

Anti-APE1 Antibody

Catalog # AP95671

Product Information

Application	WB, IHC, FC
Primary Accession	P27695
Reactivity	Human, Rat, Mk
Host	Mouse
Clonality	Monoclonal
Calculated MW	35555

Additional Information

Gene ID	328
Other 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
Target/Specificity	Recombinant fusion protein of human APE1 expressed in E. Coli
Dilution	WB~~WB (1/500 - 1/1000) IHC~~1:100~500 FC~~FC (1/100 - 1/200)
Format	Mouse IgG1. Liquid in PBS, pH 7.3, 30% glycerol, and 0.01% sodium azide.
Storage	Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name	APEX1
Synonyms	APE, APE1, APEX, APX, HAP1, REF1
Function	Multifunctional protein that plays a central role in the cellular response to oxidative stress. The two major activities of APEX1 are DNA repair and redox regulation of transcriptional factors (PubMed: 11118054 , PubMed: 11452037 , PubMed: 15831793 , PubMed: 18439621 , PubMed: 18579163 , PubMed: 21762700 , PubMed: 24079850 , PubMed: 8355688 , PubMed: 9108029 , PubMed: 9560228). Functions as an apurinic/apyrimidinic (AP) endodeoxyribonuclease in the base excision repair (BER) pathway of DNA lesions induced by oxidative and alkylating agents. Initiates repair of AP sites in DNA by catalyzing hydrolytic incision of the phosphodiester backbone immediately adjacent to the damage, generating a single-strand break with 5'-deoxyribose phosphate and 3'-hydroxyl ends. Also incises at AP sites in the DNA strand of DNA/RNA hybrids, single-stranded DNA regions of R-loop structures, and single-stranded RNA molecules (PubMed: 15380100 , PubMed: 16617147 , PubMed: 18439621 , PubMed: 19123919 ,

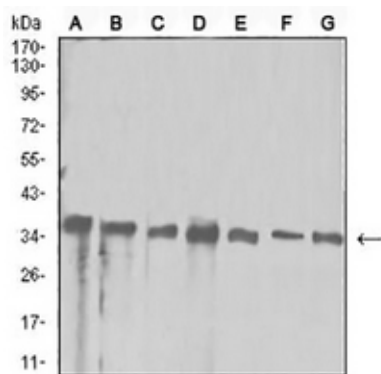
PubMed:[19188445](#), PubMed:[19934257](#), PubMed:[20699270](#), PubMed:[21762700](#), PubMed:[24079850](#), PubMed:[8932375](#), PubMed:[8995436](#), PubMed:[9804799](#)). Operates at switch sites of immunoglobulin (Ig) constant regions where it mediates Ig isotype class switch recombination. Processes AP sites induced by successive action of AICDA and UNG. Generates staggered nicks in opposite DNA strands resulting in the formation of double-strand DNA breaks that are finally resolved via non-homologous end joining repair pathway (By similarity). Has 3'-5' exodeoxyribonuclease activity on mismatched deoxyribonucleotides at the 3' termini of nicked or gapped DNA molecules during short-patch BER (PubMed:[11832948](#), PubMed:[1719477](#)). Possesses DNA 3' phosphodiesterase activity capable of removing lesions (such as phosphoglycolate and 8-oxoguanine) blocking the 3' side of DNA strand breaks (PubMed:[15831793](#), PubMed:[7516064](#)). Also acts as an endoribonuclease involved in the control of single-stranded RNA metabolism. Plays a role in regulating MYC mRNA turnover by preferentially cleaving in between UA and CA dinucleotides of the MYC coding region determinant (CRD). In association with NMD1, plays a role in the rRNA quality control process during cell cycle progression (PubMed:[19188445](#), PubMed:[19401441](#), PubMed:[21762700](#)). Acts as a loading factor for POLB onto non-incised AP sites in DNA and stimulates the 5'-terminal deoxyribose 5'-phosphate (dRp) excision activity of POLB (PubMed:[9207062](#)). Exerts reversible nuclear redox activity to regulate DNA binding affinity and transcriptional activity of transcriptional factors by controlling the redox status of their DNA-binding domain, such as the FOS/JUN AP-1 complex after exposure to IR (PubMed:[10023679](#), PubMed:[11118054](#), PubMed:[11452037](#), PubMed:[18579163](#), PubMed:[8355688](#), PubMed:[9108029](#)). Involved in calcium-dependent down-regulation of parathyroid hormone (PTH) expression by binding to negative calcium response elements (nCaREs). Together with HNRNPL or the dimer XRCC5/XRCC6, associates with nCaRE, acting as an activator of transcriptional repression (PubMed:[11809897](#), PubMed:[14633989](#), PubMed:[8621488](#)). May also play a role in the epigenetic regulation of gene expression by participating in DNA demethylation (PubMed:[21496894](#)). Stimulates the YBX1-mediated MDR1 promoter activity, when acetylated at Lys-6 and Lys-7, leading to drug resistance (PubMed:[18809583](#)). Plays a role in protection from granzyme-mediated cellular repair leading to cell death (PubMed:[18179823](#)). Binds DNA and RNA. Associates, together with YBX1, on the MDR1 promoter. Together with NPM1, associates with rRNA (PubMed:[19188445](#), PubMed:[19401441](#), PubMed:[20699270](#)).

