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Product Datasheet

Anti-Arl13b Antibody FL594 Conjugate



KO Validated

Overview

| Catalog # | 75-287-FL594 |
|--------------------------------|---|
| - | |
| Conjugate | FL594 Ex: 594 nm, Em: 615 nm |
| lsotype | IgG2a |
| Clone Number | N295B/66 |
| Size | 200 μL |
| Concentration | 0.5 mg/mL |
| Host Species | Mouse Monoclonal |
| Format | Purified by Protein A chromatography |
| Buffer | PBS with 0.09% azide |
| Applications | ICC, IHC |
| Species Reactivity | Human, Mouse, Non-Human Primate, Rat, and Zebrafish |
| Immunogen | Fusion protein amino acids 208-427 (C-terminus) of mouse Arl13b (accession number Q640N2) produced recombinantly in E. Coli |
| Molecular Weight | 50 kDa |
| Cite this Antibody | Antibodies Inc Cat# 75-287-FL594, RRID: AB_2940026 |
| | |
| | |
| Details | |
| Details Target Description | ADP-ribosylation factor-like protein 13B (ARL13B), also known as ADP-ribosylation factor-like protein 2-like 1. Arl13B is a small GTPase with both N-terminal and C-terminal guanine nucleotide- binding motifs. Arl13B is localized to the membrane of cilia and expressed in Neuroepithelial cells and developing radial glia of the developing cerebral cortex. Mutations in this gene are the cause of Joubert syndrome 8. |
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| Target Description Specificity | protein 2-like 1. Arl13B is a small GTPase with both N-terminal and C-terminal guanine nucleotide- binding motifs. Arl13B is localized to the membrane of cilia and expressed in Neuroepithelial cells and developing radial glia of the developing cerebral cortex. Mutations in this gene are the cause of Joubert syndrome 8. Does not cross-react with Arl13a (based on KO validation results) Produced by in vitro bioreactor culture of hybridoma line followed by Protein A affinity |

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