antibodies

Datasheet for ABIN6258535 anti-Aquaporin 12A antibody (C-Term)

2 Images



Overview

| Quantity: | 100 µL |
|----------------------|--|
| Target: | Aquaporin 12A (AQP12A) |
| Binding Specificity: | C-Term |
| Reactivity: | Human |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Application: | Western Blotting (WB), ELISA, Immunocytochemistry (ICC), Immunofluorescence (IF) |
| Product Details | |
| Immunogen: | A synthesized peptide derived from human AQP12, corresponding to a region within C-terminal amino acids. |
| Isotype: | lgG |
| Specificity: | AQP12 Antibody detects endogenous levels of total AQP12. |
| Purification: | The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling |

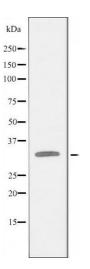
Target Details

| Target: | Aquaporin 12A (AQP12A) |
|-------------------|---|
| Alternative Name: | AQP12A,AQP12B (AQP12A Products) |
| Background: | Description: Aquaporins facilitate the transport of water and small neutral solutes across cell |
| | membranes. |

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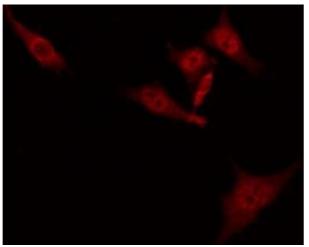
| Target Details | |
|---------------------|--|
| | Gene: AQP12A |
| Molecular Weight: | 34 kDa |
| Gene ID: | 375318, 653437 |
| UniProt: | Q8IXF9, A6NM10 |
| Application Details | |
| Application Notes: | WB 1:500-1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000 |
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Liquid |
| Concentration: | 1 mg/mL |
| Buffer: | Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol. |
| 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: | Store at -20 °C. Stable for 12 months from date of receipt. |
| Expiry Date: | 12 months |

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Western Blotting

Image 1. Western blot analysis of extracts from Jurkat cells, using AQP12 antibody.



Immunofluorescence (fixed cells)

Image 2. ABIN6274860 staining Hela by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.

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