

Product Data Sheet

Anti-ZNF638 Antibody

Catalog # Source Reactivity Applications

CPA2631 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to ZNF638

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human ZNF638. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of ZNF638 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 ZNF638

Alternative Names NP220; ZFML; Zinc finger protein 638; Cutaneous T-cell lymphoma-associated

antigen se33-1; CTCL-associated antigen se33-1; Nuclear protein 220; Zinc finger

matrin-like protein

Entrez Gene 27332 (Human); 18139 (Mouse)

SwissProt Q14966 (Human); Q61464 (Mouse)

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

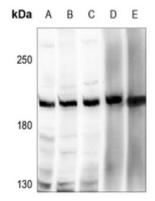
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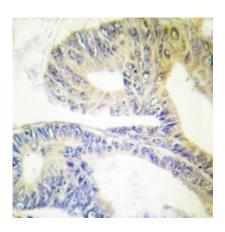




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Western blot analysis of ZNF638 expression in PC3 (A), A549 (B), HEK293T (C), rat brain (D), mouse brain (E) whole cell lysates. (Predicted band size: 220 kD; Observed band size: 220; 120 kD)



Immunohistochemical analysis of ZNF638 staining in human colon 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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