For Research Use Only HPSE Polyclonal antibody Catalog Number: 16673-1-AP 2 Publications

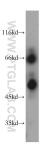


Basic Information	Catalog Number: 16673-1-AP	GenBank Accession Number: BC051321		rification Method: ntigen affinity purification
	Size:	GenelD (NCBI):		commended Dilutions:
	150ul , Concentration: 300 ug/ml by	10855	W	B 1:200-1:1000
	Nanodrop and 207 ug/ml by Bradford method using BSA as the standard;	onin Korrib.	IH	C 1:20-1:200
	Source:	Q9Y251		
	Rabbit	Full Name: heparanase		
	Isotype:	Calculated MW:		
	IgG	543 aa, 61 kDa		
	Immunogen Catalog Number: AG10167	Observed MW: 65 kDa, 50 kDa		
Applications	Tested Applications:	Pc	ositive Controls	:
	WB, IHC, ELISA	WB : DU 145 cells,		,
	Cited Applications: WB, IF		IHC : human liver cancer tissue, human placenta	
	Species Specificity:	tissue, mouse kidney tissue		
	human, mouse, rat			
	Cited Species: human, mouse			
	Note-IHC: suggested antigen r TE buffer pH 9.0; (*) Alternativ retrieval may be performed w buffer pH 6.0	vely, antigen		
	HPSE(Heparanase) is also named as HEP, HPA, HPA1, HPR1, HPSE1, HSE1 and belongs to the glycosyl hydrolase 79 family. It is a endoglycosidase that cleaves heparan sulfate proteoglycans (HSPGs) into heparan sulfate side chair and core proteoglycans. HPSE is essential in the disassembly of the extracellular matrix (ECM) by invading cells. I has 3 isoforms produced by alternative splicing with the molecular weight of 61 kDa, 55 kDa and 53 kDa. The full length protein has six glycosylation sites. The cleavage of the 65 kDa form leads to the generation of a linker peptide, and 8 kDa and 50 kDa products. The active form, the 8/50 kDa heterodimer, is resistant to degradation and glycosylation of the 50 kDa subunit appears to be essential for its solubility.			
Background Information	has 3 isoforms produced by alternativ length protein has six glycosylation s peptide, and 8 kDa and 50 kDa produc	ve splicing with the molect sites. The cleavage of the 6 cts. The active form, the 8/	55 kDa form lea 50 kDa heterod	1 kDa, 55 kDa and 53 kDa. The full ds to the generation of a linker
Background Information	has 3 isoforms produced by alternativ length protein has six glycosylation s peptide, and 8 kDa and 50 kDa produc glycosylation of the 50 kDa subunit a	ve splicing with the molect sites. The cleavage of the 6 cts. The active form, the 8/	55 kDa form lea 50 kDa heterod	1 kDa, 55 kDa and 53 kDa. The full ds to the generation of a linker
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Notable Publications	has 3 isoforms produced by alternative length protein has six glycosylation is peptide, and 8 kDa and 50 kDa produce glycosylation of the 50 kDa subunit a Author Pute Guang Xu 280 Altex M Agelidis 287 Storage: Storage: Storage Buffer: Storage Buffer:	ve splicing with the molect sites. The cleavage of the 6 cts. The active form, the 8/ ppears to be essential for i omed ID Journal 081450 Diabetes 700944 Cell Rep er shipment. % glycerol pH 7.3.	55 kDa form lea 50 kDa heterod its solubility.	1 kDa, 55 kDa and 53 kDa. The full ds to the generation of a linker imer, is resistant to degradation a Application WB

For technical support and original validation data for this product please contact: T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free E: proteintech@ptglab.com in USA), or 1(312) 455-8498 (outside USA) W: ptglab.com

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Selected Validation Data





DU 145 cells were subjected to SDS PAGE followed by western blot with 16673-1-AP (HPSE antibody) at dilution of 1:300 incubated at room temperature for 1.5 hours. Immunohistochemical analysis of paraffinembedded human liver cancer using 16673-1-AP (HPSE antibody) at dilution of 1:50 (under 10x lens). Immunohistochemical analysis of paraffinembedded human liver cancer using 16673-1-AP (HPSE antibody) at dilution of 1:50 (under 40x lens).