



4.1R Monoclonal Antibody

Catalog No	YP-mAb-03666
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	EPB41
Immunogen	The antiserum was produced against synthesized peptide derived from human EPB41. AA range:626-675
Specificity	4.1R Monoclonal Antibody detects endogenous levels of 4.1R protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	4.1R;41_HUMAN;Band 4.1;E41P;EL 1;EL1;EL1 gene;Elliptocytosis 1; Elliptocytosis 1 RH linked;EPB 4.1;EPB 41;EPB4.1;Epb41;Erythrocyte membrane protein band 4.1 (elliptocytosis 1 RH linked);Erythrocyte membrane protein band 4.1;Erythrocyte surface protein band 4.1;HE;P4.1;Protein 4.1;red blood cell type
Observed Band	60kD
Calculated Molecular Weight	60kD
Cell Pathway	Cytoplasm, cytoskeleton . Cytoplasm, cell cortex . Nucleus .
Tissue Specificity	Brain,PCR rescued clones,Reticulocyte,Spleen,
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 spectrin-actin interaction. Recruits DLG1 to membranes.,PTM:O-glycosylated; contains N-acetylglucosamine side chains in the C-ter

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],

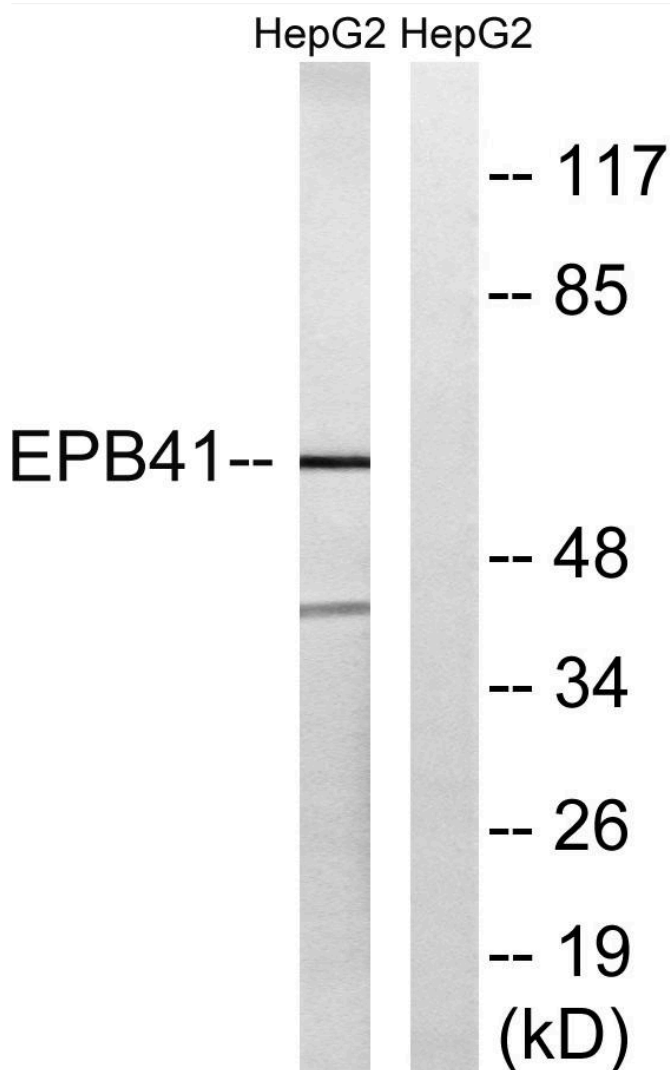
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using 4.1R Monoclonal Antibody