

À des fins de recherche uniquement

Anticorps Polyclonal de lapin anti-PSAP



Numéro de catalogue: 10801-1-AP

Phare

18 Publications

Informations de base

Numéro de catalogue:	BC001503	Méthode de purification:
10801-1-AP		Purification par affinité contre l'antigène
Taille:	5660	Dilutions recommandées:
150ul , Concentration: 700 µg/ml by Nanodrop and 350 µg/ml by Bradford method using BSA as the standard;	Nom complet: prosaposin	WB 1:1000-1:8000 IHC 1:50-1:500 IF 1:200-1:800
Hôte:	MW calculé	
Lapin	58 kDa	
Isotype:	MW observés:	
IgG	60 kDa	
Immunogen Catalog Number:		
AG1194		

Applications

Applications testées:	Contrôles positifs:
IF, IHC, WB, ELISA	WB : cellules A431, cellules HeLa, cellules HepG2, cellules MCF-7
Demandes citées:	IHC : tissu de cancer du sein humain, tissu de cervelet de souris, tissu splénique de souris
ELISA, IF, IHC, WB	IF : cellules HeLa, cellules HEK-293, cellules U-251
Spécificité de l'espèce:	
Humain	
Espèces citées:	
Humain, souris	
<i>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.</i>	

Informations générales

The PSAP gene encodes prosaposin, a precursor of four small nonenzymatic glycoproteins termed 'sphingolipid activator proteins' (SAPs) that assist in the lysosomal hydrolysis of sphingolipids. After proteolytic processing of the prosaposin protein, these 4 released polypeptides are functional activators. Saposin A is encoded by residues 60 to 143 of PSAP, saposin B by 195 to 275, saposin C by 311 to 390, and saposin D by 405 to 487. There are four 12-14 kDa heatstable glycoproteins. Saposins A-D localize primarily to the lysosomal compartment where they facilitate the catabolism of glycosphingolipids with short oligosaccharide groups. Saposins A-D are required for the hydrolysis of certain sphingolipids by specific lysosomal hydrolases. (PMID: 2001789) Defects in PSAP are the cause of Gaucher disease, Tay-Sachs disease, and metachromatic leukodystrophy (PubMed: 2060627, PMID: 15773042). This PSAP antibody (10801-1-AP) is expected to recognize both saposin A and B.

Publications notables

Autrice	Pubmed ID	Journal	Application
Costain Willard J WJ	20718007	Proteomics	IHC
Yachao He	36233357	Int J Mol Sci	WB, IF
Sebastian Boland	36207292	Nat Commun	WB

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'aliquotage n'est pas nécessaire pour le stockage à -20°C

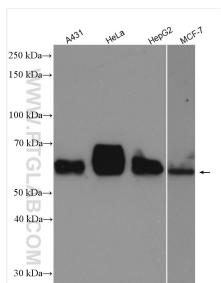
*** Les 20ul contiennent 0,1% de BSA.

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)

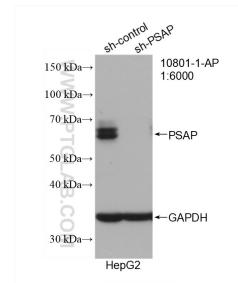
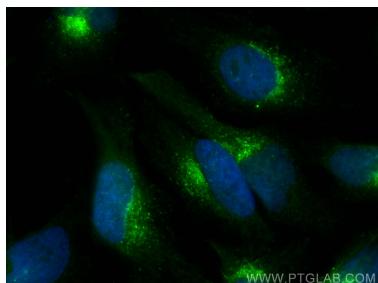
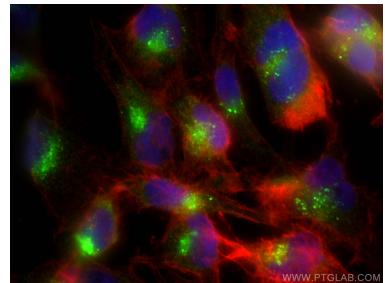
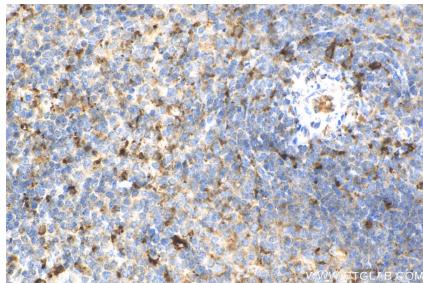
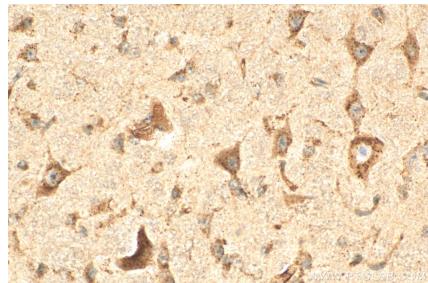
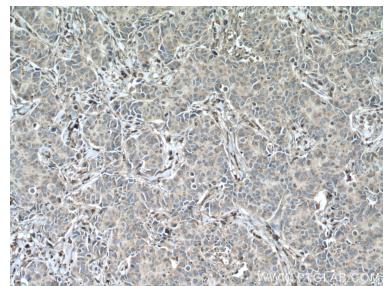
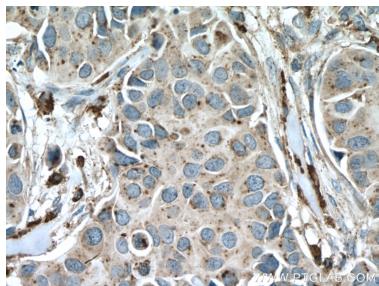
E: proteintech@ptglab.com
W: ptglab.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

Données de validation sélectionnées



Various lysates were subjected to SDS PAGE followed by western blot with 10801-1-AP (PSAP antibody) at dilution of 1:4000 incubated at room temperature for 1.5 hours.



WB result of PSAP antibody (10801-1-AP; 1:6000; incubated at room temperature for 1.5 hours) with sh-Control and sh-PSAP transfected HepG2 cells.