

Product Data Sheet

Anti-CLNS1A Antibody

Catalog # Source Reactivity Applications

CPA1245 Rabbit H, M, R, D WB, IH

Description Rabbit polyclonal antibody to CLNS1A

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

region of human CLNS1A. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of CLNS1A 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 CLNS1A

Alternative Names CLCI; ICLN; Methylosome subunit pICln; Chloride channel, nucleotide sensitive 1A;

Chloride conductance regulatory protein ICln; I(Cln); Chloride ion current inducer

protein; CICI; Reticulocyte pICIn

Entrez Gene 1207 (Human)

SwissProt P54105 (Human)

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

freeze/thaw cycles.

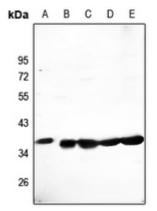
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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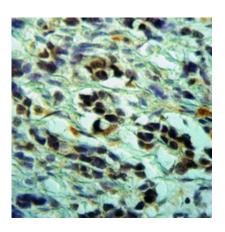
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Western blot analysis of CLNS1A expression in H9C2 (A), AML12 (B), Jurkat (C), HepG2 (D), A549 (E) whole cell lysates. (Predicted band size: 26 kD; Observed band size: 39 kD)



Immunohistochemical analysis of CLNS1A staining in human stomach 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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