

4.1R (phospho Tyr660) Polyclonal Antibody

Catalog No: YP0468

Reactivity: Human; Mouse

Applications: WB;IHC;IF;ELISA

Target: 4.1R

Gene Name: EPB41

Protein Name: Protein 4.1

P11171

P48193

Human Gene Id: 2035

Human Swiss Prot

No:

Mouse Gene ld: 269587

Mouse Swiss Prot

No:

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

EPB41 around the phosphorylation site of Tyr660/418. AA range:626-675

Specificity: Phospho-4.1R (Y660) Polyclonal Antibody detects endogenous levels of 4.1R

protein only when phosphorylated at Y660.

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. IHC 1:100 - 1:300. ELISA: 1:5000.. IF 1:50-200

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

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

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

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Observed Band: 60kD

Cell Pathway : Tight junction;

Background:

The protein encoded by this gene, together with spectrin and actin, constitute the red cell membrane cytoskeletal network. This complex plays a critical role in erythrocyte shape and deformability. Mutations in this gene are associated with type 1 elliptocytosis (EL1). Alternatively spliced transcript variants encoding different isoforms have been described for this gene.[provided by RefSeq, Oct 2009],

Function:

disease:Defects in EPB41 are a cause of hereditary pyropoikilocytosis (HPP) [MIM:266140]. HPP is an autosomal recessive hematologic disorder characterized by hemolytic anemia, microspherocytosis, poikilocytosis, and an unusual thermal sensitivity of red cells.,disease:Defects in EPB41 are the cause of elliptocytosis type 1 (EL1) [MIM:611804]. EL1 is a Rhesus-linked form of hereditary elliptocytosis, a genetically heterogeneous, autosomal dominant, hematologic disorder. It is characterized by variable hemolytic anemia and elliptical or oval red cell shape.,function:Protein 4.1 is a major structural element of the erythrocyte membrane skeleton. It plays a key role in regulating membrane physical properties of mechanical stability and deformability by stabilizing spectrinactin interaction. Recruits DLG1 to membranes.,PTM:O-glycosylated; contains N-acetylglucosamine side chains in the C-ter

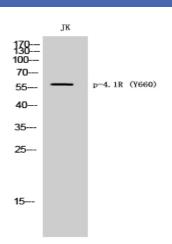
Subcellular Location:

Cytoplasm, cytoskeleton. Cytoplasm, cell cortex. Nucleus.

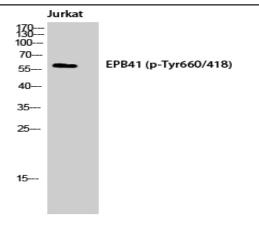
Expression:

Brain, PCR rescued clones, Reticulocyte, Spleen,

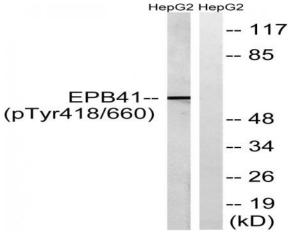
Products Images



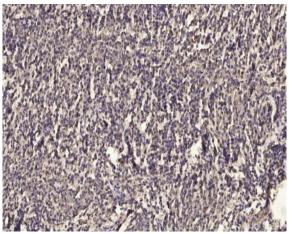
Western Blot analysis of JK cells using Phospho-4.1R (Y660) Polyclonal Antibody



Western Blot analysis of Jurkat cells using Phospho-4.1R (Y660) Polyclonal Antibody



Western blot analysis of lysates from HepG2 cells treated with PMA 125ng/ml 30', using EPB41 (Phospho-Tyr660/418) Antibody. The lane on the right is blocked with the phospho peptide.



Immunohistochemical analysis of paraffin-embedded human meningioma. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).