

Datasheet for ABIN1416511 anti-LRRC41 antibody (AA 331-430) (Cy5.5)



Overview

| Quantity: | 100 µL | |
|----------------------|--|--|
| Target: | LRRC41 | |
| Binding Specificity: | AA 331-430 | |
| Reactivity: | Human, Rat, Mouse | |
| Host: | Rabbit | |
| Clonality: | Polyclonal | |
| Conjugate: | This LRRC41 antibody is conjugated to Cy5.5 | |
| Application: | Western Blotting (WB), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) | |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human LRRC41 | |
|-----------------------|--|--|
| lsotype: | lgG | |
| Cross-Reactivity: | Human, Mouse, Rat | |
| Predicted Reactivity: | Dog,Cow,Sheep,Pig,Horse | |
| Purification: | Purified by Protein A. | |
| Target Details | | |
| Target: | LRRC41 | |
| Alternative Name: | LRRC41 (LRRC41 Products) | |

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| Background: | Synonyms: elongin BC interacting leucine rich repeat protein, Leucine rich repeat containing 41, | |
|---------------------|---|--|
| | Leucine rich repeat containing protein 41, MUF1, PP7759, LRC41_HUMAN. | |
| | 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. | |
| Gene ID: | 10489 | |
| 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. | |
| | -20 °C | |
| Storage: | -20 °C | |

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Expiry Date:

12 months

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