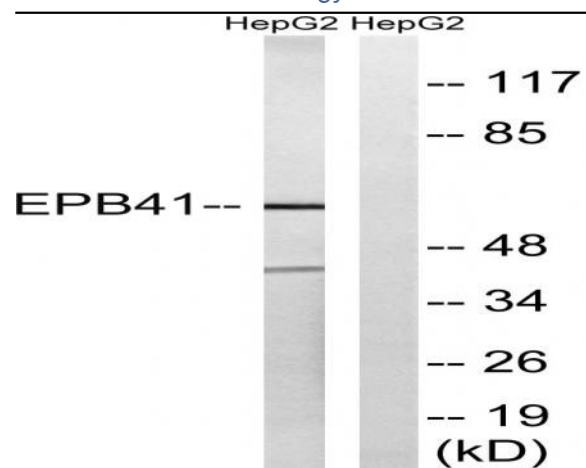


4.1R Polyclonal Antibody

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|------------------------------|---|
| Catalog No : | YT0017 |
| Reactivity : | Human;Mouse |
| Applications : | WB;ELISA |
| Target : | 4.1R |
| Gene Name : | EPB41 |
| Protein Name : | Protein 4.1 |
| Human Gene Id : | 2035 |
| Human Swiss Prot No : | P11171 |
| Mouse Gene Id : | 269587 |
| Mouse Swiss Prot No : | P48193 |
| Immunogen : | The antiserum was produced against synthesized peptide derived from human EPB41. AA range:626-675 |
| Specificity : | 4.1R Polyclonal Antibody detects endogenous levels of 4.1R protein. |
| 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: 1:5000. 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 |
| 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 spectrin-actin 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, |
| Tag : | orthogonal |
| Sort : | 1512 |
| No4 : | 1 |
| Host : | Rabbit |
| Modifications : | Unmodified |

Products Images



Western blot analysis of lysates from HepG2 cells treated with PMA 125ng/ml 30', using EPB41 Antibody. The lane on the right is blocked with the synthesized peptide.