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Datasheet for ABIN6256820 anti-NONO antibody (C-Term)

2 Images

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Overview

Quantity:	100 µL
Target:	NONO
Binding Specificity:	C-Term
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This NONO antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)

Product Details

Immunogen:	A synthesized peptide derived from human Nono, corresponding to a region within C-terminal amino acids.
Isotype:	lgG
Specificity:	Nono Antibody detects endogenous levels of total Nono.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).

Target Details

Target:	NONO	
	Order at www.antibodies-online.com www.antikoerper-online.de www.anticorps-enligne.fr www.antibodies-online.cn International: +49 (0)241 95 163 153 USA & Canada: +1 877 302 8632 support@antibodies-online.com	

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Target Details	
Alternative Name:	NONO (NONO Products)
Background:	Description: DNA- and RNA binding protein, involved in several nuclear processes. Binds the
	conventional octamer sequence in double-stranded DNA. Also binds single-stranded DNA and
	RNA at a site independent of the duplex site. Involved in pre-mRNA splicing, probably as a
	heterodimer with SFPQ. Interacts with U5 snRNA, probably by binding to a purine-rich sequence
	located on the 3' side of U5 snRNA stem 1b. Together with PSPC1, required for the formation of
	nuclear paraspeckles. The SFPQ-NONO heteromer associated with MATR3 may play a role in
	nuclear retention of defective RNAs. The SFPQ-NONO heteromer may be involved in DNA
	unwinding by modulating the function of topoisomerase I/TOP1. The SFPQ-NONO heteromer
	may be involved in DNA non-homologous end joining (NHEJ) required for double-strand break
	repair and V(D)J recombination and may stabilize paired DNA ends. In vitro, the complex
	strongly stimulates DNA end joining, binds directly to the DNA substrates and cooperates with
	the Ku70/G22P1-Ku80/XRCC5 (Ku) dimer to establish a functional preligation complex. NONO
	is involved in transcriptional regulation. The SFPQ-NONO-NR5A1 complex binds to the CYP17
	promoter and regulates basal and cAMP-dependent transcriptional activity. NONO binds to an
	enhancer element in long terminal repeats of endogenous intracisternal A particles (IAPs) and
	activates transcription. Regulates the circadian clock by repressing the transcriptional activator
	activity of the CLOCK-ARNTL/BMAL1 heterodimer. Important for the functional organization of
	GABAergic synapses. Plays a specific and important role in the regulation of synaptic RNAs and
	GPHN/gephyrin scaffold structure, through the regulation of GABRA2 transcript. Plays a role in
	the regulation of DNA virus-mediated innate immune response by assembling into the HDP-
	RNP complex, a complex that serves as a platform for IRF3 phosphorylation and subsequent
	innate immune response activation through the cGAS-STING pathway (PubMed:28712728).
	Gene: NONO
Molecular Weight:	54kDa
Gene ID:	4841
UniProt:	Q15233

Application Details

Application Notes: WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000

Restrictions:

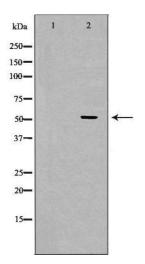
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Handling

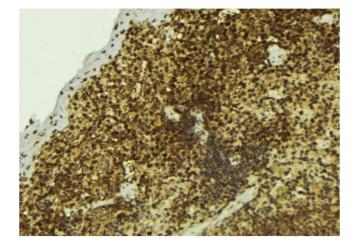
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months

Images



Western Blotting

Image 1. Western blot analysis of Hela whole cell lysates, using Nono Antibody. The lane on the left is treated with the antigen-specific peptide.



Immunohistochemistry

Image 2. ABIN6277498 at 1/100 staining Mouse pancreas tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22jãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary

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