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## Datasheet for ABIN1385421 anti-C9orf153 antibody



Overview

| Quantity:    | 100 µL  |
|--------------|---|
| Target:      | C9orf153  |
| Reactivity:  | Human   |
| Host:        | Rabbit  |
| Clonality:   | Polyclonal  |
| Conjugate:   | This C9orf153 antibody is un-conjugated   |
| Application: | Immunofluorescence (Paraffin-embedded Sections) (IF (p)), Western Blotting (WB),<br>Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)) |

## Product Details

| Immunogen:        | KLH conjugated synthetic peptide derived from human C9orf153 |
|-------------------|--|
| Isotype:          | lgG  |
| Cross-Reactivity: | Human  |
| Purification:     | Purified by Protein A.                                       |

## Target Details

| Target:           | C9orf153  |
|-------------------|---|
| Alternative Name: | C9orf153 (C9orf153 Products)  |
| Background:       | Synonyms: bA507D14.1, Chromosome 9 open reading frame 153, Hypothetical protein LOC389766, MGC131702, Uncharacterized protein C9orf153, |
|                   | Background: C9orf153 is a 101 amino acid protein that exists as two alternatively spliced   |

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|  | isoforms. The gene encoding C9orf153 maps to human chromosome 9q21.33. 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:                                       | 389766   |
| Application Details                            |  |
| Application Notes:                             | WB 1:300-5000  |
|  | IHC-P 1:200-400  |
|  | IF(IHC-P) 1:50-200   |
| Restrictions:                                  | For Research Use only  |
| Handling                                       |  |
| Format:  | Liquid   |
|  |  |
| Concentration:                                 | 1 μg/μL  |
| Concentration:<br>Buffer:                      | 1 μg/μL<br>0.01M TBS( pH 7.4) with 1 % BSA, 0.02 % Proclin300 and 50 % Glycerol.   |
|  |  |
| Buffer:  | 0.01M TBS( pH 7.4) with 1 % BSA, 0.02 % Proclin300 and 50 % Glycerol.  |
| Buffer:<br>Preservative:                       | 0.01M TBS( pH 7.4) with 1 % BSA, 0.02 % Proclin300 and 50 % Glycerol.<br>ProClin   |
| Buffer:<br>Preservative:                       | 0.01M TBS( pH 7.4) with 1 % BSA, 0.02 % Proclin300 and 50 % Glycerol.         ProClin         This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be  |
| Buffer:<br>Preservative:<br>Precaution of Use: | <ul> <li>0.01M TBS( pH 7.4) with 1 % BSA, 0.02 % Proclin300 and 50 % Glycerol.</li> <li>ProClin</li> <li>This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only.</li> </ul> |

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