

Product Data Sheet

Anti-GNE Antibody

Catalog # Source Reactivity Applications

CPA5661 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to GNE

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human GNE. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of GNE 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 - 1/1000), IH (1/50 - 1/100)

Gene Symbol GNE

Alternative Names GLCNE; Bifunctional UDP-N-acetylglucosamine 2-epimerase/N-acetylmannosamine

kinase; UDP-GlcNAc-2-epimerase/ManAc kinase

Entrez Gene 10020 (Human); 50798 (Mouse); 114711 (Rat)

SwissProt Q9Y223 (Human); Q91WG8 (Mouse); O35826 (Rat)

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

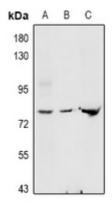
freeze/thaw cycles.

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

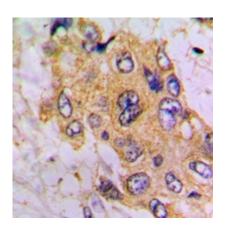
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Western blot analysis of GNE expression in LO2 (A), mouse liver (B), rat liver (C) whole cell lysates. (Predicted band size: 79 kD; Observed band size: 79 kD)



Immunohistochemical analysis of GNE staining in human breast cancer 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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