

Catalog #: APRab04464

Summary

Production Name c-Kit (phospho Tyr721) Rabbit Polyclonal Antibody

Description Rabbit Polyclonal Antibody

Host Rabbit
Application WB,ELISA

Reactivity Human, Mouse, Rat

Performance

ConjugationUnconjugatedModificationPhosphorylated

Isotype IgG

Clonality Polyclonal Form Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw Storage

cycles.

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% New type

preservative N.

Purification Affinity purification

Immunogen

Gene Name KIT

KIT; SCFR; Mast/stem cell growth factor receptor Kit; SCFR; Piebald trait protein; PBT;

Alternative Names Proto-oncogene c-Kit; Tyrosine-protein kinase Kit; p145 c-kit; v-kit Hardy-Zuckerman 4

feline sarcoma viral oncogene homolog; CD antigen CD117

Gene ID 3815.0

P10721.The antiserum was produced against synthesized peptide derived from human **SwissProt ID**

c-Kit around the phosphorylation site of Tyr721. AA range:688-737

Application

Dilution Ratio WB 1:500-1:2000, ELISA 1:10000.Not yet tested in other applications.

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838



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Molecular Weight

110kDa

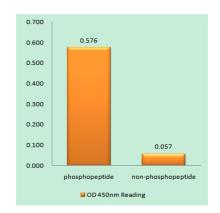
Background

This gene encodes the human homolog of the proto-oncogene c-kit. C-kit was first identified as the cellular homolog of the feline sarcoma viral oncogene v-kit. This protein is a type 3 transmembrane receptor for MGF (mast cell growth factor, also known as stem cell factor). Mutations in this gene are associated with gastrointestinal stromal tumors, mast cell disease, acute myelogenous lukemia, and piebaldism. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008], catalytic activity:ATP + a [protein]-L-tyrosine = ADP + a [protein]-Ltyrosine phosphate, disease: Defects in KIT are a cause of gastrointestinal stromal tumor (GIST) [MIM:606764], disease: Defects in KIT are a cause of piebaldism [MIM:172800]. Piebaldism is an autosomal dominant genetic developmental abnormality of pigmentation characterized by congenital patches of white skin and hair that lack melanocytes, disease: Defects in KIT have been associated with testicular tumors [MIM:273300]. It includes germ cell tumor (GCT) or testicular germ cell tumor (TGCT), function: This is the receptor for stem cell factor (mast cell growth factor). It has a tyrosine-protein kinase activity. Binding of the ligands leads to the autophosphorylation of KIT and its association with substrates such as phosphatidylinositol 3-kinase (Pi3K).,online information:CD117 entry,similarity:Belongs to the protein kinase superfamily. Tyr protein kinase family, similarity: Belongs to the protein kinase superfamily. Tyr protein kinase family. CSF-1/PDGF receptor subfamily, similarity: Contains 1 protein kinase domain, similarity: Contains 5 Iq-like C2-type (immunoglobulin-like) domains., subunit: Interacts with APS. Interacts with MPDZ (via the tenth PDZ domain). Interacts with PTPRU.,

Research Area

Cytokine-cytokine receptor interaction; Endocytosis; Hematopoietic cell lineage; Melanogenesis; Pathways in cancer; Acute myeloid leukemia;

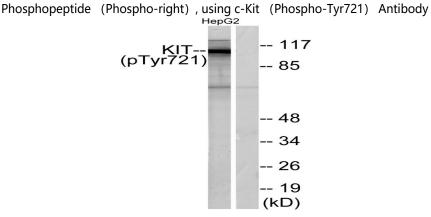
Image Data



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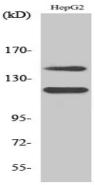
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Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-



Western blot analysis of lysates from HepG2 cells treated with EGF 200ng/ml 30 ', using c-Kit (Phospho-Tyr721)

Antibody. The lane on the right is blocked with the phospho peptide.



Western Blot analysis of various cells using Phospho-c-Kit (Y721) Polyclonal Antibody

Note

For research use only.