad2 (also designated Madr2 or JV18-1) and Smad3 are involved in TGF $\beta$  and activin-mediated growth modulation. Smad4 (also designated DPC4) has been shown to mediate all of the above activities through interaction with various Smad family members. Smad6 and Smad7 regulate the response to activin/TGF $\beta$  signaling by interfering with TGF $\beta$ -mediated phosphorylation of other Smad family members.

#### SOURCE

Smad1/5 (T-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping within the amino terminal domain of Smad1 of human origin.

#### 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-6201 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-6201 X, 200  $\mu g/0.1$  ml.

#### **APPLICATIONS**

Smad1/5 (T-20) is recommended for detection of Smad1, Smad5 and, to a lesser extent, Smad8 and Smad9 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).

Smad1/5 (T-20) is also recommended for detection of Smad1, Smad5 and, to a lesser extent, Smad8 and Smad9 in additional species, including equine, bovine, porcine and avian.

Smad1 (T-20) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of Smad1: 52-56 kDa.

Molecular Weight of Smad5: 43 kDa.

Positive Controls: NIH/3T3 whole cell lysate: sc-2210.

#### **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.

#### SELECT PRODUCT CITATIONS

- Wang, W., et al. 2000. Ski represses bone morphogenic protein signaling in *Xenopus* and mammalian cells. Proc. Natl. Acad. Sci. USA 97: 14394-14399.
- Bau, B., et al. 2002. Bone morphogenetic protein-mediating receptor-associated Smads as well as common Smad are expressed in human articular chondrocytes but not upregulated or downregulated in osteoarthritic cartilage. J. Bone Miner. Res. 17: 2141-2150.
- Oxburgh, L., et al. 2002. Dynamic regulation of Smad expression during mesenchyme to epithelium transition in the metanephric kidney. Mech. Dev. 112: 207-211.
- 4. Liu, X., et al. 2003. Smad7 but not Smad6 cooperates with oncogenic Ras to cause malignant conversion in a mouse model for squamous cell carcinoma. Brain Res. 63: 7760-7768.
- Raju, G.P., et al. 2003. SANE, a novel LEM domain protein, regulates bone morphogenetic protein signaling through interaction with Smad1. J. Biol. Chem. 278: 428-437.
- Munoz, O., et al. 2004. TGFβ-mediated activation of Smad1 in B-cell non-Hodgkin's lymphoma and effect on cell proliferation. Leukemia 18: 2015-2025.
- 7. Maire, M., et al. 2005. Alteration of transforming growth factor- $\beta$  signaling system expression in adult rat germ cells with a chronic apoptotic cell death process after fetal androgen disruption. Endocrinology 146: 5135-5143.
- Ji, M., et al. 2005. High-level activation of cyclic AMP signaling attenuates bone morphogenetic protein 2-induced sympathoadrenal lineage development and promotes melanogenesis in neural crest cultures. Mol. Cell. Biol. 25: 5134-5145.
- Fukuda, T., et al. 2009. Constitutively activated ALK2 and increased Smad1/5 cooperatively induce bone morphogenetic protein signaling in fibrodysplasia ossificans progressiva. J. Biol. Chem. 114: 7149-7156.
- Yamazaki, M., et al. 2009. Tumor necrosis factor α represses bone morphogenetic protein (BMP) signaling by interfering with the DNA binding of Smads through the activation of NFκB. J. Biol. Chem. 284: 35987-35995.
- 11. Gramley, F., et al. 2010. Atrial fibrosis and atrial fibrillation: the role of the TGF- $\beta$ 1 signaling pathway. Int. J. Cardiol. 143: 405-413.
- Alfaro, M.P., et al. 2010. sFRP2 suppression of bone morphogenic protein (BMP) and Wnt signaling mediates mesenchymal stem cell (MSC) selfrenewal promoting engraftment and myocardial repair. J. Biol. Chem. 285: 35645-35653.

## MONOS Satisfation Guaranteed

Try **Smad1 (A-4): sc-7965**, our highly recommended monoclonal aternatives to Smad1/5 (T-20). Also, for AC, HRP, FITC, PE, Alexa Fluor<sup>®</sup> 488 and Alexa Fluor<sup>®</sup> 647 conjugates, see **Smad1 (A-4): sc-7965**.