Datasheet for ABIN887459 anti-CCDC17 antibody (AA 351-450) (Biotin)

Overview

| Quantity:            | 100 µL   |  |
|----------------------|--|--|
| Target:              | CCDC17   |  |
| Binding Specificity: | AA 351-450   |  |
| Reactivity:          | Rat  |  |
| Host:                | Rabbit   |  |
| Clonality:           | Polyclonal   |  |
| Conjugate:           | This CCDC17 antibody is conjugated to Biotin   |  |
| Application:         | ELISA, Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Immunohistochemistry (Frozen Sections) (IHC (fro)) |  |

## **Product Details**

| Immunogen:            | KLH conjugated synthetic peptide derived from human CCDC17 |
|-----------------------|--|
| Isotype:              | lgG  |
| Cross-Reactivity:     | Rat  |
| Predicted Reactivity: | Human,Mouse,Cow,Sheep,Pig,Horse,Rabbit                     |
| Purification:         | Purified by Protein A.                                     |
| Target Details        |  |
| Target:               | CCDC17   |
| Alternative Name:     | CCDC17 (CCDC17 Products)                                   |

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| Target Details      |  |  |
|---------------------|--|--|
| Background:         | Synonyms: CCD17_HUMAN, CCDC17, Coiled coil domain containing 17, Coiled-coil domain-         |  |
|                     | containing protein 17, RP23-233B9.8, RP4-697E16.4.   |  |
|                     | Background: CCDC17, also known as FLJ17921 or RP4-697E16.4, is a 622 amino acid protein      |  |
|                     | expressed as four isoforms and encoded by a gene mapping to human chromosome 1.              |  |
|                     | Chromosome 1 is the largest human chromosome spanning about 260 million base pairs and       |  |
|                     | making up 8 % of the human genome. There are about 3,000 genes on chromosome 1, and          |  |
|                     | considering the great number of genes there are also a large number of diseases associated   |  |
|                     | with chromosome 1. Notably, the rare aging disease Hutchinson-Gilford progeria is associated |  |
|                     | with the LMNA gene which encodes lamin A. When defective, the LMNA gene product can build    |  |
|                     | up in the nucleus and cause characteristic nuclear blebs. The mechanism of rapidly enhanced  |  |
|                     | aging is unclear and is a topic of continuing exploration. The MUTYH gene is located on      |  |
|                     | chromosome 1 and is partially responsible for familial adenomatous polyposis. Stickler       |  |
|                     | syndrome, Parkinsons, Gaucher disease and Usher syndrome are also associated with            |  |
|                     | chromosome 1. A breakpoint has been identified in 1q which disrupts the DISC1 gene and is    |  |
|                     | linked to schizophrenia. Aberrations in chromosome 1 are found in a variety of cancers       |  |
|                     | including head and neck cancer, malignant melanoma and multiple myeloma.                     |  |
| Gene ID:            | 149483   |  |
| Application Details |  |  |
| Application Notes:  | 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   |

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 Storage Comment:
 Store at -20°C for 12 months.

Expiry Date:

12 months

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