

#### **UK Office**

#### **Everest Biotech Ltd**

Cherwell Innovation Centre 77 Heyford Park Upper Heyford Oxfordshire OX25 5HD

UK

**Enquiries:** 

info@everestbiotech.com

Sales:

sales@everestbiotech.com

Tech support:

support@everestbiotech.com

Tel: +44 (0)1869 238326

www.everestbiotech.com

Research Use Only. Not for diagnostic or therapeutic use.

# EB08237 - Goat Anti-EBPL41L5 Antibody

Size: 100µg specific antibody in 200µl



### **Target Protein**

Principal Names: EPB41L5, erythrocyte membrane protein band 4.1 like 5, BE37,

FLJ12957, KIAA1548 Official Symbol: EPB41L5

Accession Number(s): NP\_065960.2; NP\_001171866.1; NP\_001171868.1;

NP\_001317239.1

Human GenelD(s): 57669

### **Immunogen**

Peptide with sequence C-ENLPQSPGTDQHD, from the internal region of the protein sequence according to NP\_065960.2; NP\_001171866.1; NP\_001171868.1; NP\_001317239.1.

Please note the peptide is available for sale.

### **Purification and Storage**

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

## **Applications Tested**

Peptide ELISA: antibody detection limit dilution 1:8000.

**Western blot:** Approx. 80kDa band observed in lysates of cell line A431 and approx. 75kDa in lysates of cell line MCF7 (calculated MW of 81.9kDa according to NP\_065960.2). Recommended concentration: 0.3-1µg/ml. Primary incubation 1 hour at room temperature.

**Immunofluorescence:** Strong expression of the protein seen in U2OS and A549 cells. Recommended concentration: 10µg/ml.

**Flow Cytometry:** Flow cytometric analysis of A549 cells. Recommended concentration: 10ug/ml.

### **Species Reactivity**

Tested: Human

Expected from sequence similarity: Human

EB08237 (1μg/ml) staining of A431 (A) and (0.3ug/ml) MCF7 (B) cell lysate (35μg protein in RIPA buffer).

Detected by chemiluminescence.

EB08237 Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing nuclear, plasma membrane and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

EB08237 Immunofluorescence analysis of paraformaldehyde fixed A549 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).

EB08237 Flow cytometric analysis of paraformaldehyde fixed A549 cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control:

Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.