

NBN Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP11776b

Specification

NBN Antibody (C-term) - Product Information

Application WB, FC, IF, E **Primary Accession** 060934 NP 002476.2 Other Accession Reactivity Human Host **Rabbit** Clonality **Polyclonal** Isotype Rabbit IgG Calculated MW 84959 Antigen Region 602-630

NBN Antibody (C-term) - Additional Information

Gene ID 4683

Other Names

Nibrin, Cell cycle regulatory protein p95, Nijmegen breakage syndrome protein 1, NBN, NBS, NBS1, P95

Target/Specificity

This NBN antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 602-630 amino acids from the C-terminal region of human NBN.

Dilution

WB~~1:1000 FC~~1:10~50 IF~~1:10~50

E~~Use at an assay dependent concentration.

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

NBN Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

NBN Antibody (C-term) - Protein Information



Name NBN (HGNC:7652)

Function Component of the MRN complex, which plays a central role in double-strand break (DSB) repair, DNA recombination, maintenance of telomere integrity and meiosis (PubMed:10888888, PubMed: 15616588, PubMed: 18411307, PubMed: 18583988, PubMed: 18678890, PubMed: <u>19759395</u>, PubMed: <u>23115235</u>, PubMed: <u>28216226</u>, PubMed: <u>28867292</u>, PubMed: <u>9705271</u>). The MRN complex is involved in the repair of DNA double-strand breaks (DSBs) via homologous recombination (HR), an error-free mechanism which primarily occurs during S and G2 phases (PubMed: 19759395, PubMed: 28867292, PubMed: 9705271). The complex (1) mediates the end resection of damaged DNA, which generates proper single-stranded DNA, a key initial steps in HR, and is (2) required for the recruitment of other repair factors and efficient activation of ATM and ATR upon DNA damage (PubMed: 19759395, PubMed: 9705271). The MRN complex possesses single-strand endonuclease activity and double-strand-specific 3'-5' exonuclease activity, which are provided by MRE11, to initiate end resection, which is required for single-strand invasion and recombination (PubMed:19759395, PubMed:28867292, PubMed:9705271). Within the MRN complex, NBN acts as a protein-protein adapter, which specifically recognizes and binds phosphorylated proteins, promoting their recruitment to DNA damage sites (PubMed: 12419185, PubMed: 15616588, PubMed: 18411307, PubMed: 18582474, PubMed: 18583988, PubMed: 18678890, PubMed: 19759395, PubMed: 19804756, PubMed: 23762398, PubMed: <u>24534091</u>, PubMed: <u>27814491</u>, PubMed: <u>27889449</u>, PubMed: <u>33836577</u>). Recruits MRE11 and RAD50 components of the MRN complex to DSBs in response to DNA damage (PubMed:12419185, PubMed:18411307, PubMed:18583988, PubMed:18678890, PubMed: 24534091, PubMed: 26438602). Promotes the recruitment of PI3/PI4-kinase family members ATM, ATR, and probably DNA-PKcs to the DNA damage sites, activating their functions (PubMed: 15064416, PubMed: 15616588, PubMed: 15790808, PubMed: 16622404, PubMed: <u>22464731</u>, PubMed: <u>30952868</u>, PubMed: <u>35076389</u>). Mediates the recruitment of phosphorylated RBBP8/CtIP to DSBs, leading to cooperation between the MRN complex and RBBP8/CtIP to initiate end resection (PubMed:19759395, PubMed:27814491, PubMed:27889449, PubMed: 33836577). RBBP8/CtIP specifically promotes the endonuclease activity of the MRN complex to clear DNA ends containing protein adducts (PubMed: 27814491, PubMed: 27889449, PubMed: 30787182, PubMed: 33836577). The MRN complex is also required for the processing of R-loops (PubMed:31537797). NBN also functions in telomere length maintenance via its interaction with TERF2: interaction with TERF2 during G1 phase preventing recruitment of DCLRE1B/Apollo to telomeres (PubMed: 10888888, PubMed: 28216226). NBN also promotes DNA repair choice at dysfunctional telomeres: NBN phosphorylation by CDK2 promotes non-homologous end joining repair at telomeres, while unphosphorylated NBN promotes microhomology-mediated end-joining (MMEJ) repair (PubMed: 28216226). Enhances AKT1 phosphorylation possibly by association with the mTORC2 complex (PubMed: 23762398).

Cellular Location

Nucleus. Chromosome. Nucleus, PML body. Chromosome, telomere Note=Localizes to discrete nuclear foci after treatment with genotoxic agents (PubMed:10783165, PubMed:26215093, PubMed:26438602). Localizes to DNA double-strand breaks (DSBs); recruited to DNA damage sites via association with phosphorylated proteins, such as phosphorylated H2AX, phosphorylated MDC1 and phosphorylated RAD17 (PubMed:12419185, PubMed:18411307, PubMed:18582474, PubMed:18583988, PubMed:18678890, PubMed:19338747, PubMed:23115235, PubMed:24534091, PubMed:26438602) Acetylation of 'Lys-5' of histone H2AX (H2AXK5ac) promotes NBN/NBS1 assembly at the sites of DNA damage (PubMed:26438602)

Tissue Location

Ubiquitous (PubMed:9590180). Expressed at high levels in testis (PubMed:9590180).

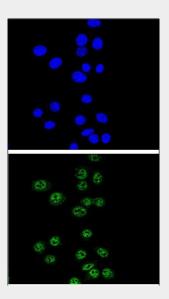
NBN Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

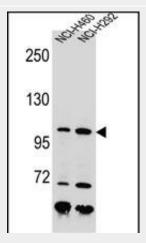


- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

NBN Antibody (C-term) - Images

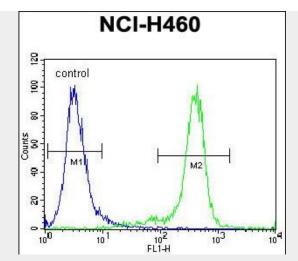


Confocal immunofluorescent analysis of NBN Antibody (C-term) (Cat. #AP11776b) with Hela cell followed by Alexa Fluor® 488-conjugated goat anti-rabbit IgG (green). DAPI was used to stain the cell nuclear (blue).



NBN Antibody (C-term) (Cat. #AP11776b) western blot analysis in NCI-H460,NCI-H292 cell line lysates (35ug/lane). This demonstrates the NBN antibody detected the NBN protein (arrow).





NBN Antibody (C-term) (Cat. #AP11776b) flow cytometric analysis of NCI-H460 cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

NBN Antibody (C-term) - Background

Mutations in this gene are associated with Nijmegen breakage syndrome, an autosomal recessive chromosomal instability syndrome characterized by microcephaly, growth retardation, immunodeficiency, and cancer predisposition. The encoded protein is a member of the MRE11/RAD50 double-strand break repair complex which consists of 5 proteins. This gene product is thought to be involved in DNA double-strand break repair and DNA damage-induced checkpoint activation.

NBN Antibody (C-term) - References

Liu, Y., et al. Carcinogenesis 31(10):1762-1769(2010) Kavitha, C.V., et al. Biochem. Biophys. Res. Commun. 399(4):575-580(2010) Ho-Pun-Cheung, A., et al. Pharmacogenomics J. (2010) In press: Jelonek, K., et al. J. Appl. Genet. 51(3):343-352(2010) Jugessur, A., et al. PLoS ONE 5 (7), E11493 (2010):