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Datasheet for ABIN1711758 anti-ZNF503 antibody (AA 41-140) (HRP)



Overview

| Quantity: | 100 μL |
|----------------------|--|
| Target: | ZNF503 |
| Binding Specificity: | AA 41-140 |
| Reactivity: | Mouse |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This ZNF503 antibody is conjugated to HRP |
| Application: | Western Blotting (WB), ELISA, Immunohistochemistry (Frozen Sections) (IHC (fro)), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)) |

Product Details

| Immunogen: | KLH conjugated synthetic peptide derived from human ZNF503/NOLZ1 |
|-----------------------|--|
| Isotype: | lgG |
| Cross-Reactivity: | Mouse |
| Predicted Reactivity: | Human,Rat,Dog,Cow,Pig,Horse,Chicken |
| Purification: | Purified by Protein A. |
| Target Details | |
| | |

| Target: | ZNF503 |
|-------------------|--------------------------------|
| Alternative Name: | ZNF503/NOLZ1 (ZNF503 Products) |

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| Background: | Synonyms: FLJ45745, MGC2555, NOLZ 1, NOLZ1, Zinc finger protein 503, ZN503_HUMAN, znf503. |
|---------------------|--|
| | |
| | Background: Nolz 1 is a 646 amino acid nuclear protein that is thought to function as a |
| | transcriptional repressor and is highly expressed in developing striatum. Additionally, Nolz-1 |
| | has been suggested to play a role in neural differentiation. A member of the Elbow/Noc family, |
| | Nolz-1 exists as three alternatively spliced isoforms and contains one C2H2-type zinc finger. |
| | The gene encoding Nolz-1 maps to human chromosome 10, which makes up approximately |
| | 4.5 % of total DNA in cells and encodes nearly 1,200 genes. Several protein-coding genes, |
| | including those that encode for chemokines, cadherins, excision repair proteins, early growth |
| | response factors (Egrs) and fibroblast growth receptors (FGFRs), are located on chromosome |
| | 10. Defects in some of the genes that map to chromosome 10 are associated with Charcot- |
| | Marie Tooth disease, Jackson-Weiss syndrome, Usher syndrome, nonsyndromatic deafness, |
| | Wolman?s syndrome, Cowden syndrome, multiple endocrine neoplasia type 2 and porphyria. |
| Gene ID: | 84858 |
| UniProt: | Q96F45 |
| Pathways: | SARS-CoV-2 Protein Interactome |
| Application Details | |
| Application Notes: | WB 1:300-5000 |
| | IHC-P 1:200-400 |
| | IHC-F 1:100-500 |
| 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. |
| | |

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Handling

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