

SMN2 monoclonal antibody (M02), clone 1A3-2B9

Catalog # H00006607-M02 Size 100 ug

Applications

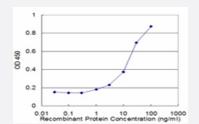


Western Blot (Transfected lysate)

Western Blot analysis of SMN2 expression in transfected 293T cell line by SMN2 monoclonal antibody (M02), clone 1A3-2B9.

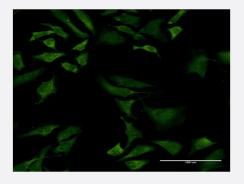
Lane 1: SMN2 transfected lysate (Predicted MW: 31.8 KDa).

Lane 2: Non-transfected lysate.



Sandwich ELISA (Recombinant protein)

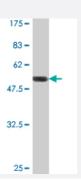
Detection limit for recombinant GST tagged SMN2 is approximately 1ng/ml as a capture antibody.



Immunofluorescence

Immunofluorescence of monoclonal antibody to SMN2 on HeLa cell . [antibody concentration 10 ug/ml]





Western Blot detection against Immunogen (56.76 KDa).

Specification	
Product Description	Mouse monoclonal antibody raised against a full-length recombinant SMN2.
Immunogen	SMN2 (AAH00908, 1 a.a. ~ 282 a.a) full-length recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
Sequence	MAMSSGGSGGVPEQEDSVLFRRGTGQSDDSDIWDDTALIKAYDKAVASFKHALKNGDICETSG KPKTTPKRKPAKKNKSQKKNTAASLQQWKVGDKCSAIWSEDGCIYPATIASIDFKRETCVVVYTG YGNREEQNLSDLLSPICEVANNIEQNAQENENESQVSTDESENSRSPGNKSDNIKPKSAPWNSFL PPPPPMPGPRLGPGKPGLKFNGPPPPPPPPPPPPHLLSCWLPPFPSGPPIIPPPPPICPDSLDDAD ALGSMLISWYMSGYHTGYYMEMLA
Host	Mouse
Reactivity	Human
Isotype	lgG2a Kappa
Quality Control Testing	Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (56.76 KDa).
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

Western Blot (Transfected lysate)

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Lane 2: Non-transfected lysate.

Protocol Download



Western Blot (Recombinant protein)

Protocol Download

Sandwich ELISA (Recombinant protein)

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Protocol Download

- ELISA
- Immunofluorescence

Immunofluorescence of monoclonal antibody to SMN2 on HeLa cell . [antibody concentration 10 ug/ml]

Gene Info — SMN2	
Entrez GeneID	6607
GeneBank Accession#	BC000908
Protein Accession#	<u>AAH00908</u>
Gene Name	SMN2
Gene Alias	BCD541, C-BCD541, FLJ76644, MGC20996, MGC5208, SMNC
Gene Description	survival of motor neuron 2, centromeric
Omim ID	601627
Gene Ontology	<u>Hyperlink</u>



Product Information

Gene Summary

This gene is part of a 500 kb inverted duplication on chromosome 5q13. This duplicated region c ontains at least four genes and repetitive elements which make it prone to rearrangements and d eletions. The repetitiveness and complexity of the sequence have also caused difficulty in determi ning the organization of this genomic region. The telomeric and centromeric copies of this gene a re nearly identical and encode the same protein. While mutations in the telomeric copy are associ ated with spinal muscular atrophy, mutations in this gene, the centromeric copy, do not lead to dis ease. This gene may be a modifier of disease caused by mutation in the telomeric copy. The criti cal sequence difference between the two genes is a single nucleotide in exon 7, which is thought t o be an exon splice enhancer. Note that the nine exons of both the telomeric and centromeric copi es are designated historically as exon 1, 2a, 2b, and 3-8. It is thought that gene conversion events may involve the two genes, leading to varying copy numbers of each gene. The full length protein e ncoded by this gene localizes to both the cytoplasm and the nucleus. Within the nucleus, the protei n localizes to subnuclear bodies called gems which are found near coiled bodies containing high concentrations of small ribonucleoproteins (snRNPs). This protein forms heteromeric complexes with proteins such as SIP1 and GEMIN4, and also interacts with several proteins known to be invo lved in the biogenesis of snRNPs, such as hnRNP U protein and the small nucleolar RNA binding protein. Four transcript variants encoding distinct isoforms have been described. [provided by Ref Seq

Other Designations

OTTHUMP00000125236|OTTHUMP00000125237|gemin 1

Disease

- Amyotrophic lateral sclerosis
- Disease Progression
- Genetic Predisposition to Disease
- Muscular Atrophy
- Nerve Degeneration
- Spinal Muscular Atrophies of Childhood
- Spinal muscular atrophy