

Product Data Sheet

Anti-APE1 Antibody

Catalog # Source Reactivity Applications

CPA1058 Rabbit H, M, R, B, P WB, IH

Description Rabbit polyclonal antibody to APE1

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

region of human APE1. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of APE1 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 (1/100 - 1/200)

Gene Symbol APEX1

Alternative Names APE; APE1; APEX; APX; HAP1; REF1; DNA-(apurinic or apyrimidinic site) lyase; APEX

nuclease; APEN; Apurinic-apyrimidinic endonuclease 1; AP endonuclease 1; APE-1;

REF-1; Redox factor-1

Entrez Gene 328 (Human); 11792 (Mouse); 79116 (Rat)

SwissProt P27695 (Human); P28352 (Mouse); P43138 (Rat)

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

freeze/thaw cycles.

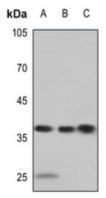
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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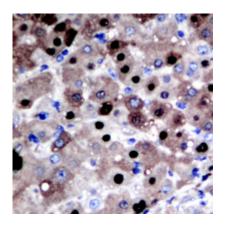
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Western blot analysis of APE1 expression in Hela (A), A549 (B), rat lung (C) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 37 kD)



Immunohistochemical analysis of APE1 staining in human liver 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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