

Product datasheet

Recombinant Human DNA polymerase eta protein
ab132167

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	<p>MATGQDRVVALVDMDCFFVQVEQRQNPHLRNKPCAVVQ YKSWKGGGIIAV SYEARAFGVTRSMWADDAKKLCPDLLLAQVRESRGKAN LTKYREASVEVM EIMSRFAVIERASIDEAYVDLTSAVQERLQKLQGQPISADLL PSTYIEGL PQGPTTAEETVQKEGMRKQGLFQWLDLQIDNLTSPDLQ LTVGAVVEEM RAAIERETGFQCSAGISHNKVLAKLACGLNKNRQTLVSH GSVPQLFSQM PIRKIRSLGGKLGASVIEILGIEYMGELTQFTESQLQSHFGE KNGSWLYA MCRGIEHDPVKPRQLPKTIGCSKNFPGKTALATREQVQW WLLQLAQELEE RLTKDRNDNDRVATQLVVSIRVQGDKRLSSLRRCALTRY DAHKMSHDAF TVIKNCNTSGIQTE</p>
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

	SDS-PAGE
	ELISA
Form	Liquid
Additional notes	

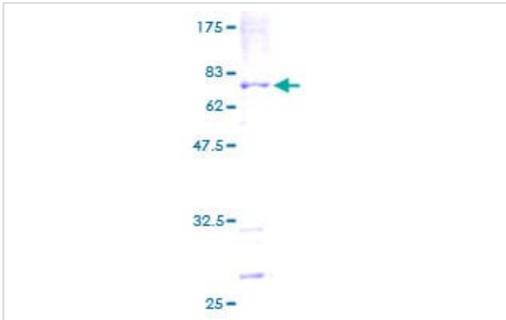
Preparation and Storage

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 HCl
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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.

SDS-PAGE - Recombinant Human DNA polymerase eta protein (ab132167)

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