TNAP (D-3): sc-137213



The Power to Overtio

BACKGROUND

Alkaline phosphatases (AP) are glycosyl-phosphatidylinositol (GPI)-anchored, dimeric, Zn²⁺ metallated glycoproteins that catalyze the hydrolysis of phosphomonoesters into an inorganic phosphate and an alcohol. There are at least four distinct but related alkaline phosphatases: intestinal (IAP), placental (PLAP), placental-like (ALP-1 or GCAP) and tissue non-specific (TNAP). The first three are located together on chromosome 2 while the tissue non-specific form is located on chromosome 1. TNAP is widely expressed in liver, kidney, bone, stomach and colon, and is therefore referred to as the tissue non-specific form of AP. TNAP, in conjuntion with plasma cell membrane glycoprotein-1, function in bone mineralization; however, mice that lack a functional form of TNAP show normal skeletal development. This enzyme has been linked directly to a disorder known as hypophosphatasia, a rare inborn disorder that is characterized by defective bone mineraliation and includes skeletal defects. Human gene encoding TNAP maps to chromosome 1p36.12.

CHROMOSOMAL LOCATION

Genetic locus: ALPL (human) mapping to 1p36.12; Alpl (mouse) mapping to 4 D3.

SOURCE

TNAP (D-3) is a mouse monoclonal antibody raised against amino acids 18-317 mapping near the N-terminus of TNAP of human origin.

PRODUCT

Each vial contains 200 $\mu g \ lgG_1$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

TNAP (D-3) is recommended for detection of TNAP 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 TNAP siRNA (h): sc-38921, TNAP siRNA (m): sc-38922, TNAP shRNA Plasmid (h): sc-38921-SH, TNAP shRNA Plasmid (m): sc-38922-SH, TNAP shRNA (h) Lentiviral Particles: sc-38921-V and TNAP shRNA (m) Lentiviral Particles: sc-38922-V.

Molecular Weight of TNAP: 80 kDa.

Molecular Weight of PLAP: 70 kDa.

Molecular Weight of GCAP: 23 kDa.

Molecular Weight of IAP: 57 kDa.

Positive Controls: Hep G2 cell lysate: sc-2227, Jurkat whole cell lysate: sc-2204 or HeLa whole cell lysate: sc-2200.

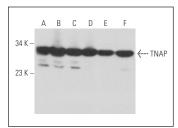
STORAGE

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

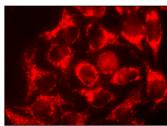
RESEARCH USE

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

DATA



TNAP (D-3): sc-137213. Western blot analysis of TNAP expression in Hep G2 ($\bf A$), HeLa ($\bf B$), A-431 ($\bf C$), MCF7 ($\bf D$), Saos-2 ($\bf E$) and Jurkat ($\bf F$) whole cell lysates.



TNAP (D-3): sc-137213. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization.

SELECT PRODUCT CITATIONS

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- Mentrup, B., et al. 2011. Functional characterization of a novel mutation localized in the start codon of the tissue-nonspecific alkaline phosphatase gene. Bone 48: 1401-1408.
- 3. Yosef, N. and Ubogu, E.E. 2013. An immortalized human blood-nerve barrier endothelial cell line for *in vitro* permeability studies. Cell. Mol. Neurobiol. 33: 175-186.
- 4. Wong, M.Y., et al. 2014. 11 kGy γ irradiated demineralized bone matrix enhances osteoclast activity. Eur. J. Orthop. Surg. Traumatol. 24: 655-661.
- Lin, T., et al. 2016. Dorsomorphin homologue 1, a highly selective smallmolecule bone morphogenetic protein inhibitor, suppresses medial artery calcification. J. Vasc. Surg. 66: 586-593.
- Hulshof, F.F.B., et al. 2017. Mining for osteogenic surface topographies: in silico design to *in vivo* osseo-integration. Biomaterials 137: 49-60.
- 7. Maximiano, W.M.A., et al. 2017. Mast cell mediators inhibit osteoblastic differentiation and extracellular matrix mineralization. J. Histochem. Cytochem. 65: 723-741.
- 8. Zhao, L., et al. 2018. Function of GCN5 in the TGF-β1-induced epithelial-to-mesenchymal transition in breast cancer. Oncol. Lett. 16: 3955-3963.
- 9. Wilson, K.M., et al. 2018. Glycans modify mesenchymal stem cell differentiation to impact on the function of resulting osteoblasts. J. Cell Sci. 131: jcs209452.
- Hoshikawa, S., et al. 2020. Phosphorylation-dependent osterix degradation negatively regulates osteoblast differentiation. FASEB J. 34: 14930-14945.



See **PLAP (8B6):** sc-47691 for PLAP antibody conjugates, including AC, HRP, FITC, PE, and Alexa Fluor[®] 488, 546, 594, 647, 680 and 790.