

For Research Use Only

# NBN / NBS1 Recombinant antibody

Catalog Number: 82884-1-RR



## Basic Information

<b>Catalog Number:</b> 82884-1-RR	<b>GenBank Accession Number:</b> BC136803	<b>Purification Method:</b> Protein A purification
<b>Size:</b> 100ul , Concentration: 1000 µg/ml by Nanodrop;	<b>GeneID (NCBI):</b> 4683	<b>CloneNo.:</b> 1C16
<b>Source:</b> Rabbit	<b>Full Name:</b> nibrin	<b>Recommended Dilutions:</b> WB 1:2000-1:10200
<b>Isotype:</b> IgG	<b>Calculated MW:</b> 85 kDa	
	<b>Observed MW:</b> 90-95 kDa	

## Applications

<b>Tested Applications:</b> WB, ELISA	<b>Positive Controls:</b> WB: Jurkat cells, HeLa cells, A549 cells, A431 cells,
<b>Species Specificity:</b> Human	

## Background Information

NBN, also named as NBS, NBS1, and P95, is a component of the MRE11/RAD50/NBN (MRN complex) which plays a critical role in the cellular response to DNA damage and the maintenance of chromosome integrity. The complex is involved in double-strand break (DSB) repair, DNA recombination, maintenance of telomere integrity, cell cycle checkpoint control, and meiosis. The complex possesses single-strand endonuclease activity and double-strand-specific 3'-5' exonuclease activity, which are provided by MRE11A. NBN modulates the DNA damage signal sensing by recruiting PI3/PI4-kinase family members ATM, ATR, and probably DNA-PKcs to the DNA damage sites and activating their functions. NBN also functions in telomere length maintenance by generating the 3' overhang which serves as a primer for telomerase-dependent telomere elongation. NBN is a major player in the control of intra-S-phase checkpoints and there is some evidence that NBN is involved in G1 and G2 checkpoints. Defects in NBN are the cause of Nijmegen breakage syndrome (NBS). Defects in NBN are a cause of genetic susceptibility to breast cancer (BC). Defects in NBN may be associated with aplastic anemia. Defects in NBN might play a role in the pathogenesis of childhood acute lymphoblastic leukemia (ALL). The antibody is specific to NBN. The full-length NBN protein, with an apparent molecular weight of 95 kDa and the two protein fragments of 26 and 70 kDa arising from the c.657\_661del5 (p.K219fsX19) mutation, and the 80 kDa protein found in patient RR with the mutation c.742\_743insGG leading to excision of exons 6 and 7 from the NBN mRNA are shown. (PMID: 26265251) The predicted molecular weight of NBN protein (p95) is 85kDa, and actually detection result is about 95kDa (PMID: 23762398).

## Storage

**Storage:**  
Store at -20°C. Stable for one year after shipment.  
**Storage Buffer:**  
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.  
Aliquoting is unnecessary for -20°C storage

\*\*\* 20ul sizes contain 0.1% BSA

For technical support and original validation data for this product please contact:  
T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA) E: proteintech@ptglab.com W: ptglab.com

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## Selected Validation Data