

**NBN Antibody (monoclonal) (M01)****Mouse monoclonal antibody raised against a partial recombinant NBN.****Catalog # AT2976a****Specification**

---

**NBN Antibody (monoclonal) (M01) - Product Information**

Application	WB, E
Primary Accession	<a href="#">O60934</a>
Other Accession	<a href="#">NM_002485</a>
Reactivity	Human
Host	mouse
Clonality	Monoclonal
Isotype	IgG2a Kappa
Calculated MW	84959

**NBN Antibody (monoclonal) (M01) - Additional Information****Gene ID** 4683**Other Names**

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

**Target/Specificity**

NBN (NP\_002476, 645 a.a. ~ 754 a.a) partial recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

**Dilution**

WB~~1:500~1000

E~~N/A

**Format**

Clear, colorless solution in phosphate buffered saline, pH 7.2 .

**Storage**

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

**Precautions**

NBN Antibody (monoclonal) (M01) is for research use only and not for use in diagnostic or therapeutic procedures.

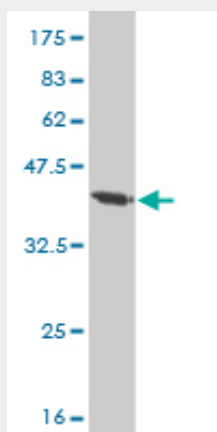
**NBN Antibody (monoclonal) (M01) - Protocols**

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

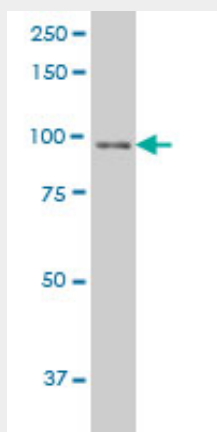
- [Western Blot](#)
- [Blocking Peptides](#)

- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

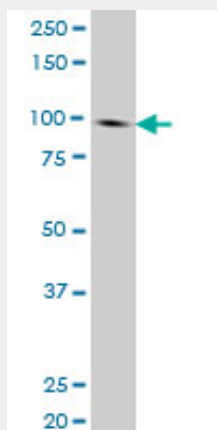
### NBN Antibody (monoclonal) (M01) - Images



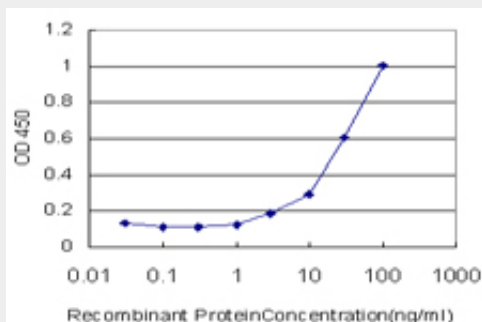
Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (37.84 kDa) .



NBN monoclonal antibody (M01), clone 3E4 Western Blot analysis of NBN expression in COLO 320 HSR ( (Cat # AT2976a )



NBN monoclonal antibody (M01), clone 3E4 Western Blot analysis of NBN expression in HL-60 (Cat # AT2976a )



Detection limit for recombinant GST tagged NBN is approximately 3ng/ml as a capture antibody.

### **NBN Antibody (monoclonal) (M01) - 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 (monoclonal) (M01) - References**

Association between single-nucleotide polymorphisms of selected genes involved in the response to DNA damage and risk of colon, head and neck, and breast cancers in a Polish population. Jelonek K, et al. J Appl Genet, 2010. PMID 20720310. A large-scale candidate gene approach identifies SNPs in SOD2 and IL13 as predictive markers of response to preoperative chemoradiation in rectal cancer. Ho-Pun-Cheung A, et al. Pharmacogenomics J, 2010 Jul 20. PMID 20644561. Maternal genes and facial clefts in offspring: a comprehensive search for genetic associations in two population-based cleft studies from Scandinavia. Jugessur A, et al. PLoS One, 2010 Jul 9. PMID 20634891. Gamma-Radiation Sensitivity and Polymorphisms in RAD51L1 Modulate Glioma Risk. Liu Y, et al. Carcinogenesis, 2010 Jul 7. PMID 20610542. The NBS1 genetic polymorphisms and the risk of the systemic lupus erythematosus in Taiwanese patients. Lin YJ, et al. J Clin Immunol, 2010 Sep. PMID 20571895.