

NAT10 Antibody (N-Term)

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP22316a

Product Information

Application Primary Accession	WB, FC, E <u>Q9H0A0</u> 08K224
Other Accession Reactivity	<u>Q8K224</u> Human, Mouse
Predicted	Mouse
Host	Rabbit
Clonality	polyclonal
Isotype	Rabbit IgG
Clone Names	RB57638
Calculated MW	115730

Additional Information

Gene ID	55226
Other Names	N-acetyltransferase 10, 2.3.1, NAT10, ALP, KIAA1709
Target/Specificity	This NAT10 antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 146-178 amino acids from the mouse region of mouse NAT10.
Dilution	WB~~1:2000 FC~~1:25 E~~Use at an assay dependent concentration.
Format	Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Precautions	NAT10 Antibody (N-Term) is for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name	NAT10 {ECO:0000255 HAMAP-Rule:MF_03211}
Function	RNA cytidine acetyltransferase that catalyzes the formation of N(4)-acetylcytidine (ac4C) modification on mRNAs, 18S rRNA and tRNAs (PubMed: <u>25411247</u> , PubMed: <u>25653167</u> , PubMed: <u>30449621</u> , PubMed: <u>35679869</u>). Catalyzes ac4C modification of a broad range of mRNAs,

	enhancing mRNA stability and translation (PubMed: <u>30449621</u> , PubMed: <u>35679869</u>). mRNA ac4C modification is frequently present within wobble cytidine sites and promotes translation efficiency (PubMed: <u>30449621</u>). Mediates the formation of ac4C at position 1842 in 185 rRNA (PubMed: <u>25411247</u>). May also catalyze the formation of ac4C at position 1337 in 185 rRNA (By similarity). Required for early nucleolar cleavages of precursor rRNA at sites A0, A1 and A2 during 185 rRNA synthesis (PubMed: <u>25411247</u> , PubMed: <u>25653167</u>). Catalyzes the formation of ac4C in serine and leucine tRNAs (By similarity). Requires the tRNA-binding adapter protein THUMPD1 for full tRNA acetyltransferase activity but not for 185 rRNA acetylation (PubMed: <u>25653167</u>). In addition to RNA acetyltransferase activity, also able to acetylate lysine residues of proteins, such as histones, microtubules, p53/TP53 and MDM2, in vitro (PubMed: <u>14592445</u> , PubMed: <u>17631499</u> , PubMed: <u>1903003</u> , PubMed: <u>14592445</u> , PubMed: <u>27993683</u> , PubMed: <u>30165671</u>). The relevance of the protein lysine acetyltransferase activity is however unsure in vivo (PubMed: <u>30449621</u>). Activates telomerase function by affecting the balance of telomerase subunit assembly, disassembly, and localization (PubMed: <u>14592445</u> , PubMed: <u>18082603</u>). Involved in the regulation of centrosome duplication by acetylating CENATAC during mitosis, promoting SASS6 proteasome degradation (PubMed: <u>31722219</u>). Part of the small subunit (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the small eukaryotic ribosomal subunit. (SSU) processome, first precursor of the
Cellular Location	Nucleus, nucleolus {ECO:0000255 HAMAP- Rule:MF_03211, ECO:0000269 PubMed:12429849, ECO:0000269 PubMed:14592445, ECO:0000269 PubMed:19303003, ECO:0000269 PubMed:24786082, ECO:0000269 PubMed:25653167, ECO:0000269 PubMed:30165671, ECO:0000269 PubMed:34516797}. Midbody {ECO:0000255 HAMAP-Rule:MF_03211, ECO:0000269 PubMed:19303003} Note=Nucleolar in interphase and redistributes to the perichromosomal layer and to the midbody during telophase

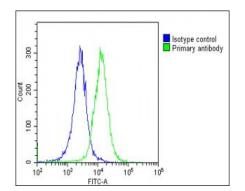
Background

Has protein acetyltransferase activity in vitro. Can acetylate both histones and microtubules. Histone acetylation may regulate transcription and mitotic chromosome de-condensation. Activates telomerase activity by stimulating the transcription of TERT, and may also regulate telomerase function by affecting the balance of telomerase subunit assembly, disassembly, and localization. Acetylates alpha-tubulin, which may affect microtubule stability and cell division.

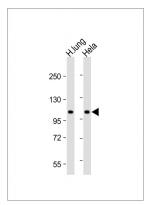
References

Nagase T.,et al.DNA Res. 7:347-355(2000). Wiemann S.,et al.Genome Res. 11:422-435(2001). Ota T.,et al.Nat. Genet. 36:40-45(2004). Taylor T.D.,et al.Nature 440:497-500(2006). Lv J.,et al.Biochem. Biophys. Res. Commun. 311:506-513(2003).

Images



Overlay histogram showing Hela cells stained with AP22316a(green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody (AP22316a, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Goat-Anti-Rabbit IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed(OE188374) at 1/200 dilution for 40 min at 37°C. Isotype control antibody (blue line) was rabbit IgG1 (1µg/1x10^6 cells) used under the same conditions. Acquisition of >10, 000 events was performed.



All lanes : Anti-NAT10 Antibody (N-Term) at 1:2000 dilution Lane 1: Human lung lysate Lane 2: Hela whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 116 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.