For Research Use Only

CoraLite®594-conjugated SMN-Exon7 Monoclonal antibody



Purification Method:

Catalog Number: CL594-60255

Basic Information

Catalog Number: GenBank Accession Number: CL594-60255 BC062723

Protein G purification GeneID (NCBI): CloneNo.:

Size: 1000 µg/ml 6606 3A8G11 **UNIPROT ID:** Recommended Dilutions: Source:

Mouse Q16637 IF 1:50-1:500

Full Name: Excitation/Emission maxima Isotype:

lgG1 survival of motor neuron 1, telomeric wavelengths:

588 nm / 604 nm Calculated MW: Immunogen Catalog Number:

AG16615 294 aa, 32 kDa

Observed MW: 40 kDa

Applications

Tested Applications:

IF/ICC

Species Specificity: human, mouse

Positive Controls:

IF: HepG2 cells,

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.

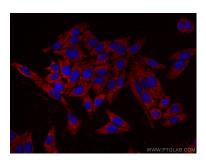
Storage

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

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.