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Datasheet for ABIN6980034 anti-SBN01 antibody (AA 331-430) (Alexa Fluor 647)



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 647 |
| Application: | Western Blotting (WB), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

Product Details

Alternative Name:

| Sheep,Pig,Horse,Chicken,Rabbit,Zebrafish |
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| 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 |
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SBN01 (SBN01 Products)

| 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. |

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