

EPB41 (Human) Recombinant Protein (Q01)

Catalog # H00002035-Q01 Size 25 ug, 10 ug

Applications



Specification	
Product Description	Human EPB41 partial ORF (AAH39079.1, 116 a.a 225 a.a.) recombinant protein with GST tag at N-terminal.
Sequence	IEFGTSLDEEIILKAPIAAPEPELKTDPSLDLHSLSSAETQPAQEELREDPDFEIKEGEGLEECSKIE VKEESPQSKAETELKASQKPIRKHRNMHCKVSLLDDTVYECV
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	37.84
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.

Applications



- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — EPB41	
Entrez GenelD	2035
GeneBank Accession#	BC039079.1
Protein Accession#	AAH39079.1
Gene Name	EPB41
Gene Alias	4.1R, EL1, HE
Gene Description	erythrocyte membrane protein band 4.1 (elliptocytosis 1, RH-linked)
Omim ID	130500
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Elliptocytosis is a hematologic disorder characterized by elliptically shaped erythrocytes and a var iable degree of hemolytic anemia. Inherited as an autosomal dominant, elliptocytosis results from mutation in any one of several genes encoding proteins of the red cell membrane skeleton. The form discussed here is the one found in the 1950s to be linked to Rh blood group and more recently shown to be caused by a defect in protein 4.1. 'Rh-unlinked' forms of elliptocytosis are caused by mutation in the alpha-spectrin gene (MIM 182860), the beta-spectrin gene (MIM 182870), or the b and 3 gene (MIM 109270).[supplied by OMIM
Other Designations	OTTHUMP0000003772 OTTHUMP0000003773 OTTHUMP00000003774 erythrocyte surface protein band 4.1

Pathway

• Tight junction