For Research Use Only

CoraLite® Plus 488-conjugated SMN-Exon7 Monoclonal antibody



Catalog Number: CL488-60255

Basic Information

Catalog Number: GenBank Accession Number: Purification Method: Protein G purification

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

Nanodrop; Full Name: Recommended Dilutions:

Source: survival of motor neuron 1, telomeric IF 1:50-1:500

Mouse Calculated MW: Excitation/Emission maxima

 Isotype:
 294 aa, 32 kDa
 wavelengths:

 IgG1
 Observed MW:
 488 nm / 515 nm

Immunogen Catalog Number: 40 kDa

AG16615

Applications Tested Applications:

IF

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

Storage:

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

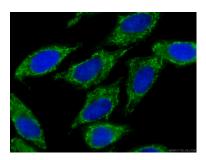
Storage Buffer

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

Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

Selected Validation Data



Immunofluorescent analysis of (-20°C Ethanol) fixed HepG2 cells using CL488-60255 (SMN-Exon7 antibody) at dilution of 1:100.