# antibodies .- online.com





# anti-C7orf10 antibody (AA 351-445) (Alexa Fluor 555)



Go to Product page

| $\sim$ |    |    |    |             |   |
|--------|----|----|----|-------------|---|
| ()     | VE | ۲۱ | /1 | $\triangle$ | Λ |

| Quantity:            | 100 μL   |
|----------------------|--|
| Target:              | C7orf10  |
| Binding Specificity: | AA 351-445   |
| Reactivity:          | Human  |
| Host:                | Rabbit   |
| Clonality:           | Polyclonal   |
| Conjugate:           | This C7orf10 antibody is conjugated to Alexa Fluor 555   |
| 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 C7orf10 |
|-----------------------|---|
| Isotype:              | IgG   |
| Predicted Reactivity: | Human, Mouse, Rat, Dog, Cow, Sheep, Horse, Rabbit           |
| Purification:         | Purified by Protein A.                                      |

## **Target Details**

| Target:           | C7orf10  |
|-------------------|--|
| Alternative Name: | C7orf10 (C7orf10 Products)   |
| Background:       | Synonyms: Chromosome 7 open reading frame 10, Dermal papilla derived protein 13, DERP13, |

| FLJ11808, Hypothetica   | protein LOC79783, ORF19, Rus | ssel-Silver syndrome candidate, |
|-------------------------|------------------------------|---------------------------------|
| Uncharacterized protein | C7orf10,CG010_HUMAN.         |                                 |

Background: Chromosome 7 has been linked to Osteogenesis imperfecta, Pendred syndrome, Lissencephaly, Citrullinemia and Shwachman-Diamond syndrome. The deletion of a portion of the q arm of chromosome 7 is associated with Williams-Beuren syndrome, a condition characterized by mild mental retardation, an unusual comfort and friendliness with strangers and an elfin appearance. Deletions of portions of the q arm of chromosome 7 are also seen in a number of myeloid disorders including cases of acute myelogenous leukemia and myelodysplasia. The C7orf10 gene product has been provisionally designated C7orf10 pending further characterization.

Gene ID:

79783

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