Driving Innovation in Antibody Development

The company, mAbProtein, has been established to help patients, doctors, and researchers.



Human Nucleophosmin (NPM1) monoclonal antibody

Category monoclonal antibody

Catalog No. R-N-001

Applications WB, IHC, IF, IP

Reactivity Human, Dog No cross reaction: Mouse, Chicken

Immunogen information

Immunogen Recombinant protein of human NPM1 (P06748-1)

UniProt ID P06748

Synonyms B23, NP38, Numatrin

Gene ID 4869

Product information

Source Mouse **Clone No.** 9-2.6

Epitope This clone recognized NPM (114–219), a central region of NPM1, as well as

full length NPM1 (P06748-1), NPM1.3 (P06748-3) and NPM1(mut-A).

Isotype lgG1

Purification method DEAE ion-exchange purification

Lot No. 001

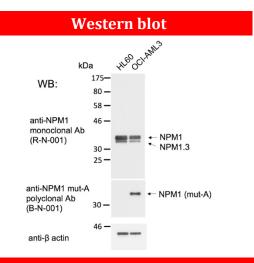
Concentration 1.0 mg/mL

Buffer 50% glycerol/PBS, pH7.4, w/o sodium azide

Storage Store at -20°C.

Recommended dilutions

WB 1:1000 – 1:2000
IHC 1:50 – 1:200
IF 1:50 – 1:200
IP 1:20 – 1:50



Detection of Endogenous NPM1 in Human leukemia Cell Lines by Western Blot.

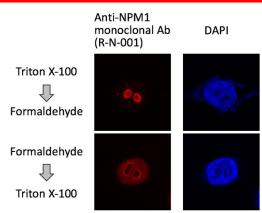
Endogenous NPM1 in HL-60 and OCI-AML3 leukemia cells were detected by Western blotting with mouse antihuman NPM1 monoclonal antibody (Catalog # R-N-001, upper panel).

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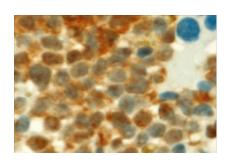
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Immunofluorescence



Immunohistochemistry



Background

Nucleophosmin was originally identified as a highly phosphorylated protein in the nucleolus. NPM1 also localizes to the nucleoplasm, and shuttles between the nucleus and the cytoplasm. NPM1 is involved in various biological processes such as ribosome biogenesis, centrosome duplication, genome instability and apoptosis. The overexpression of NPM1 has been observed in many types of solid tumors, including gastric, prostate, liver and colon. Translocation in the NPM1 gene has been reported for several hematopoietic malignancies; for example, t(2;5)(p23;q35) in 75% of anaplastic lymphoma kinase-positive anaplastic large cell lymphoma. Furthermore, approximately one-third of acute myeloid leukemia (AML) patients harbor frameshift mutations in exon 12 of the NPM1 gene, resulting in the generation of a nuclear export signal in the C-terminal region of NPM1 and localization of the mutant NPM1 to the cytoplasm. NPM1 type A mutation (NPM1 mut-A; TCTG insertion at the 960th nucleotide in exon 12) may be a good molecular marker for assessing the clinical status and predicting the outcomes in AML patients.

References for human NPM1 monoclonal antibody (R-N-001)

rarely occurs in myelodysplastic syndromes.

| PMID: | 29507312 | Journal: | Scientific Reports |
|--------------|---|------------|---------------------|
| Application: | WB, IF, IP | IF (2020): | 4.380 |
| Title: | Analysis of the oligomeric states of nucleophosmin using size exclusion chromatography. | | |
| PMID: | 19052985 | Journal: | Leukemia & Lymphoma |
| Application: | WB, IHC | IF (2020): | 3.280 |
| Title: | Abnormal cytoplasmic dyslocalisation and/or reduction of nucleophosmin protein level | | |