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## Datasheet for ABIN1421681 **anti-C17orf49 antibody (Cy3)**

### Overview

|              |   |
|--------------|---|
| Quantity:    | 100 µL  |
| Target:      | C17orf49  |
| Reactivity:  | Human, Mouse, Rat   |
| Host:        | Rabbit  |
| Clonality:   | Polyclonal  |
| Conjugate:   | This C17orf49 antibody is conjugated to Cy3                                     |
| Application: | Western Blotting (WB), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

### Product Details

|                   |  |
|-------------------|--|
| Immunogen:        | KLH conjugated synthetic peptide derived from human C17orf49 |
| Isotype:          | IgG  |
| Cross-Reactivity: | Human, Mouse, Rat  |
| Purification:     | Purified by Protein A.                                       |

### Target Details

|                   |  |
|-------------------|--|
| Target:           | C17orf49   |
| Alternative Name: | C17orf49 ( <a href="#">C17orf49 Products</a> )   |
| Background:       | <p>Synonyms: Bap18, BAP18_HUMAN, BPTF-associated protein of 18 kDa, Chromatin complexes subunit BAP18, Chromosome 17 open reading frame 49, Hypothetical protein LOC124944, MGC49942, Uncharacterized potential DNA binding protein C17orf49.</p> <p>Background: C17orf49 is a 172 amino acid protein that is encoded by a gene mapping to</p> |

## Target Details

human chromosome 17. Chromosome 17 makes up over 2.5 % of the human genome with about 81 million bases encoding over 1,200 genes. Two key tumor suppressor genes are associated with chromosome 17, namely, p53 and BRCA1. Tumor suppressor p53 is necessary for maintenance of cellular genetic integrity by moderating cell fate through DNA repair versus cell death. Malfunction or loss of p53 expression is associated with malignant cell growth and Li-Fraumeni syndrome. Like p53, BRCA1 is directly involved in DNA repair, specifically it is recognized as a genetic determinant of early onset breast cancer and predisposition to cancers of the ovary, colon, prostate gland and fallopian tubes. Chromosome 17 is also linked to neurofibromatosis, a condition characterized by neural and epidermal lesions, and dysregulated Schwann cell growth. Alexander disease, Birt-Hogg-Dube syndrome and Canavan disease are also associated with chromosome 17.

Gene ID: 124944

## 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