



Neurofibromin Monoclonal Antibody

Catalog No	YP-mAb-00460
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	NF1
Immunogen	The antiserum was produced against synthesized peptide derived from human NF1. AA range:1551-1600
Specificity	Neurofibromin Monoclonal Antibody detects endogenous levels of Neurofibromin protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	Neurofibromin;NF1;NF 1;NF-1;Neurofibromatosis-related protein NF-1
Observed Band	319kD
Calculated Molecular Weight	319kD
Cell Pathway	Nucleus . Nucleus, nucleolus .
Tissue Specificity	Detected in brain, peripheral nerve, lung, colon and muscle.
Function	alternative products:Experimental confirmation may be lacking for some isoforms,caution:Was originally (PubMed:8807336) thought to be associated with LEOPARD (LS), an autosomal dominant syndrome.,disease:Defects in NF1 are a cause of familial spinal neurofibromatosis (spinal NF) [MIM:162210]. Familial spinal NF is considered to be an alternative form of neurofibromatosis, showing multiple spinal tumors.,disease:Defects in NF1 are a cause of juvenile myelomonocytic leukemia (JMML) [MIM:607785]. JMML is a pediatric myelodysplastic syndrome that constitutes approximately 30% of childhood cases of myelodysplastic syndrome (MDS) and 2% of leukemia. Germline mutations of NF1 account for the association of JMML with type 1 neurofibromatosis (NF1).,disease:Defects in NF1 are a cause of



neurofibromatosis-Noonan syndrome (NFNS) [MIM:601321]. NFNS is characterized by manifestations of both NF1 and

Background

This gene product appears to function as a negative regulator of the ras signal transduction pathway. Mutations in this gene have been linked to neurofibromatosis type 1, juvenile myelomonocytic leukemia and Watson syndrome. The mRNA for this gene is subject to RNA editing (CGA>UGA->Arg1306Term) resulting in premature translation termination. Alternatively spliced transcript variants encoding different isoforms have also been described for this gene. [provided by RefSeq, Jul 2008],

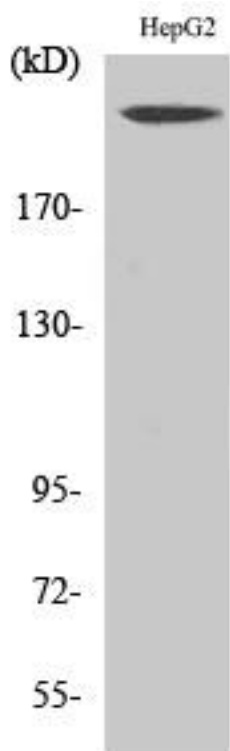
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using Neurofibromin Monoclonal Antibody