

PDGFR- α (C-9): sc-398206

BACKGROUND

Platelet-derived growth factor (PDGF) is a mitogen for mesenchyme- and gliaderived cells. PDGF consists of two chains, A and B, which dimerize to form functionally distinct isoforms PDGF-AA, PDGF-AB and PDGF-BB. These three isoforms bind with different affinities to two receptor types, PDGFR- α and - β , which are endowed with protein tyrosine kinase domains. PDGFR- α can bind to both A and B subunits of PDGF, while PDGFR- β can only bind the B subunit. Ligand binding promotes either homo- or heterodimerization of the PDGF receptors in a specific manner. PDGF-AA induces the dimerization of two α receptors, PDGF-AB induces dimerization of $\alpha\alpha$ and $\alpha\beta$ and PDGF-BB induces the formation of three types of dimers, $\alpha\alpha$, $\alpha\beta$ and $\beta\beta$. The genes encoding PDGFR- α and - β map to human chromosome 4q12 and 5q33.1, respectively. Translocation of the PDGFR- β gene with the TEL gene is linked with chronic myelomonocytic leukemia (CMML), a myelodysplastic syndrome, and demonstrates the oncogenic potential of the PDGF receptors.

REFERENCES

- Ross, R., et al. 1986. The biology of platelet-derived growth factor. *Cell* 46: 155-169.
- Hart, C.E., et al. 1988. Two classes of PDGF receptor recognize different isoforms of PDGF. *Science* 240: 1529-1531.
- Heldin, C., et al. 1988. Binding of different dimeric forms of PDGF to human fibroblasts: evidence for two separate receptor types. *EMBO J.* 7: 1387-1393.

CHROMOSOMAL LOCATION

Genetic locus: PDGFRA (human) mapping to 4q12; Pdgfra (mouse) mapping to 5 C3.3.

SOURCE

PDGFR- α (C-9) is a mouse monoclonal antibody raised against amino acids 951-1089 of PDGFR- α of human origin.

PRODUCT

Each vial contains 200 μ g IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

PDGFR- α (C-9) is available conjugated to agarose (sc-398206 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-398206 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-398206 PE), fluorescein (sc-398206 FITC), Alexa Fluor[®] 488 (sc-398206 AF488), Alexa Fluor[®] 546 (sc-398206 AF546), Alexa Fluor[®] 594 (sc-398206 AF594) or Alexa Fluor[®] 647 (sc-398206 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-398206 AF680) or Alexa Fluor[®] 790 (sc-398206 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

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STORAGE

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

APPLICATIONS

PDGFR- α (C-9) is recommended for detection of PDGFR- α of mouse, rat and human origin by Western Blotting (starting dilution 1:100, 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).

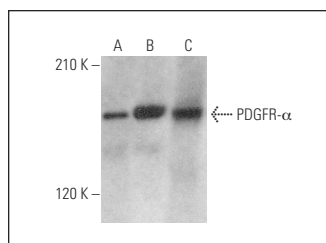
PDGFR- α (C-9) is also recommended for detection of PDGFR- α in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for PDGFR- α siRNA (h): sc-29443, PDGFR- α siRNA (m): sc-29444, PDGFR- α shRNA Plasmid (h): sc-29443-SH, PDGFR- α shRNA Plasmid (m): sc-29444-SH, PDGFR- α shRNA (h) Lentiviral Particles: sc-29443-V and PDGFR- α shRNA (m) Lentiviral Particles: sc-29444-V.

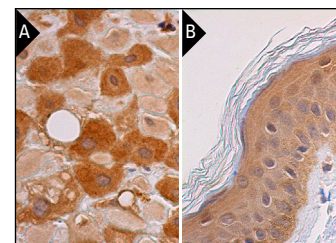
Molecular Weight of PDGFR- α : 170 kDa.

Positive Controls: U-2 OS cell lysate: sc-2295, HEK293 whole cell lysate: sc-45136 or NIH/3T3 whole cell lysate: sc-2210.

DATA



PDGFR- α (C-9): sc-398206. Western blot analysis of PDGFR- α expression in U-2 OS (A), HEK293 (B) and NIH/3T3 (C) whole cell lysates.



PDGFR- α (C-9): sc-398206. Immunoperoxidase staining of formalin fixed, paraffin-embedded human placenta tissue showing cytoplasmic staining of decidual cells (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human skin tissue showing cytoplasmic staining of keratinocytes, fibroblasts and melanocytes (B).

SELECT PRODUCT CITATIONS

- Rusu, M.C., et al. 2016. Subsets of telocytes: myocardial telocytes. *Ann. Anat.* 209: 37-44.
- Akama, T. and Chun, T.H. 2018. Transcription factor 21 (TCF21) promotes proinflammatory interleukin 6 expression and extracellular matrix remodeling in visceral adipose stem cells. *J. Biol. Chem.* 293: 6603-6610.
- Das, L., et al. 2019. Idiopathic gigantomastia: newer mechanistic insights implicating the paracrine milieu. *Endocrine* 66: 166-177.
- Santini, M.P., et al. 2020. Tissue-resident PDGFR α ⁺ progenitor cells contribute to fibrosis versus healing in a context- and spatiotemporally dependent manner. *Cell Rep.* 30: 555-570.

RESEARCH USE

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