Cellular Location

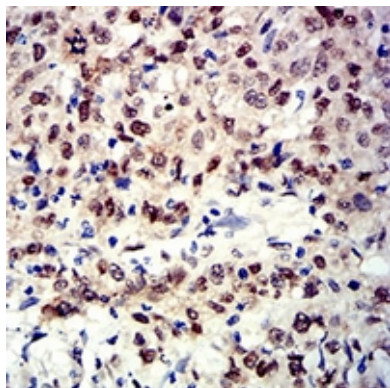
Nucleus {ECO:0000255|PROSITE-ProRule:PRU00764}. Nucleus, nucleolus. Nucleus speckle. Endoplasmic reticulum. Cytoplasm Note=Detected in the cytoplasm of B-cells stimulated to switch (By similarity). Colocalized with SIRT1 in the nucleus. Colocalized with YBX1 in nuclear speckles after genotoxic stress. Together with OGG1 is recruited to nuclear speckles in UVA-irradiated cells. Colocalized with nucleolin and NPM1 in the nucleolus. Its nucleolar localization is cell cycle dependent and requires active rRNA transcription. Colocalized with calreticulin in the endoplasmic reticulum. Translocation from the nucleus to the cytoplasm is stimulated in presence of nitric oxide (NO) and function in a CRM1-dependent manner, possibly as a consequence of demasking a nuclear export signal (amino acid position 64-80). S-nitrosylation at Cys-93 and Cys-310 regulates its nuclear-cytosolic shuttling. Ubiquitinated form is localized predominantly in the cytoplasm.

Images

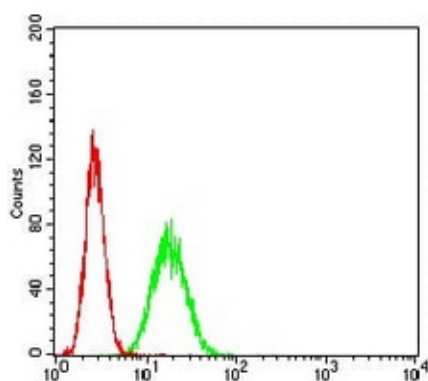
Western blot analysis of APE1 expression in Hela (A), Jurkat (B), SW480 (C), A431 (D), HepG2 (E), NIH/3T3 (F),



PC12 (G) whole cell lysates. (Predicted band size: 36 kD;
Observed band size: 36 kD)



Immunohistochemical analysis of APE1 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.



Flow cytometric analysis of HeLa cells using Anti-APE1
Antibody (green) and negative control (red).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.