.-online.com antibodies

# Datasheet for ABIN896007 anti-FAHD2A antibody (AA 121-230) (Alexa Fluor 488)



Overview

Quantity:	100 μL	
Target:	FAHD2A	
Binding Specificity:	AA 121-230	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This FAHD2A antibody is conjugated to Alexa Fluor 488	
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 FAHD2A	
Isotype:	lgG	
Predicted Reactivity:	Human,Mouse,Rat,Horse,Rabbit	
Purification:	Purified by Protein A.	

#### Target Details

Target:	FAHD2A	
Alternative Name:	FAHD2A (FAHD2A Products)	
Background: Synonyms: CGI 105, FAH2A_HUMAN, FAHD 2A, FAHD2A, Fumarylacetoacetate hydrolase		

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 ABIN896007 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

domain containing 1, Fumarylacetoacetate hydrolase domain containing 2A,
Fumarylacetoacetate hydrolase domain containing protein 2A, Fumarylacetoacetate hydrolase
domain-containing protein 2A.
Background: The FAH family contains two highly homologous 314 amino acid proteins,
designated FAHD2A (fumarylacetoacetate hydrolase domain-containing protein 2A) and
FAHD2B (fumarylacetoacetate hydrolase domain-containing protein 2A). FAHD2A and B utilize
calcium and magnesium as cofactors, and may possess hydrolase activity. The genes
encoding FAHD2A/B map to human chromosome 2, the second largest human chromosome,
which consists of 237 million bases, encodes over 1,400 genes and makes up approximately
8 % of the human genome. A number of genetic diseases are linked to genes on chromosome
2. Harlequin icthyosis, a rare and morbid 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 also associated with
mutations to chromosome 2.

Gene ID:

51011

## Application Details

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

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 ABIN896007 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

1.1	(	1:
Н	land	ling
		3

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 ABIN896007 | 03/07/2024 | Copyright antibodies-online. All rights reserved.