

## PDGFRb (Phospho Tyr857) Polyclonal Antibody

Catalog No: YP1235

**Reactivity:** Human; Mouse; Rat

**Applications:** IHC;IF;WB

Target: PDGFR-β

**Fields:** >>EGFR tyrosine kinase inhibitor resistance;>>MAPK signaling pathway;>>Ras

signaling pathway;>>Rap1 signaling pathway;>>Calcium signaling pathway;>>Phospholipase D signaling pathway;>>Pl3K-Akt signaling pathway;>>Focal adhesion;>>Gap junction;>>JAK-STAT signaling pathway;>>Regulation of actin cytoskeleton;>>Human papillomavirus

infection;>>Pathways in cancer;>>MicroRNAs in cancer;>>Glioma;>>Prostate cancer:>>Melanoma:>>Central carbon metabolism in cancer:>>Choline

metabolism in cancer

Gene Name: PDGFRB PDGFR PDGFR1

**Protein Name:** PDGFRb (Phospho-Tyr857)

P09619

Human Gene Id: 5159

**Human Swiss Prot** 

No:

Immunogen: Synthesized peptide derived from human PDGFRb (Phospho-Tyr857)

**Specificity:** This antibody detects endogenous phospho levels of PDGFRb (Phospho-

Tyr857) at Human: Y857, Mouse: Y856, Rat: Y856

**Formulation:** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

**Source:** Polyclonal, Rabbit, lgG

**Dilution:** IHC 1:50-200, WB 1:500-2000. IF 1:50-200

**Purification:** The antibody was affinity-purified from rabbit serum by affinity-chromatography

using specific immunogen.

**Concentration**: 1 mg/ml

**Storage Stability:** -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 135-180kD

**Background :** This gene encodes 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. This gene is flanked on chromosome 5 by the genes for granulocyte-macrophage colony-stimulating factor and macrophage-colony stimulating factor receptor; all three genes may be implicated in the 5-q syndrome. A translocation between chromosomes 5 and 12, that fuses this gene

to that of the translocation, ETV6, leukemia gene, results in chronic

myeloproliferative disorder with eosinophilia. [provided by RefSeq, Jul 2008],

**Function :** catalytic activity:ATP + a [protein]-L-tyrosine = ADP + a [protein]-L-tyrosine

phosphate., disease: A chromosomal aberration involving PDGFRB is a cause in many instances of chronic myeloproliferative disorder with eosinophilia (MPE) [MIM:131440]. Translocation t(5;12) with ETV6 on chromosome 12 creating an PDGFRB-ETV6 fusion protein., disease: A chromosomal aberration involving PDGFRB is found in a form of chronic myelomonocytic leukemia (CMML). Translocation t(5;12)(q33;p13) with EVT6/TEL. It is characterized by abnormal clonal myeloid proliferation and by progression to acute myelogenous leukemia (AML)., disease: A chromosomal aberration involving PDGFRB may be a cause of acute myelogenous leukemia. Translocation t(5;14)(q33;q32) with TRIP11. The

fusion protein may be involved in clonal evolution of leukemia and

eosinophilia.,disease:A chromosomal aberration involving PDGFRB may be a

cause

Subcellular Location:

Cell membrane; Single-pass type I membrane protein. Cytoplasmic vesicle. Lysosome lumen. After ligand binding, the autophosphorylated receptor is

ubiquitinated and internalized, leading to its degradation.

**Expression :** Brain, Spleen,

**Sort :** 11752

No4: 1

Host: Rabbit

Modifications: Phospho



## **Products Images**