

# PDGFRA Antibody (C-term E1063)

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP7666B

## Specification

## PDGFRA Antibody (C-term E1063) - Product Information

Application Primary Accession Other Accession Reactivity Predicted Host Clonality Isotype Calculated MW Antigen Region FC, IHC-P, WB,E <u>P16234</u> <u>P20786</u>, <u>P26618</u>, <u>O9PUF6</u> Human, Mouse Chicken, Rat Rabbit Polyclonal Rabbit IgG 122670 1048-1077

## PDGFRA Antibody (C-term E1063) - Additional Information

## Gene ID 5156

### **Other Names**

Platelet-derived growth factor receptor alpha, PDGF-R-alpha, PDGFR-alpha, Alpha platelet-derived growth factor receptor, Alpha-type platelet-derived growth factor receptor, CD140 antigen-like family member A, CD140a antigen, Platelet-derived growth factor alpha receptor, Platelet-derived growth factor receptor 2, PDGFR-2, CD140a, PDGFRA, PDGFR2, RHEPDGFRA

## Target/Specificity

This PDGFRA antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 1048-1077 amino acids from the C-terminal region of human PDGFRA.

Dilution FC~~1:10~50 IHC-P~~1:50~100 WB~~1:1000 E~~Use at an assay dependent concentration.

#### Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.

#### Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

#### Precautions

PDGFRA Antibody (C-term E1063) is for research use only and not for use in diagnostic or therapeutic procedures.



## PDGFRA Antibody (C-term E1063) - Protein Information

Name PDGFRA

Synonyms PDGFR2, RHEPDGFRA

Function Tyrosine-protein kinase that acts as a cell-surface receptor for PDGFA, PDGFB and PDGFC and plays an essential role in the regulation of embryonic development, cell proliferation, survival and chemotaxis. Depending on the context, promotes or inhibits cell proliferation and cell migration. Plays an important role in the differentiation of bone marrow-derived mesenchymal stem cells. Required for normal skeleton development and cephalic closure during embryonic development. Required for normal development of the mucosa lining the gastrointestinal tract, and for recruitment of mesenchymal cells and normal development of intestinal villi. Plays a role in cell migration and chemotaxis in wound healing. Plays a role in platelet activation, secretion of agonists from platelet granules, and in thrombin-induced platelet aggregation. Binding of its cognate ligands - homodimeric PDGFA, homodimeric PDGFB, heterodimers formed by PDGFA and PDGFB or homodimeric PDGFC -leads to the activation of several signaling cascades; the response depends on the nature of the bound ligand and is modulated by the formation of heterodimers between PDGFRA and PDGFRB. Phosphorylates PIK3R1, PLCG1, and PTPN11. Activation of PLCG1 leads to the production of the cellular signaling molecules diacylglycerol and inositol 1,4,5-trisphosphate, mobilization of cytosolic Ca(2+) and the activation of protein kinase C. Phosphorylates PIK3R1, the regulatory subunit of phosphatidylinositol 3-kinase, and thereby mediates activation of the AKT1 signaling pathway. Mediates activation of HRAS and of the MAP kinases MAPK1/ERK2 and/or MAPK3/ERK1. Promotes activation of STAT family members STAT1, STAT3 and STAT5A and/or STAT5B. Receptor signaling is down-regulated by protein phosphatases that dephosphorylate the receptor and its down-stream effectors, and by rapid internalization of the activated receptor.

#### **Cellular Location**

Cell membrane; Single-pass type I membrane protein. Cell projection, cilium {ECO:0000250|UniProtKB:P26618}. Golgi apparatus {ECO:0000250|UniProtKB:P26618}

#### Tissue Location

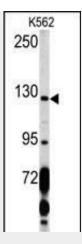
Detected in platelets (at protein level). Widely expressed. Detected in brain, fibroblasts, smooth muscle, heart, and embryo. Expressed in primary and metastatic colon tumors and in normal colon tissue.

## PDGFRA Antibody (C-term E1063) - Protocols

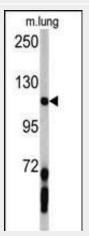
Provided below are standard protocols that you may find useful for product applications.

- <u>Western Blot</u>
- Blocking Peptides
- <u>Dot Blot</u>
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

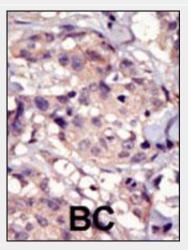
PDGFRA Antibody (C-term E1063) - Images



Western blot analysis of PDGFRA Antibody (C-term E1063) (Cat.# AP7666b) in K562 cell line lysates (35ug/lane). PDGFRA (arrow) was detected using the purified Pab.

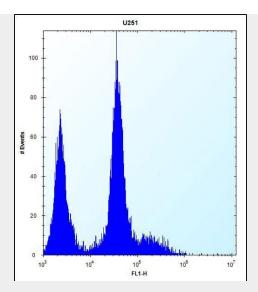


Western blot analysis of PDGFRA antibody (C-term E1063) (Cat.# AP7666b) in mouse lung tissue lysates (35ug/lane). PDGFRA (arrow) was detected using the purified Pab.



Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated. BC = breast carcinoma; HC = hepatocarcinoma.





PDGFRA Antibody (C-term E1063) (Cat. #AP7666b) flow cytometric analysis of U251 cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

## PDGFRA Antibody (C-term E1063) - Background

PDGFRA is a cell surface tyrosine kinase receptor for members of the platelet-derived growth factor family. These growth factors are mitogens for cells of mesenchymal origin. The identity of the growth factor bound to a receptor monomer determines whether the functional receptor is a homodimer or a heterodimer, composed of both platelet-derived growth factor receptor alpha and beta polypeptides. Studies in knockout mice, where homozygosity is lethal, indicate that the alpha form of the platelet-derived growth factor receptor exhibit defective kidney phenotypes.

# PDGFRA Antibody (C-term E1063) - References

Cools, J., et al., N. Engl. J. Med. 348(13):1201-1214 (2003). Heinrich, M.C., et al., Science 299(5607):708-710 (2003). Andrae, J., et al., Biochem. Biophys. Res. Commun. 296(3):604-611 (2002). Ribom, D., et al., J. Neurol. Neurosurg. Psychiatr. 72(6):782-787 (2002). Kawagishi, J., et al., Genomics 30(2):224-232 (1995).