antibodies.com

# Datasheet for ABIN888081 anti-CCDC69 antibody (AA 41-140) (Alexa Fluor 555)



Overview

Quantity:	100 µL
Target:	CCDC69
Binding Specificity:	AA 41-140
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This CCDC69 antibody is conjugated to Alexa Fluor 555
Application:	Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p))

## Product Details

Isotype:IgGCross-Reactivity:HumanPredicted Reactivity:Dog,Cow,Sheep,Rabbit	Immunogen:	KLH conjugated synthetic peptide derived from human CCDC69
	lsotype:	lgG
Predicted Reactivity: Dog,Cow,Sheep,Rabbit	Cross-Reactivity:	Human
	Predicted Reactivity:	Dog,Cow,Sheep,Rabbit
Purification: Purified by Protein A.	Purification:	Purified by Protein A.

#### Target Details

Target:	CCDC69
Alternative Name:	CCDC69 (CCDC69 Products)

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 1/2 | Product datasheet for ABIN888081 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

Target Details	
Background:	Synonyms: CCD69_HUMAN, ccdc69, Coiled coil domain containing 69, Coiled-coil domain-
	containing protein 69.
	Background: The coiled-coil domain is a structural motif found in proteins that are involved in a
	diverse array of biological functions such as the regulation of gene expression, cell division,
	membrane fusion and drug extrusion and delivery. CCDC69 (Coiled-coil domain-containing
	protein 69) is a 296 amino acid protein that is encoded by a gene which maps to human
	chromosome 5, which contains 181 million base pairs and comprises nearly 6 % of the human
	genome. Chromosome 5 is associated with Cockayne syndrome through the ERCC8 gene and
	familial adenomatous polyposis through the adenomatous polyposis coli (APC) tumor
	suppressor gene. Treacher Collins syndrome is also chromosome 5-associated and is caused
	by insertions or deletions within the TCOF1 gene. Deletion of the p arm of chromosome 5 leads
	to Cri du chat syndrome, while deletion of the q arm or of chromosome 5 altogether is commo
	in therapy-related acute myelogenous leukemias and myelodysplastic syndrome.
Gene ID:	26112

# 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.
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 2/2 | Product datasheet for ABIN888081 | 03/07/2024 | Copyright antibodies-online. All rights reserved.