For Research Use Only

CoraLite®594-conjugated SMN-Exon7 Monoclonal antibody

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Catalog Number: CL594-60255

Catalog Number: GenBank Accession Number: **Purification Method: Basic Information** CL594-60255 BC062723 Protein G purification

> GeneID (NCBI): CloneNo.: 100ul, Concentration: 1000 ug/ml by 6606 3A8G11

Nanodrop: **UNIPROT ID:** Recommended Dilutions: Q16637 IF/ICC 1:50-1:500

Mouse Full Name: Excitation/Emission maxima

Isotype: survival of motor neuron 1, telomeric wavelengths: 588 nm / 604 nm lgG1 Calculated MW:

Immunogen Catalog Number: 294 aa, 32 kDa AG16615 Observed MW: 40 kDa

Applications

Positive Controls: **Tested Applications:** IF/ICC

IF/ICC: HepG2 cells,

Species Specificity: human, mouse

Background Information

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-lg, raised against the C-terminal region (275-294aa) encoded by the exon 7.

Storage

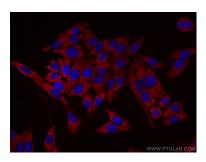
Store at -20°C. Avoid exposure to light. Stable for one year after shipment.

PBS with 50% Glycerol, 0.05% Proclin300, 0.5% BSA, pH 7.3.

Aliquoting is unnecessary for -20°C storage

in USA), or 1(312) 455-8498 (outside USA)

Selected Validation Data



Immunofluorescent analysis of (-20°C Ethanol) fixed HepG2 cells using CoraLite®594 SMN-Exon7 antibody (CL594-60255, Clone: 3A8G11) at dilution of 1:200.