

Bcr (phospho Tyr177) Polyclonal Antibody

Catalog No: YP0036

Reactivity: Human; Mouse

Applications: WB;ELISA

Target: Bcr

Fields: >>Pathways in cancer;>>Chronic myeloid leukemia

Gene Name: BCR

Protein Name: Breakpoint cluster region protein

Human Gene Id: 613

Human Swiss Prot

P11274

Q6PAJ1

No:

Mouse Gene ld: 110279

Mouse Swiss Prot

No:

Immunogen: The antiserum was produced against synthesized peptide derived from human

Bcr around the phosphorylation site of Tyr177. AA range:144-193

Specificity: Phospho-Bcr (Y177) Polyclonal Antibody detects endogenous levels of Bcr

protein only when phosphorylated at Y177.

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

Source: Polyclonal, Rabbit, IgG

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

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

1/3



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

Observed Band: 160kD

Cell Pathway: Pathways in cancer; Chronic myeloid leukemia;

Background: A reciprocal translocation between chromosomes 22 and 9 produces the

Philadelphia chromosome, which is often found in patients with chronic myelogenous leukemia. The chromosome 22 breakpoint for this translocation is located within the BCR gene. The translocation produces a fusion protein which is encoded by sequence from both BCR and ABL, the gene at the chromosome 9 breakpoint. Although the BCR-ABL fusion protein has been extensively studied, the function of the normal BCR gene product is not clear. The protein has serine/threonine kinase activity and is a GTPase-activating protein for p21rac.

Two transcript variants encoding different isoforms have been found for this gene.

[provided by RefSeq, Jul 2008],

Function: catalytic activity:ATP + a protein = ADP + a phosphoprotein.,disease:A

chromosomal aberration involving BCR is a cause of chronic myeloid leukemia

(CML) [MIM:608232]. Translocation t(9;22)(q34;q11) with ABL1. The

translocation produces a BCR-ABL found also in acute myeloid leukemia (AML) and acute lymphoblastic leukemia (ALL)..domain:The DH domain is involved in

interaction with CCPG1.,domain:The region involved in binding to ABL1

SH2-domain is rich in serine residues and needs to be Ser/Thr phosphorylated prior to SH2 binding. This region is essential for the activation of the ABL1

tyrosine kinase and transforming potential of the chimeric BCR-ABL

oncogene.,function:GTPase-activating protein for RAC1 and CDC42. Promotes the exchange of RAC or CDC42-bound GDP by GTP, thereby activating them.

Displays serine/threonine kinase

activity.,PTM:Autophosphorylated.,similarity:Contains 1 C2 domai

Subcellular

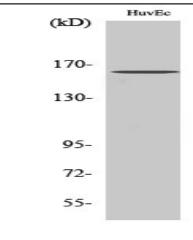
Location:

Cell junction, synapse, postsynaptic density . Cell projection, dendritic spine .

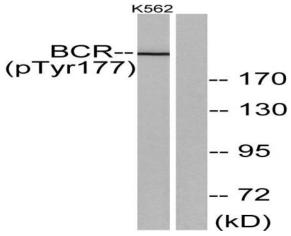
Cell projection, axon . Cell junction, synapse .

Expression: Brain, Epithelium, Platelet, Renal cell carcinoma, T-cell,

Products Images



Western Blot analysis of various cells using Phospho-Bcr (Y177) Polyclonal Antibody



Western blot analysis of lysates from K562 cells, using Bcr (Phospho-Tyr177) Antibody. The lane on the right is blocked with the phospho peptide.