antibodies

Datasheet for ABIN1405630 anti-C12orf53 antibody (Alexa Fluor 488)



| Overview | |
|---|---|
| Quantity: | 100 µL |
| Target: | C12orf53 |
| Reactivity: | Human, Mouse, Rat |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This C12orf53 antibody is conjugated to Alexa Fluor 488 |
| Application: | Western Blotting (WB), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |
| Product Details | |
| Immunogen: | KLH conjugated synthetic peptide derived from human C12ORF53 |
| lsotype: | |
| | IgG |
| Cross-Reactivity: | Human, Mouse, Rat |
| | |
| Cross-Reactivity: | Human, Mouse, Rat |
| Cross-Reactivity: Purification: | Human, Mouse, Rat |
| Cross-Reactivity: Purification: Target Details | Human, Mouse, Rat Purified by Protein A. |
| Cross-Reactivity: Purification: Target Details Target: | Human, Mouse, Rat Purified by Protein A. C12orf53 |

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 ABIN1405630 | 03/06/2024 | Copyright antibodies-online. All rights reserved.

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 varying in severity depending on the extent of |
| | mosaicism and is most severe in cases of complete trisomy. The C12orf53 gene product has been provisionally designated C12orf53 pending further characterization. |
| | |
| Gene ID: | 196500 |
| Application Details | |
| Application Notes: | IF(IHC-P) 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 |
| | |