

Product Data Sheet

Anti-P-LAP Antibody

Catalog # Source Reactivity Applications

CQA4962 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to P-LAP

Immunogen Recombinant fusion protein of human P-LAP. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of P-LAP protein

Clonality Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/2000), IH (1/50 - 1/200)

Gene Symbol LNPEP

Alternative Names OTASE; Leucyl-cystinyl aminopeptidase; Cystinyl aminopeptidase; Insulin-regulated

membrane aminopeptidase; Insulin-responsive aminopeptidase; IRAP; Oxytocinase;

OTase; Placental leucine aminopeptidase; P-LAP

Entrez Gene 4012 (Human); 240028 (Mouse); 100912445171105 (Rat)

SwissProt Q9UIQ6 (Human); Q8C129 (Mouse); P97629 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

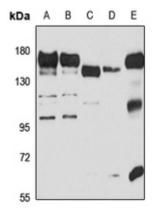
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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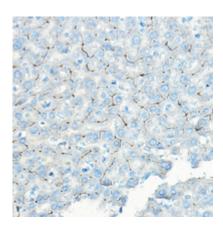
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Western blot analysis of P-LAP expression in HeLa (A), Jurkat (B), mouse brain (C), mouse liver (D), rat heart (E) whole cell lysates. (Predicted band size: 115; 117 kD; Observed band size: 165 kD)



Immunohistochemical analysis of P-LAP staining in rat liver formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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