

À des fins de recherche uniquement

Anticorps Polyclonal de lapin anti-SMN



Numéro de catalogue: 11708-1-AP

Phare

11 Publications

Informations de base

Numéro de catalogue:

11708-1-AP

Taille:

150ul, Concentration: 500 µg/ml by Nanodrop;

Hôte:

Lapin

Isotype:

IgG

Immunogen Catalog Number:

AG2260

Numéro d'acquisition GenBank:

BC000908

Identification du gène (NCBI):

6607

Nom complet:

survival of motor neuron 2, centromeric

MW calculé

282 aa, 30 kDa

MW observés:

38 kDa, 66-70 kDa

Méthode de purification:

Purification par affinité contre l'antigène

Dilutions recommandées:

WB 1:1000-1:6000

IP 0.5-4.0 µg for IP and 1:500-1:2000 for WB

IHC 1:50-1:200

IF 1:400-1:1600

Applications

Applications testées:

IF, IHC, IP, WB, ELISA

Demandes citées:

ELISA, IF, IP, WB

Spécificité de l'espèce:

Humain, rat, souris

Espèces citées:

Humain, rat, souris

Remarque-IHC: il est suggéré de démasquer l'antigène avec un tampon de TE buffer pH 9,0; (*) À défaut, le démasquage de l'antigène peut être effectué avec un tampon citrate pH 6,0.

Contrôles positifs:

WB : cellules HEK-293, cellules HeLa, cellules HepG2, cellules Jurkat, cellules K-562, tissu testiculaire de souris

IP : cellules HEK-293,

IHC : tissu rénal humain, tissu cardiaque humain, tissu cérébral humain, tissu cutané humain, tissu ovarien humain, tissu placentaire humain, tissu pulmonaire humain, tissu splénique humain, tissu testiculaire humain

IF : cellules HepG2,

Informations générales

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, 11708-1-AP, raised against the recombinant full-length human SMN2 protein, recognizes all isoforms of SMN protein.

Publications notables

Autrice	Pubmed ID	Journal	Application
Vicki L McGovern	33084884	Hum Mol Genet	WB
Yuhong Zhang	34628513	J Mol Med (Berl)	WB,IF,IP
Phillip Zaworski	26953792	PLoS One	

Stockage

Stockage:

Stocker à -20°C. Stable pendant un an après l'expédition.

Tampon de stockage:

PBS avec azoture de sodium à 0,02 % et glycérol à 50 % pH 7,3

L'aliquote n'est pas nécessaire pour le stockage à -20C

*** Les 20ul contiennent 0,1% de BSA.

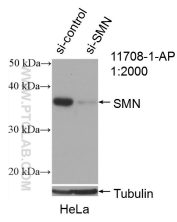
For technical support and original validation data for this product please contact:

T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA)

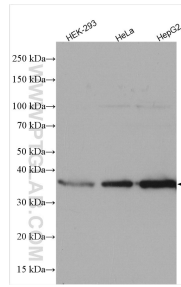
E: proteintech@ptglab.com
W: ptglab.com

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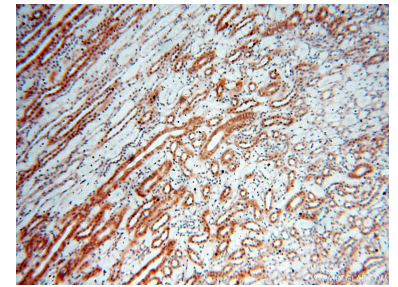
Données de validation sélectionnées



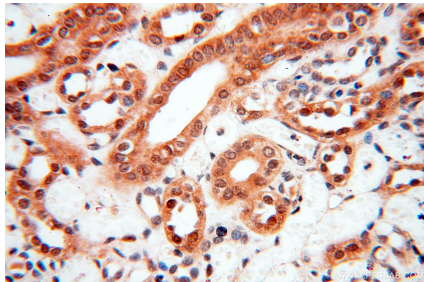
WB result of SMN antibody (11708-1-AP; 1:2000; incubated at room temperature for 1.5 hours) with sh-Control and sh-SMN transfected HeLa cells.



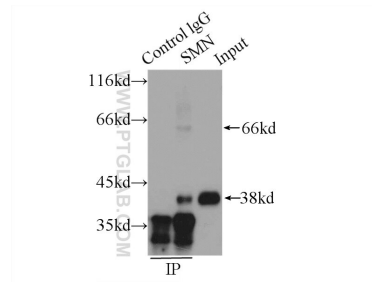
Various lysates were subjected to SDS PAGE followed by western blot with 11708-1-AP (SMN antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours.



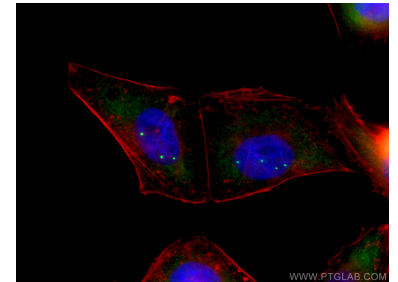
Immunohistochemical analysis of paraffin-embedded human kidney using 11708-1-AP (SMN antibody) at dilution of 1:100 (under 10x lens).



Immunohistochemical analysis of paraffin-embedded human kidney using 11708-1-AP (SMN antibody) at dilution of 1:100 (under 40x lens).



IP Result of anti-SMN (IP:11708-1-AP, 3ug; Detection:11708-1-AP 1:1000) with HEK-293 cells lysate 1000ug.



Immunofluorescent analysis of (4% PFA) fixed HepG2 cells using SMN antibody (11708-1-AP) at dilution of 1:800 and CoraLite@488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L), CL594-Phalloidin (red).