

NUP98 mouse mAb

Catalog No :	YM1318
Reactivity :	Human
Applications :	WB;IP
Target :	NUP98
Fields :	>>Nucleocytoplasmic transport;>>Amyotrophic lateral sclerosis;>>Influenza A
Gene Name :	nup98
Human Gene Id :	4928
Human Swiss Prot No :	P52948
Immunogen :	Purified recombinant human NUP98 protein fragments expressed in E.coli.
Specificity :	This antibody detects endogenous levels of NUP98 and does not cross-react with related proteins.
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Monoclonal, Mouse
Dilution :	wb 1:1000
Purification :	The antibody was affinity-purified from mouse ascites by affinity-chromatography using epitope-specific immunogen.
Concentration :	1 mg/ml
Storage Stability :	-15°C to -25°C/1 year(Do not lower than -25°C)
Observed Band :	98kD
Background :	Nuclear pore complexes (NPCs) regulate the transport of macromolecules between the nucleus and cytoplasm, and are composed of many polypeptide

subunits, many of which belong to the nucleoporin family. This gene belongs to the nucleoporin gene family and encodes a 186 kDa precursor protein that undergoes autoproteolytic cleavage to generate a 98 kDa nucleoporin and 96 kDa nucleoporin. The 98 kDa nucleoporin contains a Gly-Leu-Phe-Gly (GLGF) repeat domain and participates in many cellular processes, including nuclear import, nuclear export, mitotic progression, and regulation of gene expression. The 96 kDa nucleoporin is a scaffold component of the NPC. Proteolytic cleavage is important for targeting of the proteins to the NPC. Translocations between this gene and many other partner genes have been observed in different leukemias. Rearrangements typically result in chimeras with the N-terminal GLGF domain of

Function :

disease:A chromosomal aberration involving NUP98 is associated with pediatric acute myeloid leukemia (AML) with intermediate characteristics between M2-M3 French-American-British (FAB) subtypes. Translocation t(9;11)(p22;p15) with PSIP1/LEDGF. The chimeric transcript is an in-frame fusion of NUP98 exon 8 to PSIP1/LEDGF exon 4.,disease:A chromosomal aberration involving NUP98 is found in a form of acute myeloid leukemia. Translocation t(7;11)(p15;p15) with HOXA9. Translocation t(11;17)(p15;p13) with PHF23.,disease:A chromosomal aberration involving NUP98 is found in a form of T-cell acute lymphoblastic leukemia (T-ALL). Translocation t(3;11)(q12.2;p15.4) with LNP1.,disease:A chromosomal aberration involving NUP98 is found in a form of therapy-related myelodysplastic syndrome. Translocation t(11;20)(p15;q11) with TOP1.,disease:A chromosomal aberration involving NUP98 is found in childhood

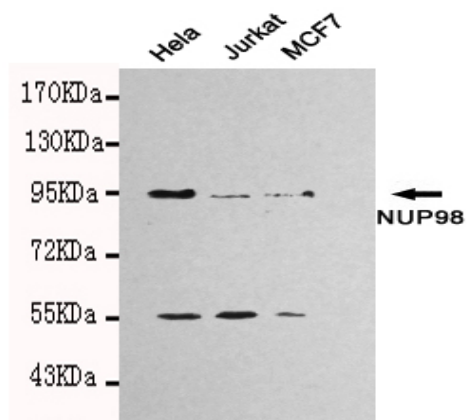
Subcellular Location :

Nucleus membrane ; Peripheral membrane protein; Nucleoplasmic side . Nucleus, nuclear pore complex . Nucleus, nucleoplasm . Localized to the nucleoplasmic side of the nuclear pore complex (NPC), at or near the nucleoplasmic basket (PubMed:11839768). Dissociates from the disassembled NPC structure early during prophase of mitosis (PubMed:12802065). Colocalized with NUP153 and TPR to the nuclear basket of NPC (PubMed:11839768). Colocalized with DHX9 in diffuse and discrete intranuclear foci (GLFG-body) (PubMed:11839768, PubMed:28221134). . ; Nucleus membrane . (Microbial infection) Remains localized to the nuclear membrane after poliovirus (PV) infection. .

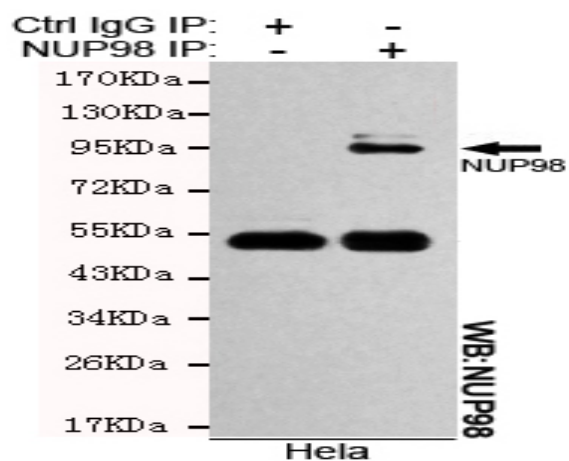
Expression :

Brain,Epithelium,Liver,Lung,Peripheral blood,Testis,

Products Images



Western blot detection of NUP98 in HeLa, Jurkat and MCF7 cell lysates using NUP98 mouse mAb (1:1000 diluted). Predicted band size: 98kDa. Observed band size: 98kDa.



Immunoprecipitation analysis of HeLa cell lysates using NUP98 mouse mAb.