SANTA CRUZ BIOTECHNOLOGY, INC.

FGFR-3 (C-15): sc-123



BACKGROUND

Acidic and basic fibroblast growth factors (FGFs) are members of a family of multifunctional polypeptide growth factors that stimulate proliferation of cells of mesenchymal, epithelial and neuroectodermal origin. Like other growth factors, FGFs act by binding and activating specific cell surface receptors. These include the Flg receptor or FGFR-1, the Bek receptor or FGFR-2, FGFR-3, FGFR-4, FGFR-5 and FGFR-6. These receptors usually contain an extracellular ligand-binding region containing three immunoglobulin-like domains, a transmembrane domain and a cytoplasmic tyrosine kinase domain. The gene encoding human FGFR-3 maps to chromosome 4p16.3 and is alternatively spliced to produce three isoforms that are expressed in brain, kidney and testis. Defects in FGFR-3 are associated with several diseases, including Crouzon syndrome, achondroplasia, thanatophoric dysplasia, craniosynostosis adelaide type and hypochondroplasia. Mutations in FGFR-3 are also a cause of some bladder and cervical cancers.

CHROMOSOMAL LOCATION

Genetic locus: FGFR3 (human) mapping to 4p16.3; Fgfr3 (mouse) mapping to 5 B2.

SOURCE

FGFR-3 (C-15) is an affinity purified rabbit polyclonal antibody raised against a peptide mapping at the C-terminus of FGFR-3 of human origin.

PRODUCT

Each vial contains 100 μg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-123 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

Available as agarose conjugate for immunoprecipitation, sc-123 AC, 500 μ g/0.25 ml agarose in 1 ml.

APPLICATIONS

FGFR-3 (C-15) is recommended for detection of FGFR-3 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (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 FGFR-3 siRNA (h): sc-29314, FGFR-3 siRNA (m): sc-35367, FGFR-3 shRNA Plasmid (h): sc-29314-SH, FGFR-3 shRNA Plasmid (m): sc-35367-SH, FGFR-3 shRNA (h) Lentiviral Particles: sc-29314-V and FGFR-3 shRNA (m) Lentiviral Particles: sc-35367-V.

Molecular Weight of non-glycosylated FGFR-3: 97 kDa.

Molecular Weight of FGFR-3 precursor: 125 kDa.

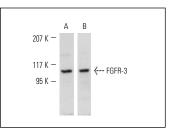
Molecular Weight of mature FGFR-3: 135 kDa.

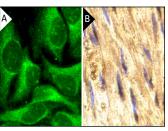
Positive Controls: K-562 whole cell lysate: sc-2203, T-47D cell lysate: sc-2293 or A549 cell lysate: sc-2413.

STORAGE

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

DATA





FGFR-3 (C-15): sc-123. Western blot analysis of FGFR-3 expression in A549 $({\rm A})$ and T-47D $({\rm B})$ whole cell lysates.

FGFR-3 (C-15): sc-123. Immunofluorescence staining of methanol-fixed HeLa cells showing cyroplasmic localization (**A**). Immunoperoxidase staining of formalin fixed, parafin-embedded human colon tissue showing cytoplasmic staining of smooth muscle cells (**B**).

SELECT PRODUCT CITATIONS

- 1. Webster, M.K., et al. 1996. Constitutive activation of fibroblast growth factor receptor 3 by the transmembrane domain point mutation found in achondroplasia. EMBO J. 15: 520-527.
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- 3. You, H., et al. 2011. Construction of eukaryotic expression plasmid of hTGF- β 3 and its inducing effect on differentiation of precartilaginous stem cells into chondroblasts. J. Huazhong Univ. Sci. Technol. Med. Sci. 31: 524-529.
- Ikner, A., et al. 2011. TWEAK induces apoptosis through a death-signaling complex comprising receptor-interacting protein 1 (RIP1), Fas-associated death domain (FADD), and caspase-8. J. Biol. Chem. 286: 21546-21554.
- Calvo, F., et al. 2011. RasGRF suppresses Cdc42-mediated tumour cell movement, cytoskeletal dynamics and transformation. Nat. Cell Biol. 13: 819-826.
- 6. Zemskov, E.A., et al. 2012. Tissue transglutaminase promotes PDGF/ PDGFR-mediated signaling and responses in vascular smooth muscle cells. J. Cell. Physiol. 227: 2089-2096.
- Lim, K., et al. 2012. Vascular Klotho deficiency potentiates the development of human artery calcification and mediates resistance to fibroblast growth factor 23. Circulation 125: 2243-2255.
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RESEARCH USE

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