

Product datasheet

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ARG59016 anti-PARN antibody

Package: 50 μg Store at: -20°C

Summary

Product Description Rabbit Polyclonal antibody recognizes PARN

Tested Reactivity Hu, Ms, Rat

Tested Application FACS, ICC/IF, IHC-P, WB

Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name PARN

Species Human

Immunogen Recombinant protein corresponding to M1-Y301 of Human PARN.

Conjugation Un-conjugated

Alternate Names DAN; DKCB6; PFBMFT4; Poly(A)-specific ribonuclease PARN; EC 3.1.13.4; Deadenylating nuclease;

Deadenylation nuclease; Polyadenylate-specific ribonuclease

Application Instructions

Application table	Application	Dilution
	FACS	1:150 - 1:500
	ICC/IF	1:200 - 1:1000
	IHC-P	0.5 - 1 μg/ml
	WB	0.1 - 0.5 μg/ml
Application Note	IHC-P: Antigen Retrieval: Heat mediation was performed in Citrate buffer (pH 6.0) for 20 min. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form	Liquid	
Purification	Affinity purification with immunogen.	
Buffer	0.2% Na2HPO4, 0.9% NaCl, 0.05% Sodium azide and 4% Trehalose.	
Preservative	0.05% Sodium azide	
Stabilizer	4% Trehalose	
Concentration	0.5 mg/ml	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed	

For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol PARN

Gene Full Name poly(A)-specific ribonuclease

Background The protein encoded by this gene is a 3'-exoribonuclease, with similarity to the RNase D family of

3'-exonucleases. It prefers poly(A) as the substrate, hence, efficiently degrades poly(A) tails of mRNAs. Exonucleolytic degradation of the poly(A) tail is often the first step in the decay of eukaryotic mRNAs. This protein is also involved in silencing of certain maternal mRNAs during oocyte maturation and early embryonic development, as well as in nonsense-mediated decay (NMD) of mRNAs that contain premature stop codons. Alternatively spliced transcript variants encoding different isoforms have been

found for this gene. [provided by RefSeq, Aug 2008]

Function 3'-exoribonuclease that has a preference for poly(A) tails of mRNAs, thereby efficiently degrading

poly(A) tails. Exonucleolytic degradation of the poly(A) tail is often the first step in the decay of eukaryotic mRNAs and is also used to silence certain maternal mRNAs translationally during oocyte maturation and early embryonic development. Interacts with both the 3'-end poly(A) tail and the 5'-end cap structure during degradation, the interaction with the cap structure being required for an efficient degradation of poly(A) tails. Involved in nonsense-mediated mRNA decay, a critical process of selective degradation of mRNAs that contain premature stop codons. Also involved in degradation of inherently unstable mRNAs that contain AU-rich elements (AREs) in their 3'-UTR, possibly via its interaction with KHSRP. Probably mediates the removal of poly(A) tails of AREs mRNAs, which constitutes the first step

of destabilization. [UniProt]

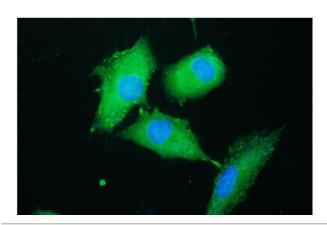
Calculated Mw 73 kDa

PTM Phosphorylation by MAPKAPK2, preventing GADD45A mRNA degradation after genotoxic stress.

[UniProt]

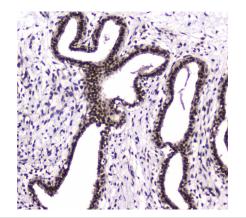
Cellular Localization Nucleus. [UniProt]

Images



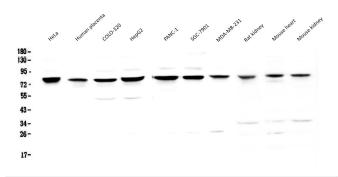
ARG59016 anti-PARN antibody ICC/IF image

Immunofluorescence: A549 cells were blocked with 10% goat serum and then stained with ARG59016 anti-PARN antibody (green) at 2 μ g/ml dilution, overnight at 4°C. DAPI (blue) for nuclear staining.



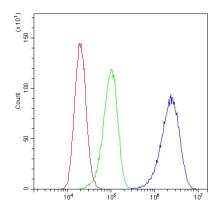
ARG59016 anti-PARN antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human mammary cancer tissue. Antigen Retrieval: Heat mediated was performed in Citrate buffer (pH 6.0, epitope retrieval solution) for 20 min. The tissue section was blocked with 10% goat serum. The tissue section was then stained with ARG59016 anti-PARN antibody at 1 $\mu g/ml$ dilution, overnight at 4°C.



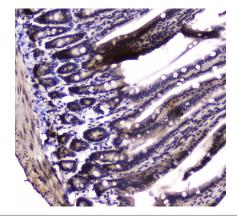
ARG59016 anti-PARN antibody WB image

Western blot: $50 \mu g$ of samples under reducing conditions. HeLa, Human placenta, COLO-320, HepG2, PANC-1, SGC-7901, MDA-MB-231, Rat kidney, Mouse heart and Mouse kidney lysates stained with ARG59016 anti-PARN antibody at $0.5 \mu g/ml$, overnight at 4° C.



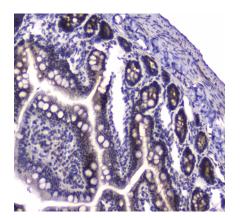
ARG59016 anti-PARN antibody FACS image

Flow Cytometry: A431 cells were blocked with 10% normal goat serum and then stained with ARG59016 anti-PARN antibody (blue) at 1 μ g/10^6 cells for 30 min at 20°C, followed by incubation with DyLight®488 labelled secondary antibody. Isotype control antibody (green) was rabbit IgG (1 μ g/10^6 cells) used under the same conditions. Unlabelled sample (red) was also used as a control.



ARG59016 anti-PARN antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Mouse small intestine tissue. Antigen Retrieval: Heat mediated was performed in Citrate buffer (pH 6.0, epitope retrieval solution) for 20 min. The tissue section was blocked with 10% goat serum. The tissue section was then stained with ARG59016 anti-PARN antibody at 1 $\mu g/ml$ dilution, overnight at 4°C.



ARG59016 anti-PARN antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Rat small intestine tissue. Antigen Retrieval: Heat mediated was performed in Citrate buffer (pH 6.0, epitope retrieval solution) for 20 min. The tissue section was blocked with 10% goat serum. The tissue section was then stained with ARG59016 anti-PARN antibody at 1 $\mu g/ml$ dilution, overnight at 4°C.