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Datasheet for ABIN6989487

anti-SBNO1 antibody (AA 331-430)



Overview

| Quantity: | 100 μL |
|----------------------|---|
| Target: | SBN01 |
| Binding Specificity: | AA 331-430 |
| Reactivity: | Human |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This SBNO1 antibody is un-conjugated |
| Application: | Western Blotting (WB), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)), ELISA, Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Immunohistochemistry (Frozen Sections) (IHC (fro)), Immunocytochemistry (ICC) |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human SBN01 |
|-----------------------|--|
| Isotype: | IgG |
| Cross-Reactivity: | Human |
| Predicted Reactivity: | Mouse,Rat,Dog,Cow,Sheep,Pig,Horse,Chicken,Rabbit,Zebrafish |
| Purification: | Purified by Protein A. |

Target Details

| 9 | Target: | SBN01 |
|---|---------|-------|
|---|---------|-------|

Target Details

| Alternative Name: | 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: SBNO1 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: | WB 1:300-5000 |
| | ELISA 1:500-1000 |
| | IHC-P 1:200-400 |
| | IHC-F 1:100-500 |
| | IF(IHC-P) 1:50-200 |
| | IF(IHC-F) 1:50-200 |
| | IF(ICC) 1:50-200 |
| | ICC 1:100-500 |
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Liquid |
| Concentration: | 1 μg/μL |
| | |

Handling

| Buffer: | 0.01M TBS(pH 7.4) with 1 % BSA, 0.02 % 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: | 4 °C,-20 °C |
| Storage Comment: | Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles. |
| Expiry Date: | 12 months |