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WRNIP1 Rabbit pAb

Catalog No.: A9170

Basic Information

Observed MW

72kDa

Calculated MW

72kDa

Category

Primary antibody

Applications

ELISA,WB

Cross-Reactivity

Human

Background

Werner's syndrome is a rare autosomal recessive disorder characterized by accelerated aging that is caused by defects in the Werner syndrome ATP-dependent helicase gene (WRN). The protein encoded by this gene interacts with the exonuclease-containing N-terminal portion of the Werner protein. This protein has a ubiquitin-binding zinc-finger domain in the N-terminus, an ATPase domain, and two leucine zipper motifs in the C-terminus. It has sequence similarity to replication factor C family proteins and is conserved from E. coli to human. This protein likely accumulates at sites of DNA damage by interacting with polyubiquinated proteins and also binds to DNA polymerase delta and increases the initiation frequency of DNA polymerase delta-mediated DNA synthesis. This protein also interacts with nucleoporins at nuclear pore complexes. Two transcript variants encoding different isoforms have been isolated for this gene.

Recommended Dilutions

WB

1:1000 - 1:2000

Immunogen Information

Gene ID 56897

Swiss Prot

Q96S55

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 401-640 of human WRNIP1 (NP_569079.1).

Synonyms

WHIP; FAP93; CFAP93; bA420G6.2; WRNIP1

Contact

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Product Information

Source Rabbit **Purification**

Affinity purification

Storage

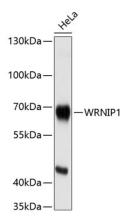
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.

Isotype

IgG

Validation Data



Western blot analysis of extracts of HeLa cells, using WRNIP1 antibody (A9170) at 1:1000 dilution.

Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.