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## anti-FAM65B antibody (AA 201-300) (Cy7)



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| Quantity:            | 100 μL   |  |
|----------------------|--|--|
| Target:              | FAM65B   |  |
| Binding Specificity: | AA 201-300   |  |
| Reactivity:          | Mouse  |  |
| Host:                | Rabbit   |  |
| Clonality:           | Polyclonal   |  |
| Conjugate:           | This FAM65B antibody is conjugated to Cy7  |  |
| 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 FAM65B |  |
|-----------------------|--|--|
| Isotype:              | IgG  |  |
| Cross-Reactivity:     | Mouse  |  |
| Predicted Reactivity: | Human,Rat,Dog,Cow,Sheep,Pig,Horse,Rabbit                   |  |
| Purification:         | Purified by Protein A.                                     |  |

#### Target Details

| Target:           | FAM65B                   |
|-------------------|--------------------------|
| Alternative Name: | FAM65B (FAM65B Products) |

#### Target Details

| Background:         | Synonyms: Ab2 162, C6orf32, DF48, FA65B_HUMAN, Fam65b, KIAA0386, PL48, Protein               |
|---------------------|--|
|                     | FAM65B.  |
|                     | Background: Making up nearly 6 % of the human genome, chromosome 6 contains around           |
|                     | 1,200 genes within 170 million base pairs of sequence. Deletion of a portion of the q arm of |
|                     | chromosome 6 is associated with early onset intestinal cancer suggesting the presence of a   |
|                     | cancer susceptibility locus. Porphyria cutanea tarda is associated with chromosome 6 through |
|                     | the HFE gene which, when mutated, predisposes an individual to developing this porphyria.    |
|                     | Notably, the PARK2 gene, which is associated with Parkinson's disease, and the genes         |
|                     | encoding the major histocompatiblity complex proteins, which are key molecular components    |
|                     | of the immune system and determine predisposition to rheumatic diseases, are also located on |
|                     | chromosome 6. Stickler syndrome, 21-hydroxylase deficiency and maple syrup urine disease     |
|                     | are also associated with genes on chromosome 6. A bipolar disorder susceptibility locus has  |
|                     | been identified on the q arm of chromosome 6. The C6orf32 gene product has been              |
|                     | provisionally designated C6orf32 pending further characterization.                           |
| Gene ID:            | 9750   |
| Dethuusus           | Transition Matellan Hamasatasia  |
| Pathways:           | Transition Metal Ion Homeostasis   |
| 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  |
| restrictions.       | 1 of Nescarott osc offly   |
| 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   |
|                     |  |

### Handling

| Storage Comment: | Store at -20°C. Aliquot into multiple vials to avoid repeated freeze-thaw cycles. |
|------------------|---|
| Expiry Date:     | 12 months   |