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# anti-FAM98A antibody (AA 251-350) (Alexa Fluor 350)



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| Quantity:            | 100 μL   |
|----------------------|--|
| Target:              | FAM98A   |
| Binding Specificity: | AA 251-350   |
| Reactivity:          | Human, Mouse   |
| Host:                | Rabbit   |
| Clonality:           | Polyclonal   |
| Conjugate:           | This FAM98A antibody is conjugated to Alexa Fluor 350  |
| Application:         | Flow Cytometry (FACS), Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

#### **Product Details**

| Immunogen:            | KLH conjugated synthetic peptide derived from human FAM98A |
|-----------------------|--|
| Isotype:              | IgG  |
| Cross-Reactivity:     | Human, Mouse   |
| Predicted Reactivity: | Rat,Cow,Sheep,Pig,Horse,Chicken                            |
| Purification:         | Purified by Protein A.                                     |

## Target Details

| Target:           | FAM98A                   |
|-------------------|--------------------------|
| Alternative Name: | FAM98A (FAM98A Products) |

## **Target Details**

| Background:         | Synonyms: FAM 98A, Family with sequence similarity 98 member A, Hypothetical protein                               |  |  |  |
|---------------------|--|--|--|--|
|                     | LOC25940, LOC25940, Protein FAM98A, FA98A_HUMAN.   |  |  |  |
|                     | Background: Encoding more than 700 genes, chromosome 15 is made up of approximately 106                            |  |  |  |
|                     | million base pairs and is about 3 % of the human genome. Angelman and Prader-Willi                                 |  |  |  |
|                     | syndromes are associated with loss of function or deletion of genes in the 15q11-q13 region. In                    |  |  |  |
|                     | the case of Angelman syndrome, this loss is due to inactivity of the maternal 15q11-q13                            |  |  |  |
|                     | encoded UBE3A gene in the brain by either chromosomal deletion or mutation. In cases of                            |  |  |  |
|                     | Prader-Willi syndrome, there is a partial or complete deletion of this region from the paternal                    |  |  |  |
|                     | copy of chromosome 15. Tay-Sachs disease is a lethal disorder associated with mutations of                         |  |  |  |
|                     | the HEXA gene, which is encoded by chromosome 15. Marfan syndrome is associated with                               |  |  |  |
|                     | chromosome 15 through the FBN1 gene. The FAM98 gene product has been provisionally                                 |  |  |  |
|                     | designated FAM98 pending further characterization.   |  |  |  |
| Gene ID:            | 25940  |  |  |  |
| Pathways:           | SARS-CoV-2 Protein Interactome   |  |  |  |
| Application Details |  |  |  |  |
| Application Notes:  | FCM 1:20-100   |  |  |  |
|                     | 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. |  |  |  |
| 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