

Product Data Sheet

Anti-Pancreatic Lipase Antibody

| Catalog # | Source | Reactivity | Applications |
|----------------|--------|--------------------------------|--|
| CQA1362 | Rabbit | H, M, R | WB, IH |
| Description | | Rabbit polyclonal antibody | to Pancreatic Lipase |
| Immunogen | | Recombinant full length pro | tein of human Pancreatic Lipase |
| Purification | | The antibody was purified b | y immunogen affinity chromatography. |
| Specificity | | Recognizes endogenous lev | els of Pancreatic Lipase protein. |
| Clonality | | Polyclonal | |
| Conjugation | | | |
| Form | | Liquid in 0.42% Potassium p | hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, |
| | | and 0.01% sodium azide. | |
| Dilution | | WB (1/500 - 1/1000), IH (1/50 |) - 1/100) |
| Gene Symbol | | PNLIP | |
| Alternative Na | ames | Pancreatic triacylglycerol lip | ase; PL; PTL; Pancreatic lipase |
| Entrez Gene | | 5406 (Human); 69060 (Mou | se); 25702 (Rat) |
| SwissProt | | P16233 (Human); Q6P8U6 (| Mouse); P27657 (Rat) |
| Storage/Stabi | lity | Shipped at 4°C. Upon delive | ry 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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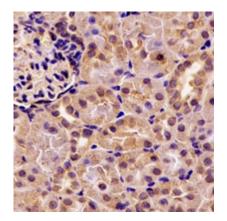
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Western blot analysis of Pancreatic Lipase expression in mouse liver (A), mouse pancreas (B) whole cell lysates. (Predicted band size: 51 kD; Observed band size: 49 kD)



Immunohistochemical analysis of Pancreatic Lipase staining in rat kidney 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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