

### Datasheet for ABIN4999234

# anti-CCDC19 antibody (AbBy Fluor® 680)



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| Quantity:    | 100 μL  |
|--------------|---|
| Target:      | CCDC19  |
| Reactivity:  | Human, Rat, Mouse                                     |
| Host:        | Rabbit  |
| Clonality:   | Polyclonal  |
| Conjugate:   | This CCDC19 antibody is conjugated to AbBy Fluor® 680 |
| Application: | Western Blotting (WB)                                 |

### **Product Details**

| Immunogen:            | KLH conjugated synthetic peptide derived from human CCDC19 |
|-----------------------|--|
| Isotype:              | IgG  |
| Predicted Reactivity: | Human,Mouse,Rat,Cow,Sheep,Horse,Rabbit                     |
| Purification:         | Purified by Protein A.                                     |

## **Target Details**

| Target:           | CCDC19  |
|-------------------|---|
| Alternative Name: | CCDC19 (CCDC19 Products)  |
| Background:       | Synonyms: Coiled coil domain containing 19, Nasopharyngeal epithelium specic protein 1,                               |
|                   | NESG1, RP11 190A12.6, CCD19_HUMAN.  Background: CCDC19 is a 466 amino acid protein 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, Parkinson?s, 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:

25790

### **Application Details**

| Application Notes: | IF(IHC-P) 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. |
| Storage:         | -20 °C   |
| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles.                          |
| Expiry Date:     | 12 months  |