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# Datasheet for ABIN4999293 anti-CCDC69 antibody (AA 41-140) (Alexa Fluor 750)



Overview

| Quantity:            | 100 µL  |
|----------------------|---|
| Target:              | CCDC69  |
| Binding Specificity: | AA 41-140   |
| Reactivity:          | Human   |
| Host:                | Rabbit  |
| Clonality:           | Polyclonal  |
| Conjugate:           | This CCDC69 antibody is conjugated to Alexa Fluor 750   |
| Application:         | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

## Product Details

| Immunogen:            | KLH conjugated synthetic peptide derived from human CCDC69 |
|-----------------------|--|
| Isotype:              | IgG  |
| Cross-Reactivity:     | Human  |
| Predicted Reactivity: | Dog,Cow,Sheep,Rabbit                                       |
| Purification:         | Purified by Protein A.                                     |
|                       |  |

#### Target Details

| Target:           | CCDC69                   |
|-------------------|--------------------------|
| Alternative Name: | CCDC69 (CCDC69 Products) |

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| Background: | Synonyms: CCD69_HUMAN, ccdc69, Coiled coil domain containing 69, Coiled-coil domain-              |
|-------------|---|
|             | containing protein 69.  |
|             | Background: The coiled-coil domain is a structural motif found in proteins that are involved in a |
|             | diverse array of biological functions such as the regulation of gene expression, cell division,   |
|             | membrane fusion and drug extrusion and delivery. CCDC69 (Coiled-coil domain-containing            |
|             | protein 69) is a 296 amino acid protein that is encoded by a gene which maps to human             |
|             | chromosome 5, which contains 181 million base pairs and comprises nearly 6 % of the human         |
|             | genome. Chromosome 5 is associated with Cockayne syndrome through the ERCC8 gene and              |
|             | familial adenomatous polyposis through the adenomatous polyposis coli (APC) tumor                 |
|             | suppressor gene. Treacher Collins syndrome is also chromosome 5-associated and is caused          |
|             | by insertions or deletions within the TCOF1 gene. Deletion of the p arm of chromosome 5 leads     |
|             | to Cri du chat syndrome, while deletion of the q arm or of chromosome 5 altogether is commo       |
|             | in therapy-related acute myelogenous leukemias and myelodysplastic syndrome.                      |
| Gene ID:    | 26112   |
|             |   |

# Application Details

| 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.                                  |
| Expiry Date:       | 12 months  |

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