For Research Use Only

RNASEH1 Polyclonal antibody Catalog Number: 15606-1-AP Featured Product 51 Pub

Featured Product

51 Publications

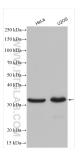


Basic Information	Catalog Number: 15606-1-AP	GenBank Accession Nun BC002973		cation Method: en affinity purification	
	Concentration: 500 ug/ml	GenelD (NCBI): 246243	Recon	Recommended Dilutions: WB 1:1000-1:5000 IP 0.5-4.0 ug for 1.0-3.0 mg of total protein lysate IF/ICC 1:50-1:500	
	Source: Rabbit	UNIPROT ID: 060930	IP 0.5 protei		
	Isotype: IgG	Full Name: ribonuclease H1	IF/IC0		
	Immunogen Catalog Number: AG7990	Calculated MW: 32 kDa			
		Observed MW: 32 kDa			
Applications	Tested Applications:	1	Positive Controls:		
	WB, IF/ICC, IP, ELISA	WB: HeLa cells, U2OS cells			
	Cited Applications: WB, IHC, IF, ChIP	I	IP : HeLa cells,		
	Species Specificity: human, mouse, rat	IF/ICC : HeLa cells,			
	Cited Species: human, mouse				
	Ribonucleases H are enzymes that cleave the RNA of RNA/DNA hybrids that form during replication and repair ar which could lead to DNA instability if they were not processed. Eukaryotic RNases H are larger and more compley than their prokaryotic counterparts [PMID:12667461]. Eukaryotic RNase H1 has acquired a Hybrid Binding Domain that confers processivity and affinity for the substrate. Because of its essential role and its similarities in structur and enzymatic mechanism to the RNase H of HIV reverse transcriptase, RNASEH1 has been implicated in mitochondrial DNA replication during mouse development [PMID:19228196].				
Background Information	than their prokaryotic counterpart that confers processivity and affir and enzymatic mechanism to the	ty if they were not processed s [PMID:12667461]. Eukaryo ity for the substrate. Because RNase H of HIV reverse trans	l. Eukaryotic RNases tic RNase H1 has acq e of its essential role scriptase, RNASEH1 h	H are larger and more comple uired a Hybrid Binding Doma and its similarities in structu	
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Background Information Notable Publications	than their prokaryotic counterpart that confers processivity and affir and enzymatic mechanism to the mitochondrial DNA replication du Author Negin Khosraviani	ty if they were not processed s [PMID:12667461]. Eukaryo ity for the substrate. Because RNase H of HIV reverse trans ring mouse development [PI Pubmed ID Journal 36178790 STAR P	I. Eukaryotic RNases tic RNase H1 has acq e of its essential role criptase, RNASEH1 h MID:19228196]. l rotoc mmun	H are larger and more comple uired a Hybrid Binding Doma and its similarities in structu as been implicated in Application WB	

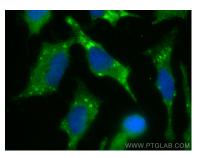
For technical support and original validation data for this product please contact: E: Proteintech-CN@ptglab.com T: 4006900926 W: ptgcn.com

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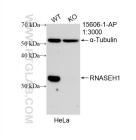
Selected Validation Data



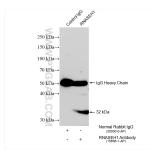
Various lysates were subjected to SDS PAGE followed by western blot with 15606-1-AP (RNASEH1 antibody) at dilution of 1:2500 incubated at room temperature for 1.5 hours.



Immunofluorescent analysis of (-20°C Methanol) fixed HeLa cells using RNASEH1 antibody (15606-1-AP) at dilution of 1:200 and CoraLite®488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).



WB result of RNASEH1 antibody (15606-1-AP; 1:3000; room temperature for 1.5 hours) with wildtype and RNASEH1 knockout HeLa cells.



IP result of anti-RNASEH1 (IP:15606-1-AP, 4ug; Detection:15606-1-AP 1:3000) with HeLa cells lysate 1750 ug.