

Datasheet for ABIN7599042

anti-CCDC115 antibody (AA 1-180)



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| Quantity: | 100 μg | |
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| Target: | CCDC115 | |
| Binding Specificity: | AA 1-180 | |
| Reactivity: | Human | |
| Host: | Rabbit | |
| Clonality: | Polyclonal | |
| Conjugate: | This CCDC115 antibody is un-conjugated | |
| Application: | Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Flow Cytometry (FACS), Immunocytochemistry (ICC) | |

Product Details

| Purpose: | Anti-CCDC115 Antibody Picoband® | |
|-----------------------------|--|--|
| Immunogen: | E.coli-derived human CCDC115 recombinant protein (Position: M1-A180). | |
| Isotype: | IgG | |
| Cross-Reactivity (Details): | No cross-reactivity with other proteins | |
| Characteristics: | Anti-CC Antibody Picoband® (ABIN7599042). Tested in ELISA, Flow Cytometry, IF, IHC, ICC, WB applications. This antibody reacts with Human. The brand Picoband indicates this is a premium antibody that guarantees superior quality, high affinity, and strong signals with minimal background in Western blot applications. Only our best-performing antibodies are designated as Picoband, ensuring unmatched performance. | |
| Purification: | Immunogen affinity purified. | |

| Target Details | |
|---------------------|--|
| Target: | CCDC115 |
| Alternative Name: | CCDC115 (CCDC115 Products) |
| Background: | Synonyms: RNA-binding protein 47,RNA-binding motif protein 47,RBM47, Tissue Specificity: Abundantly expressed in tonsil, lymph node, and trachea, strong expression in prostate, lower expression in thyroid, stomach, and colon. Background: Coiled-coil domain containing 115 is a protein that in humans is encoded by the CCDC115 gene. The protein encoded by this gene has been observed to localize to the endoplasmic reticulum (ER)-Golgi intermediate compartment (ERGIC) and coat protein complex I (COPI) vesicles in some human cells. The encoded protein shares some homology with the yeast V-ATPase assembly factor Vma22p, and the orthologous protein in mouse promotes cell proliferation and suppresses cell death. Defects in this gene are a cause of congenital disorder of glycosylation, type IIo in humans. |
| Molecular Weight: | 20 kDa |
| Gene ID: | 84317 |
| Application Details | |
| Application Notes: | Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human Immunocytochemistry/Immunofluorescence, 5 μg/mL, Human |

Flow Cytometry (Fixed), 1-3 µg/1x10⁶ cells, Human

ELISA, 0.1-0.5 µg/mL, -

1. D., Krawitz, P., Wada, Y., Ashikov, A., Perez-Cerda, C., and 31 others. CCDC115 deficiency

causes a disorder of Golgi homeostasis with abnormal protein glycosylation. Am. J. Hum.

Genet. 98: 310-321, 2016. 2. Pellicano, F., Inglis-Broadgate, S. L., Pante, G., Ansorge, W., Iwata, T.

Expression of coiled-coil protein 1, a novel gene downstream of FGF2, in the developing brain.

Gene Expr. Patterns 6: 285-293, 2006. 3. Pellicano, F., Thomson, R. E., Inman, G. J., Iwata, T.

Regulation of cell proliferation and apoptosis in neuroblastoma cells by ccp1, a FGF2

downstream gene. BMC Cancer 10: 657, 2010. Note: Electronic Article.

Restrictions:

For Research Use only

Handling

| Format: | Lyophilized |
|-----------------|---|
| Reconstitution: | Adding 0.2 mL of distilled water will yield a concentration of 500 μg/mL. |

Handling

| Concentration: | 500 μg/mL |
|------------------|--|
| Buffer: | Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na2HPO4. |
| Storage: | 4 °C,-20 °C |
| Storage Comment: | At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and thawing. |