

## Product datasheet for **TP302950L**

### WHIP (WRNIP1) (NM\_130395) Human Recombinant Protein

#### Product data:

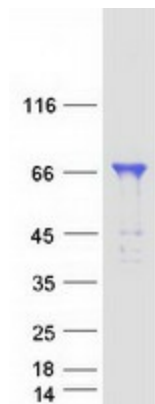
Product Type:	Recombinant Proteins
Description:	Recombinant protein of human Werner helicase interacting protein 1 (WRNIP1), transcript variant 2, 1 mg
Species:	Human
Expression Host:	HEK293T
Expression cDNA Clone or AA Sequence:	>RC202950 representing NM_130395 <b>Red</b> =Cloning site <b>Green</b> =Tags(s)
	<p>MEVSGPEDDPFLSQLHQVQCPVCQQMMPAAHINSHLDRCLLLHPAGHAEPAAAGSHRAGERAKGPSPPGAK RRRLSESSALKQPATPTAAESSEGEEGEGDDGGGETESRESYDAPPTPSGARLIPDFPVARSSSPGRKGS KRPAAAAAAGSASPRSWDEAEAQEEEEAVGDGDGDADADGEDDPGHWADAAEAATAFGASGGGRPH RALAAEEIRQMLQGKPLADTMRPDTLQDYFGQSKAVGQDTLLRSLETNEIPSLILWGPPGCGKTTLAHI IASNSKKHSIRFVTLSATNAKTNDVRDVIKQAQNEKSFKRKTILFIDEIHRFNKSQQVNAALLSRCRVI VLEKLPVEAMVTILMRAINS LGIHVLDSSRPTDPLSHSSNSSEPEAMFIEDKAVDTLAYLSDGDARAGLN GLQLAVLARLSSRKMFCCKSGQSYSPSRVLITENDVKEGLQRSHILYDRAGEEHYNCISALHKSMRGS NASLYWLARMLEGGEDPLYVARRLVRFASEDIGLADPSALTQAVAAAYQGCHFIGMPECEVLLAQC VYFA RAPKSIEVYSAYNNVKA CLRNHQGPLPPVPLHLRNAPTRLMKDLGYGKGYKYNPMYSE PVDQEYLPEELR GVDFFKQRR</p> <p><b>TRTRPLEQKLISEEDLAANDILDYKDDDDKV</b></p>
Tag:	C-Myc/DDK
Predicted MW:	69.3 kDa
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol
Preparation:	Recombinant protein was captured through anti-DDK affinity column followed by conventional chromatography steps.
Note:	For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process.
Storage:	Store at -80°C.



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<b>Stability:</b>	Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles.
<b>RefSeq:</b>	<a href="#">NP_569079</a>
<b>Locus ID:</b>	56897
<b>UniProt ID:</b>	<a href="#">Q96S55</a>
<b>RefSeq Size:</b>	2592
<b>Cytogenetics:</b>	6p25.2
<b>RefSeq ORF:</b>	1920
<b>Synonyms:</b>	bA420G6.2; CFAP93; FAP93; WHIP
<b>Summary:</b>	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 polyubiquitinated 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. [provided by RefSeq, Jul 2012]

### Product images:



Coomassie blue staining of purified WRNIP1 protein (Cat# [TP302950]). The protein was produced from HEK293T cells transfected with WRNIP1 cDNA clone (Cat# [RC202950]) using MegaTran 2.0 (Cat# [TT210002]).