For Research Use Only

XRN2 Monoclonal antibody

Catalog Number:66852-1-lg 2 Publications



Purification Method:

WB 1:1000-1:6000 IHC 1:150-1:600

IF 1:250-1:1000

Basic Information

Catalog Number: GenBank Accession Number:

66852-1-Ig BC006417 Protein G purification Size: GeneID (NCBI): CloneNo.:

150ul , Concentration: 1000 µg/ml by 22803 2C3E3
Nanodrop and 920 µg/ml by Bradford Full Name: Recommended Dilutions:

method using BSA as the standard; 5'-3' exoribonuclease 2

 Source:
 Calculated MW:

 Mouse
 104 kDa

 Isotype:
 Observed MW:

 IgG1
 109 kDa

Immunogen Catalog Number:

AG27927

Positive Controls:

IF, IHC, WB, ELISA WB: HT-29 cells, HEK-293 cells, COLO 320 cells, Jurkat

actions: cells, HSC-T6 cells, NIH/3T3 cells

IHC : human breast cancer tissue,

IF : HepG2 cells,

Applications

Cited Applications: WB

Tested Applications:

Species Specificity: Human, mouse, rat

Cited Species: human

Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0

Background Information

XRN2 is one exonuclease that degrades the Pol II associated product of poly(A) site cleavage, which is crucial for Pol II termination. During transcription termination, XRN2 cleavages at the polyadenylation site liberates a 5' fragment which is subsequently processed to form the mature mRNA and a 3' fragment which remains attached to the elongating polymerase. The processive degradation of this 3' fragment by this protein may promote termination of transcription.

Notable Publications

Author	Pubmed ID	Journal	Application
Wen-Long Xue	32186933	Am J Physiol Cell Physiol	WB
Ruihui Xie	36939377	Cancer Res	WB

Storage

Storage:

Store at -20°C. Stable for one year after shipment.

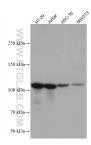
Storage Buffer:

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

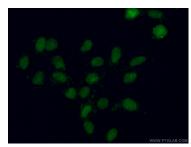
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

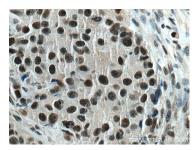
Selected Validation Data



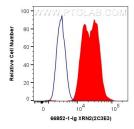
Various lysates were subjected to SDS PAGE followed by western blot with 66852-1-1g (XRN2 antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours.



Immunofluorescent analysis of (4% PFA) fixed HepG2 cells using 66852-1-Ig (XRN2 antibody) at dilution of 1:500 and CoraLite488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



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



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).