Datasheet for ABIN1705096 anti-NDUFAF7 antibody (AA 101-200) (Cy5)

-online.com antibodies



Overview

Quantity:	100 µL	
Target:	NDUFAF7	
Binding Specificity:	AA 101-200	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This NDUFAF7 antibody is conjugated to Cy5	
Application:	Western Blotting (WB), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p))	

Product Details

Immunogen:	KLH conjugated synthetic peptide derived from human C2orf56	
Isotype:	lgG	
Predicted Reactivity:	Human,Mouse,Rat,Cow,Sheep,Pig,Rabbit	
Purification:	Purified by Protein A.	

Target Details

Target:	NDUFAF7	
Alternative Name:	C2orf56 (NDUFAF7 Products)	
Background:	Synonyms: C2orf56, Chromosome 2 open reading frame 56, MidA, MIDA_HUMAN,	

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 Page 1/3 | Product datasheet for ABIN1705096 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

mitochondrial, Mitochondrial dysfunction protein A homolog, OTTHUMP00000158583,
OTTHUMP00000201359, OTTHUMP00000201362, PR01853, Protein midA homolog, Protein
midA homolog, mitochondrial.
Background: C12orf56 (chromosome 12 open reading frame 56), also known as PRO1853 or
protein midA homolog, is a 441 amino acid mitochondrial protein that belongs to the midA
family. Existing as two alternatively spliced isoforms, C12orf56 is encoded by a gene that maps
to human chromosome 2p22.2. As the second largest human chromosome, chromosome 2
makes up approximately 8 % of the human genome and contains 237 million bases encoding
over 1,400 genes. A number of genetic diseases are linked to genes on chromosome 2.
Harlequin icthyosis, a rare skin deformity, is associated with mutations in the ABCA12 gene.
The lipid metabolic disorder sitosterolemia is associated with ABCG5 and ABCG8. An extremely
rare recessive genetic disorder, Alstr syndrome, is related to mutations in the ALMS1 gene.
Chromosome 2 contains a probable vestigial second centromere as well as vestigial telomeres,
which gives credence to the hypothesis that human chromosome 2 formed as a result of an
ancient fusion of two ancestral chromosomes, which are still present in modern day apes.

Gene ID:

55471

Application Details

1-1		
Application Notes:	IF(IHC-P) 1:50-200	
	IF(IHC-F) 1:50-200	
	IF(ICC) 1:50-200	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	1 µg/µL	
Buffer:	Aqueous buffered solution containing 0.01M TBS (pH 7.4) with 1 % BSA, 0.03 % Proclin300 and	
	50 % Glycerol.	
Preservative:	ProClin	
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be	
	handled by trained staff only.	
Storage:	-20 °C	

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 Page 2/3 | Product datasheet for ABIN1705096 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

lond	lina
land	
10110	

Storage Comment:

Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles.

Expiry Date:

12 months

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 Page 3/3 | Product datasheet for ABIN1705096 | 03/07/2024 | Copyright antibodies-online. All rights reserved.