

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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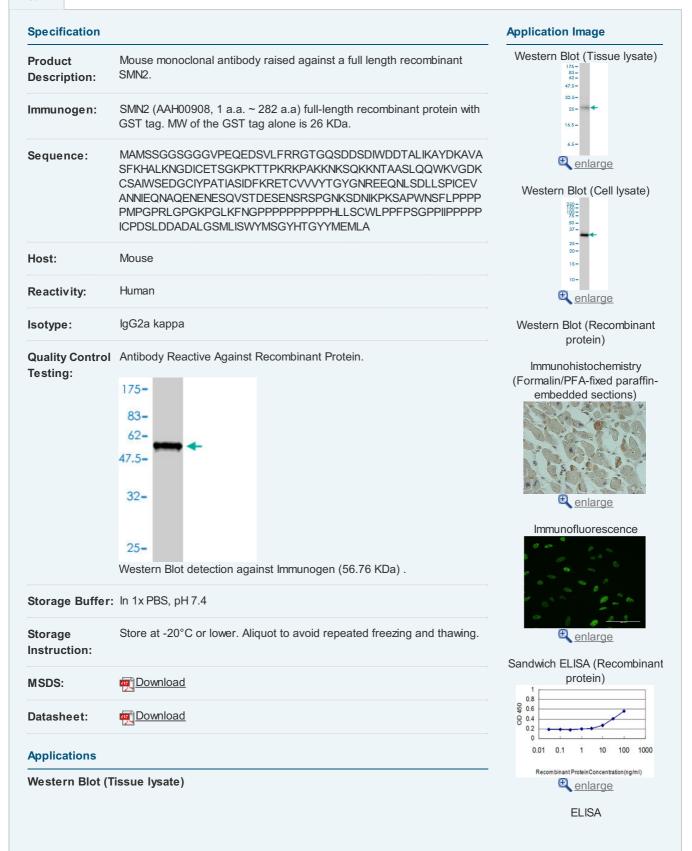




SMN2 monoclonal antibody (M01), clone 2B11-2A9

Catalog # : H00006607-M01 規格 : [100 ug]

List All



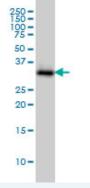
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SMN2 monoclonal antibody (M01), clone 2B11-2A9. Western Blot analysis of SMN2 expression in human colon.



Western Blot (Cell lysate)



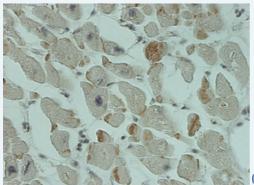
SMN2 monoclonal antibody (M01), clone 2B11-2A9 Western Blot analysis of SMN2 expression in IMR-32 (Cat # L008V1).



Western Blot (Recombinant protein)



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)



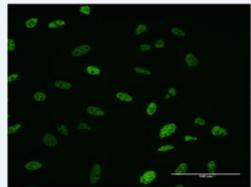
enlarge this image

Immunoperoxidase of monoclonal antibody to SMN2 on formalin-fixed paraffin-embedded human heart tissue. [antibody concentration $1 \sim 10 \text{ ug/ml}$]

Protocol Download

Immunofluorescence

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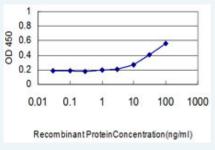


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Immunofluorescence of monoclonal antibody to SMN2 on HeLa cell. [antibody concentration 10 ug/ml]

Protocol Download

Sandwich ELISA (Recombinant protein)



Detection limit for recombinant GST tagged SMN2 is approximately 0.3ng/ml as a capture antibody.



ELISA

Gene Information

Entrez GenelD: 6607

GeneBank BC000908

Accession#:

Protein AAH00908

Accession#:

Gene Name: SMN2

BCD541,C-BCD541,FLJ76644,MGC20996,MGC5208,SMNC Gene Alias:

Gene

survival of motor neuron 2, centromeric

Description:

Omim ID: 601627

Gene Ontology: Hyperlink

Gene Summary: This gene is part of a 500 kb inverted duplication on chromosome 5q13. This duplicated region contains at least four genes and repetitive elements which make it prone to rearrangements and deletions. The repetitiveness and complexity of the sequence have also caused difficulty in determining the organization of this genomic region. The telomeric and centromeric copies of this gene are nearly identical and encode the same protein. While mutations in the telomeric copy are associated with spinal muscular atrophy, mutations in this gene, the centromeric copy, do not lead to disease. This gene may be a modifier of disease caused by mutation in the telomeric copy. The critical sequence difference between the two genes is a single nucleotide in exon 7, which is thought to be an exon splice enhancer. Note that the nine exons of both the telomeric and centromeric copies are designated

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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 encoded by this gene localizes to both the cytoplasm and the nucleus. Within the nucleus, the protein 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 involved 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 RefSeq

Other Designations:

OTTHUMP00000125236,OTTHUMP00000125237,gemin 1

Related Disease

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

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