

Allgemeine Informationen

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| Katalog-Nr.: CL594-60255 | GenBank-Zugangsnummer: BC062723 | Reinigungsmethode: Protein-G-Reinigung |
| Größe: 100ul , Konzentration: 1000 µg/ml von6606 Nanodrop; | GeneID (NCBI): von6606 | CloneNo.: 3A8G11 |
| Wirt: Maus | Vollständiger Name: survival of motor neuron 1, telomeric | Empfohlene Verdünnungen: IF 1:50-1:500 |
| Isotyp: IgG1 | Berechnete Masse: 294 aa, 32 kDa | Anregungs-/Emissionsmaxima- Wellenlängen: 588 nm / 604 nm |
| Immunogen Katalognummer: AG16615 | Beobachtete Masse: 40 kDa | |

Anwendungen

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| Geprüfte Anwendungen: IF | Positivkontrollen: IF : HepG2-Zellen, |
| Getestete Reaktivität: Human, Maus | |

Hintergrundinformationen

Spinal muscular atrophy (SMA) is an autosomal recessive neurodegenerative disease characterized by loss of anterior horn cells in the spinal cord and concomitant symmetrical muscle weakness and atrophy (PMID: 16364894). SMA is caused by deletion or mutations of the survival motor neuron (SMN1) gene. SMA patients lack a functional SMN1 gene, but they possess an intact SMN2 gene, which though nearly identical to SMN1, is only partially functional (PMID: 17355180). A large majority of SMN2 transcripts lack exon 7, resulting in production of a truncated, less stable SMN protein (PMID: 10369862). The level of SMN protein correlates with phenotypic severity of SMA. This antibody, 60255-1-ig, raised against the C-terminal region (275-294aa) encoded by the exon 7.

Lagerung

Lagerungsbedingungen:
Bei -20°C lagern. Vor Licht schützen. Nach dem Versand ein Jahr stabil.

Lagerungspuffer:
BS mit 50% Glycerin, 0,05% Proclin300, 0,5% BSA, pH 7,3.

Aliquotieren ist nicht notwendig bei -20°C Lagerung

*** 20ul-Größen enthalten 0.1% BSA

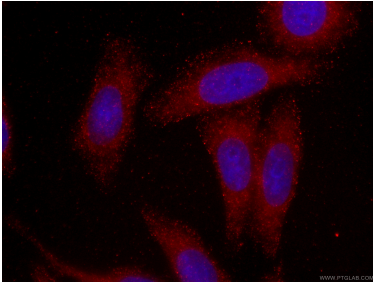
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Ausgewählte Validierungsdaten



Immunofluorescent analysis of (-20°C Ethanol) fixed HepG2 cells using CL594-60255 (SMN-Exon7 antibody) at dilution of 1:100.