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## Datasheet for ABIN1700149 **anti-CCDC171 antibody (AA 1-100) (Biotin)**

### Overview

|                      |   |
|----------------------|---|
| Quantity:            | 100 µL  |
| Target:              | CCDC171   |
| Binding Specificity: | AA 1-100  |
| Reactivity:          | Human   |
| Host:                | Rabbit  |
| Clonality:           | Polyclonal  |
| Conjugate:           | This CCDC171 antibody is conjugated to Biotin   |
| Application:         | Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Immunohistochemistry (Frozen Sections) (IHC (fro)), Western Blotting (WB), ELISA |

### Product Details

|                       |   |
|-----------------------|---|
| Immunogen:            | KLH conjugated synthetic peptide derived from human C9orf93 |
| Isotype:              | IgG   |
| Predicted Reactivity: | Human,Dog,Cow,Sheep,Pig,Horse,Rabbit                        |
| Purification:         | Purified by Protein A.                                      |

### Target Details

|                   |  |
|-------------------|--|
| Target:           | CCDC171  |
| Alternative Name: | C9orf93 ( <a href="#">CCDC171 Products</a> )                                   |
| Background:       | Synonyms: bA536D16.1, bA778P13.1, C9orf93, Chromosome 9 open reading frame 93, |

## Target Details

CC171\_HUMAN, Uncharacterized protein C9orf93.

Background: Chromosome 9 consists of about 145 million bases and 4 % of the human genome and encodes nearly 900 genes. Considered 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. The C9orf93 gene product has been provisionally designated C9orf93 pending further characterization. There are two isoforms of C9orf93 that are produced as a result of alternative splicing events.

Gene ID: 203238

## Application Details

Application Notes: WB 1:300-5000  
IHC-P 1:200-400  
IHC-F 1:100-500

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 for 12 months.

Expiry Date: 12 months