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## Datasheet for ABIN1404623 anti-C1orf163 antibody (AbBy Fluor® 555)



| - ·                              |  |
|----------------------------------|--|
| Overview                         |  |
| Quantity:                        | 100 µL   |
| Target:                          | C1orf163   |
| Reactivity:                      | Human, Rat, Mouse  |
| Host:                            | Rabbit   |
| Clonality:                       | Polyclonal   |
| Conjugate:                       | This C1orf163 antibody is conjugated to AbBy Fluor® 555  |
| Application:                     | Western Blotting (WB), Immunofluorescence (Paraffin-embedded Sections) (IF (p))  |
| Product Details                  |  |
| Immunogen:                       | KLH conjugated synthetic peptide derived from human C1orf163   |
| Isotype:                         | lgG  |
| Cross-Reactivity:                | Human, Mouse, Rat  |
| Purification:                    | Purified by Protein A.   |
| Target Details                   |  |
| Target:                          | C1orf163   |
|                                  |  |
| Alternative Name:                | C1orf163 (C1orf163 Products)   |
| Alternative Name:<br>Background: | C1orf163 (C1orf163 Products)<br>Synonyms: Chromosome 1 open reading frame 163, FLJ12439, Hcp beta lactamase like protein |
|                                  |  |

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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. The       |
|          | C1orf163 gene product has been provisionally designated C1orf163 pending further              |
|          | characterization.   |
| Gene ID: | 65260   |

## 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.        |
| 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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