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Datasheet for ABIN7299632 anti-Anoctamin 7 antibody (C-Term)

2 Images



Overview

| Quantity: | 100 μL |
|----------------------|--|
| Target: | Anoctamin 7 (ANO7) |
| Binding Specificity: | C-Term |
| Reactivity: | Human, Mouse |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This Anoctamin 7 antibody is un-conjugated |
| Application: | Western Blotting (WB), Immunohistochemistry (IHC) |
| Product Details | |
| Immunogen: | KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of |
| | human NGEP. |
| Specificity: | Recognizes endogenous levels of NGEP protein. |
| Characteristics: | Rabbit polyclonal antibody to NGEP |
| Purification: | The antibody was purified by immunogen affinity chromatography. |

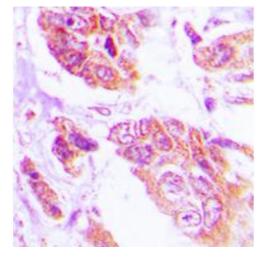
Target Details

| Target: | Anoctamin 7 (ANO7) |
|-------------------|--|
| Alternative Name: | NGEP (ANO7 Products) |
| Background: | NGEP, PCANAP5, TMEM16G, Anoctamin-7, Dresden transmembrane protein of the prostate, D- |

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| Target Details | |
|---------------------|--|
| | TMPP, IPCA-5, New gene expressed in prostate, Prostate cancer-associated protein 5, Transmembrane protein 16G |
| Gene ID: | 50636, 404545 |
| UniProt: | Q6IWH7, Q14AT5 |
| Application Details | |
| Application Notes: | WB (1:500 - 1:1000), IH (1:100 - 1:200) |
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Liquid |
| Buffer: | Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide. |
| Preservative: | Sodium azide |
| Precaution of Use: | This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. |
| Storage: | -20 °C |
| Storage Comment: | Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles. |
| Expiry Date: | 12 months |

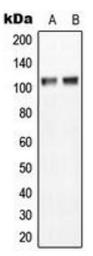
Images



Immunohistochemistry

Image 1. Immunohistochemical analysis of NGEP staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with

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haematoxylin and mounted with DPX.

Western Blotting

Image 2. Western blot analysis of NGEP expression in MCF7 (A), NIH3T3 (B) whole cell lysates.

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