

Datasheet for ABIN1692678

anti-C9orf163 antibody (AA 101-203) (AbBy Fluor® 350)



Go to Product page

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| Quantity: 100 µL Target: C9orf163 (C9ORF163) Binding Specificity: AA 101-203 Reactivity: Human Host: Rabbit Clonality: Polyclonal Conjugate: This C9orf163 antibody is conjugated to AbBy Fluor® 350 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 C9orf163 Isotype: IgG Predicted Reactivity: Human Purification: Purified by Protein A. Target Details Target: C9orf163 (C9ORF163) Alternative Name: C9orf163 (C9ORF163) Products) Background: Synonyms: C9orf163, Chromosome 9 open reading frame 163, C1163_HUMAN, FLJ36779, | Overview | |
|---|-----------------------|---|
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| Purification: Purified by Protein A. Target Details Target: C9orf163 (C9ORF163) Alternative Name: C9orf163 (C9ORF163 Products) | Isotype: | IgG |
| Target Details Target: C9orf163 (C9ORF163) Alternative Name: C9orf163 (C9ORF163 Products) | Predicted Reactivity: | Human |
| Target: C9orf163 (C9ORF163) Alternative Name: C9orf163 (C9ORF163 Products) | Purification: | Purified by Protein A. |
| Alternative Name: C9orf163 (C9ORF163 Products) | Target Details | |
| | Target: | C9orf163 (C9ORF163) |
| Background: Synonyms: C9orf163, Chromosome 9 open reading frame 163, Cl163_HUMAN, FLJ36779, | Alternative Name: | C9orf163 (C9ORF163 Products) |
| | Background: | Synonyms: C9orf163, Chromosome 9 open reading frame 163, Cl163_HUMAN, FLJ36779, |

| RP11-413M3.11 | , RP11-413M3.11-001, | Uncharacterized | protein C9orf163. |
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Background: C9orf163 (chromosome 9 open reading frame 163) is a 203 amino acid protein encoded by a gene that maps to human chromosome 9q34.3. Chromosome 9 consists of about 145 million bases, represents 4 % of the human genome and encodes nearly 900 genes. Thought to play a role in gender determination, deletion of the distal portion of 9p can lead to development of male to female sex reversal, the phenotype of a female with a male X,Y genotype. Hereditary hemorrhagic telangiectasia, which is characterized by harmful vascular defects, is associated with the chromosome 9 gene encoding endoglin protein, ENG. Familial dysautonomia is also associated with chromosome 9 though through the gene IKBKAP. Notably, chromosome 9 encompasses the largest interferon family gene cluster. Chromosome 9 is partnered with chromosome 22 in the translocation leading to the aberrant production of BCR-ABL fusion protein often found in leukemias.

Gene ID:

158055

Application Details

| Application Notes: | IF(IHC-P) 1:50-200 |
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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. |
| Expiry Date: | 12 months |