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## Datasheet for ABIN1692611 anti-C1ORF141 antibody (AA 101-200) (Alexa Fluor 350)



## Overview

| Quantity:            | 100 µL   |  |
|----------------------|--|--|
| Target:              | C10RF141   |  |
| Binding Specificity: | AA 101-200   |  |
| Reactivity:          | Human  |  |
| Host:                | Rabbit   |  |
| Clonality:           | Polyclonal   |  |
| Conjugate:           | This C10RF141 antibody is conjugated to Alexa Fluor 350  |  |
| Application:         | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)), Western Blotting (WB) |  |

## Product Details

| Immunogen:        | KLH conjugated synthetic peptide derived from human C1orf141 |  |
|-------------------|--|--|
| Isotype:          | IgG  |  |
| Cross-Reactivity: | Human  |  |
| Purification:     | Purified by Protein A.                                       |  |

## Target Details

| Target:           | C10RF141   |
|-------------------|--|
| Alternative Name: | C1orf141 (C10RF141 Products)                                       |
| Background:       | Synonyms: C1orf141, CA141_HUMAN, Uncharacterized protein C1orf141. |

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|                     | Background: 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. The       |  |
|                     | C1orf141 gene product has been provisionally designated C1orf141 pending further              |  |
|                     | characterization.   |  |
| Gene ID:            | 400757  |  |
| Application Details |   |  |
| Application Notes:  |   |  |
| Application Notes:  | IF(IHC-P) 1:50-200<br>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.             |  |
|                     |   |  |

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Expiry Date:

12 months

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