For Research Use Only

XRN2 Monoclonal antibody

Catalog Number:66852-1-lg 2 Publications

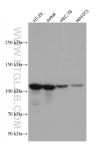


Basic Information	Catalog Number: 66852-1-lg	GenBank Accession Number: BC006417		Purification Method: Protein A purification	
	Size:			CloneNo.: 2C3E3 Recommended Dilutions: WB 1:1000-1:6000	
	150ul , Concentration: 1000 ug/ml by Nanodrop and 920 ug/ml by Bradford method using BSA as the standard;				
	Source: Mouse			IHC 1:150-1:600 IF/ICC 1:400-1:1	
	lsotype: lgG1	Calculated MW: 104 kDa			
	Immunogen Catalog Number: AG27927	Observed MW: 109 kDa			
Applications	Tested Applications:		Positive Controls:		
				9 cells, HEK-293 cells, COLO 320 cells, Jurka :-T6 cells, NIH/3T3 cells	
	WB IHC : human breast cancer tissu				e,
	Species Specificity: human, mouse, rat		IF/ICC : MCF-7 cells,		
	Cited Species: human				
	Note-IHC: suggested antigen ro TE buffer pH 9.0; (*) Alternativ retrieval may be performed w buffer pH 6.0	vely, antigen			
Background Information	XRN2 is one exonuclease that degrade II termination. During transcription te		ges at the poly	adenylation site	
	which is subsequently processed to for elongating polymerase. The processiv transcription.		0		
	elongating polymerase. The processiv transcription.		fragment by t		
<u> </u>	elongating polymerase. The processiv transcription. Author Pub	ve degradation of this 3' med ID Journa	fragment by t	his protein may p	romote termination of
	elongating polymerase. The processive transcription. Author Pub Wen-Long Xue 321	ve degradation of this 3' med ID Journa	fragment by t l nysiol Cell Ph	his protein may p	romote termination of Application
Notable Publications	elongating polymerase. The processive transcription. Author Pub Wen-Long Xue 321. Ruihui Xie 369. Storage: Storage Storage Buffer: PBS with 0.02% sodium azide and 50%	ve degradation of this 3' med ID Journa 86933 Am J Pl 139377 Cancer er shipment. % glycerol pH 7.3.	fragment by t l nysiol Cell Ph	his protein may p	romote termination of Application WB
Notable Publications Storage *** 20ul sizes contain 0.1% BSA	elongating polymerase. The processive transcription.	ve degradation of this 3' med ID Journa 86933 Am J Pl 139377 Cancer er shipment. % glycerol pH 7.3.	fragment by t l nysiol Cell Ph	his protein may p	romote termination of Application WB

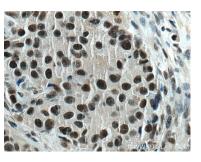
For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll freeE: proteintech@ptglab.comin USA), or 1(312) 455-8498 (outside USA)W: ptglab.com

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

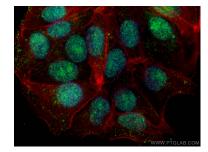
Selected Validation Data



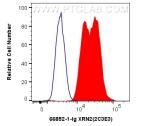
Various lysates were subjected to SDS PAGE followed by western blot with 66852-1-1g (XRN2 antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours.



Immunohistochemical analysis of paraffinembedded human breast cancer tissue slide using 66852-1-1g (XRN2 antibody) at dilution of 1:300 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunofluorescent analysis of (4% PFA) fixed MCF-7 cells using XRN2 antibody (66852-1-1g, Clone: 2C3E3) at dilution of 1:800 and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) (SA00013-1), CL594-phalloidin (red).



1X10^6 HepG2 cells were intracellularly stained with 0.4 ug Anti-Human XRN2 (66852-1-1g, Clone:2C3E3) and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000 (red), or 0.4 ug Mouse IgG1 Isotype Control (MOPC-21) (65124-1-1g, Clone: MOPC-21) (blue). Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).