

Datasheet for ABIN1415675

anti-C12ORF23 antibody (AA 68-116) (Cy5)



| \sim | | | |
|--------|------------|-----|-----|
| ()\ | / e | rVI | iew |

| Overview | |
|----------------------|---|
| Quantity: | 100 μL |
| Target: | C120RF23 |
| Binding Specificity: | AA 68-116 |
| Reactivity: | Human |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This C120RF23 antibody is conjugated to Cy5 |
| Application: | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |
| Product Details | |
| Immunogon: | KLH conjugated cynthotic pontide derived from human C12orf22 |

| Immunogen: | KLH conjugated synthetic peptide derived from human C12orf23 |
|-----------------------|--|
| Isotype: | IgG |
| Predicted Reactivity: | Human,Mouse,Rat,Dog,Cow,Pig,Horse,Chicken,Rabbit |
| Purification: | Purified by Protein A. |

Target Details

| Target: | C120RF23 | |
|-------------------|---|--|
| Alternative Name: | C12orf23 (C12ORF23 Products) | |
| Background: | Synonyms: C12orf23, Chromosome 12 open reading frame 23, CL023_HUMAN, MGC17943, | |

UPF0444 transmembrane protein C12orf23.

Background: C12orf23 (chromosome 12 open reading frame 23), also known as FLJ11721, FLJ13959 or MGC17943, is a 116 amino acid multi-pass membrane protein belonging to the UPF0444 family. C12orf23 is encoded by a gene located on human chromosome 12, which encodes over 1,100 genes and comprises approximately 4.5 % of the human genome. Chromosome 12 is associated with a number of skeletal deformities, 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 varying in severity depending on the extent of mosaicism and is most severe in cases of complete trisomy.

Gene ID:

90488

Application Details

| Application Notes: | |
|--------------------|--|
| 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. |

| | 1 | l: |
|---|------|--------|
| - | ココロロ | urnen |
| | Hand | III IU |
| | | |

Expiry Date:

12 months