abcam

Product datasheet

Recombinant Human DNA polymerase eta protein ab 132167

1 Image

Description

Product name Recombinant Human DNA polymerase eta protein

Expression system Wheat germ
Accession Q9Y253-2

Protein length Full length protein

Animal free No.

Nature Recombinant

Species Human

Sequence MATGQDRVVALVDMDCFFVQVEQRQNPHLRNKPCAVVQ

YKSWKGGGIAV

SYEARAFGVTRSMWADDAKKLCPDLLLAQVRESRGKAN

LTKYREASVEVM

EIMSRFAVIERASIDEAYVDLTSAVQERLQKLQGQPISADLL

PSTYIEGL

PQGPTTAEETVQKEGMRKQGLFQWLDSLQIDNLTSPDLQ

LTVGAVIVEEM

RAAIERETGFQCSAGISHNKVLAKLACGLNKPNRQTLVSH

GSVPQLFSQM

PIRKIRSLGGKLGASVIEILGIEYMGELTQFTESQLQSHFGE

KNGSWLYA

MCRGIEHDPVKPRQLPKTIGCSKNFPGKTALATREQVQW

WLLQLAQELEE

RLTKDRNDNDRVATQLVVSIRVQGDKRLSSLRRCCALTRY

DAHKMSHDAF TVIKNCNTSGIQTE

Predicted molecular weight 71 kDa including tags

Amino acids 1 to 414

Specifications

Our Abpromise guarantee covers the use of ab132167 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Western blot

1

SDS-PAGE

ELISA

Form

Liquid

Additional notes

Preparation and Storage

Stability and Storage

Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.31% Glutathione, 0.79% Tris HCI

General Info

Function

DNA polymerase specifically involved in DNA repair. Plays an important role in translesion synthesis, where the normal high fidelity DNA polymerases cannot proceed and DNA synthesis stalls. Plays an important role in the repair of UV-induced pyrimidine dimers. Depending on the context, it inserts the correct base, but causes frequent base transitions and transversions. May play a role in hypermutation at immunoglobulin genes. Forms a Schiff base with 5'-deoxyribose phosphate at abasic sites, but does not have lyase activity. Targets POLI to replication foci.

Involvement in disease

Defects in POLH are the cause of xeroderma pigmentosum variant type (XPV) [MIM:278750]; also designated as XP-V. Xeroderma pigmentosum (XP) is an autosomal recessive disease due to deficient nucleotide excision repair. It is characterized by hypersensitivity of the skin to sunlight, followed by high incidence of skin cancer and frequent neurologic abnormalities. XPV shows normal nucleotide excision repair, but an exaggerated delay in recovery of replicative DNA synthesis. Most XPV patients do not develop clinical symptoms and skin neoplasias until a later age. Clinical manifestations are limited to photo-induced deterioration of the skin and eyes.

Sequence similarities

Belongs to the DNA polymerase type-Y family.

Contains 1 umuC domain.

Domain

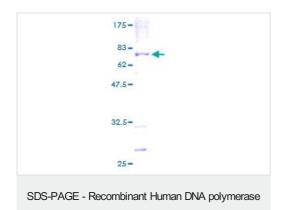
The catalytic core consists of fingers, palm and thumb subdomains, but the fingers and thumb subdomains are much smaller than in high-fidelity polymerases; residues from five sequence motifs of the Y-family cluster around an active site cleft that can accommodate DNA and nucleotide substrates with relaxed geometric constraints, with consequently higher rates of

misincorporation and low processivity.

Cellular localization

Nucleus. Accumulates at replication forks after DNA damage.

Images



12.5% SDS-PAGE analysis of ab132167 stained with Coomassie Blue.

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eta protein (ab132167)

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