

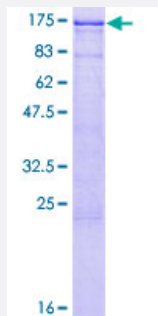
Full-Length

EPB41 (Human) Recombinant Protein (P02)

Catalog # H00002035-P02

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human EPB41 full-length ORF (AAH39079.1, 1 a.a. - 720 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MTTEKSLVTEAENSQHQQKEEGEEAINSGQQEPQQEESCQTAAEGDNWCEQKLKASNGDTPTH
EDLTKNKERTSESRLSRLFSSFLKRPKSQVSEEEGKEVESDKEKGEGGQKEIEFGTSLDEEILK
APIAAPEPELKTDPSLDLHSLSSAETQPAQEELREDPDFEIKEGEGLEESKIEVKEESPQSKAE
TELKASQKPIRKHRNMHCKVSLDDTVYECVVEKHAKGQDLLKRVCEHLNLLEEDYFGLAWDNA
TSKTWLD SAKEIKKQVRGVPWNFTFNVKFYPPDPAQLTEDITRYLCLQLRQDIVAGRLPCSFATL
ALLGSYTIQSELGDYDPELHGVDYVSDFKLAPNQTKELKVMELHKSYSRSMTPAQADLEFLNA
KKLSMYGVDLHKAKDLEGVDIILGVCSSGLLVYKDKLRINRFPWPKVLKISYKRSSFFIKIRPGEQE
QYESTIGFKLPSYRAAKKLWKVCVEHHTFFRLTSTDITPKSKFLALGSKFRYSGRTQAQTRQASALI
DRPAPHFERTASKRASRSLDGAAVDSADRSRPTSAPAITQGQVAEGGVLDASAKKTVVPAK
QKETVKA EVKKEDEPPEQAEPEPTAEWKVEKTHIEVTPTSNGDQTQKKRERLDGENIYRHSNL
MLEDLDKSQEEIKKHASISELKKNFMESVPEPRPSEWDKRLSTHSPFRTLNINGQIPTGEGVSTL
ST

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

107.6

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 GeneID	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	Hyperlink

Gene Summary	Elliptocytosis is a hematologic disorder characterized by elliptically shaped erythrocytes and a variable 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]
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Other Designations

OTTHUMP00000003772|OTTHUMP00000003773|OTTHUMP00000003774|erythrocyte surface protein band 4.1

Pathway

- [Tight junction](#)