antibodies.com

Datasheet for ABIN6978380 anti-SBNO1 antibody (AA 331-430) (Alexa Fluor 555)



Overview

Quantity:	100 µL
Target:	SBN01
Binding Specificity:	AA 331-430
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SBN01 antibody is conjugated to Alexa Fluor 555
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 SBNO1
lsotype:	lgG
Cross-Reactivity:	Human
Predicted Reactivity:	Mouse,Rat,Dog,Cow,Sheep,Pig,Horse,Chicken,Rabbit,Zebrafish
Purification:	Purified by Protein A.
Target Details	
Target:	SBN01

Target:	SBN01
Alternative Name:	SBN01 (SBN01 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 1/3 | Product datasheet for ABIN6978380 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

Background:	Synonyms: FLJ10701, FLJ10833, FLJ16176, Monocyte protein 3, MOP 3, MOP-3, MOP3, Proteir
	strawberry notch homolog 1, SBNO 1, Sbno1, SBNO1_HUMAN, Sno, Sno strawberry notch
	homolog 1, Strawberry notch homolog 1.
	Background: SBN01 is a 1,392 amino acid protein encoded by the human gene of the same
	name located on chromosome 12. Encoding over 1,100 genes within 132 million bases,
	chromosome 12 makes up about 4.5 % of the human genome. A number of skeletal
	deformities are linked to chromosome 12, including hypochondrogenesis, achondrogenesis and
	Kniest dysplasia. Noonan syndrome, which includes heart and facial developmental defects
	among the primary symptoms, is caused by a mutant form of PTPN11 gene product, SH-PTP2.
	Chromosome 12 is also home to a homeobox gene cluster, which encodes crucial transcription
	factors for morphogenesis, and the natural killer complex gene cluster, encoding C-type lectin
	proteins which mediate the NK cell response to MHC I interaction. Trisomy 12p leads to facial
	development defects, seizure disorders and a host of other symptoms which vary in severity
	depending on the extent of mosaicism. It is most severe in cases of complete trisomy.
Gene ID:	55206
UniProt:	A3KN83
Pathways:	SARS-CoV-2 Protein Interactome
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.

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 ABIN6978380 | 03/07/2024 | Copyright antibodies-online. All rights reserved.

Handling	
Storage:	-20 °C
Storage Comment:	Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles.
Expiry Date:	12 months