

Datasheet for ABIN7600723 anti-NUP133 antibody (AA 228-1156)



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Quantity:	100 μg	
Target:	NUP133	
Binding Specificity:	AA 228-1156	
Reactivity:	Human, Mouse, Rat	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This NUP133 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunoprecipitation (IP), ELISA, Immunocytochemistry (ICC), Immunofluorescence (IF)	

Product Details

Purpose:	Anti-NUP133 Antibody Picoband®	
Immunogen:	E.coli-derived human NUP133 recombinant protein (Position: Q228-I1156).	
Isotype:	IgG	
Cross-Reactivity (Details):	No cross-reactivity with other proteins.	
Characteristics:	Anti-N Antibody Picoband® (ABIN7600723). Tested in ELISA, IP, IF, ICC, WB applications. This antibody reacts with Human, Mouse, Rat. The brand Picoband indicates this is a premium antibody that guarantees superior quality, high affinity, and strong signals with minimal background in Western blot applications. Only our best-performing antibodies are designated as Picoband, ensuring unmatched performance.	
Purification:	Immunogen affinity purified.	

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Target Details

rarget Details	
Target:	NUP133
Alternative Name:	NUP133 (NUP133 Products)
Background:	Synonyms: Cytoskeleton-associated protein 5, Colonic and hepatic tumor overexpressed gene
	protein, Ch-TOG, CKAP5, KIAA0097
	Tissue Specificity: Overexpressed in hepatomas and colonic tumors. Also expressed in skeleta
	muscle, brain, heart, placenta, lung, liver, kidney and pancreas. Expression is elevated in the
	brain, highly expressed in the Purkinje cell bodies of the cerebellum.
	Background: Nuclear pore complex protein Nup133, or Nucleoporin Nup133, is a protein that in
	humans is encoded by the NUP133 gene. The nuclear envelope creates distinct nuclear and
	cytoplasmic compartments in eukaryotic cells. It consists of two concentric membranes
	perforated by nuclear pores, large protein complexes that form aqueous channels to regulate
	the flow of macromolecules between the nucleus and the cytoplasm. These complexes are
	composed of at least 100 different polypeptide subunits, many of which belong to the
	nucleoporin family. The nucleoporin protein encoded by this gene displays evolutionarily
	conserved interactions with other nucleoporins. This protein, which localizes to both sides of
	the nuclear pore complex at interphase, remains associated with the complex during mitosis
	and is targeted at early stages to the reforming nuclear envelope. This protein also localizes to
	kinetochores of mitotic cells.
Molecular Weight:	129 kDa
Gene ID:	55746
Application Details	
Application Notes:	Western blot, 0.25-0.5 μg/mL, Human, Mouse, Rat
	Immunocytochemistry/Immunofluorescence, 5 µg/mL, Human, Rat
	Immunoprecipitation, 0.5-2 μg/mL, Human
	ELISA, 0.1-0.5 μg/mL, -
	1. Belgareh, N., Rabut, G., Bai, S. W., van Overbeek, M., Beaudouin, J., Daigle, N., Zatsepina, O. V
	Pasteau, F., Labas, V., Fromont-Racine, M., Ellenberg, J., Doye, V. An evolutionarily conserved
	NPC subcomplex, which redistributes in part to kinetochores in mammalian cells. J. Cell. Biol.
	154: 1147-1160, 2001. 2. Braun, D. A., Lovric, S., Schapiro, D., Schneider, R., Marquez, J., Asif, N

Hussain, M. S., Daga, A., Widneier, E., Rao, J., Ashraf, S., Tan, W., and 46 others. Mutations in multiple components of the nuclear pore complex cause nephrotic syndrome. J. Clin. Invest.

S., Komohara, Y., Shiina, M., Nakamura, S., Kitajima, M., Tsurusaki, Y., Miyatake, S., Ogata, K.,

128: 4313-4328, 2018. 3. Fujita, A., Tsukaguchi, H., Koshimizu, E., Nakazato, H., Itoh, K., Kuraoka,

Application Details

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	lijima, K., Matsumoto, N., Miyake, N. Homozygous splicing mutation in NUP133 causes
	Galloway-Mowat syndrome. Ann. Neurol. 84: 814-828, 2018. Note: Erratum: Ann. Neurol. 85:
	462-463, 2019.
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Adding 0.2 mL of distilled water will yield a concentration of 500 µg/mL.
Concentration:	500 μg/mL
Buffer:	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na2HPO4.
Storage:	4 °C,-20 °C
Storage Comment:	At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month.
	It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and
	thawing.