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Datasheet for ABIN5007871

**anti-Neurensin 1 antibody (AA 101-195) (Alexa Fluor 680)**

## Overview

|                      |   |
|----------------------|---|
| Quantity:            | 100 µL  |
| Target:              | Neurensin 1 (NRSN1)   |
| Binding Specificity: | AA 101-195  |
| Reactivity:          | Human   |
| Host:                | Rabbit  |
| Clonality:           | Polyclonal  |
| Conjugate:           | This Neurensin 1 antibody is conjugated to Alexa Fluor 680  |
| Application:         | Immunofluorescence (Cultured Cells) (IF (cc)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)) |

## Product Details

|                       |   |
|-----------------------|---|
| Immunogen:            | KLH conjugated synthetic peptide derived from human NRSN1 |
| Isotype:              | IgG   |
| Predicted Reactivity: | Human,Mouse,Rat,Horse,Rabbit                              |
| Purification:         | Purified by Protein A.                                    |

## Target Details

|                   |   |
|-------------------|---|
| Target:           | Neurensin 1 (NRSN1)   |
| Alternative Name: | NRSN1 ( <a href="#">NRSN1 Products</a> )  |
| Background:       | Synonyms: Neurensin-1, Neuro-p24, NRSN1, NRSN1_HUMAN, p24, Vesicular membrane protein |

## Target Details

of 24 kDa, Vesicular membrane protein p24, VMP.

Background: Neurensin-1 (NRSN1), also designated Vesicular membrane protein of 24 kDa (VMP) or Neuro-p24, is a 195 amino acid multi-pass membrane protein belonging to the VMP family that is involved in the transport of neural organelle transport and in the transduction of nerve signals or in nerve growth. Expressed solely in brain, Neurensin-1 is also thought to play a role in neurite extension. The gene encoding Neurensin-2 maps to human chromosome 6, which 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, Parkinson's disease, Stickler syndrome, 21-hydroxylase deficiency and maple syrup urine disease are also associated with genes on chromosome 6.

## 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