



MALD2 Polyclonal Antibody

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| Catalog No | BYab-05709 |
| Isotype | IgG |
| Reactivity | Human;Mouse |
| Applications | WB;ELISA |
| Gene Name | MARVELD2 TRIC |
| Protein Name | MARVEL domain-containing protein 2 (Tricellulin) |
| Immunogen | Synthesized peptide derived from human protein . at AA range: 170-250 |
| Specificity | MALD2 Polyclonal Antibody detects endogenous levels of protein. |
| Formulation | Liquid in PBS containing 50% glycerol, and 0.02% sodium azide. |
| Source | Polyclonal, Rabbit,IgG |
| Purification | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen. |
| Dilution | WB 1:500-2000 ELISA 1:5000-20000 |
| Concentration | 1 mg/ml |
| Purity | ≥90% |
| Storage Stability | -20°C/1 year |
| Synonyms | |
| Observed Band | 61kD |
| Cell Pathway | Cell membrane ; Multi-pass membrane protein . Cell junction, tight junction . Located at tricellular contacts. . |
| Tissue Specificity | Brain,Epithelium,Lung,Trachea, |
| Function | disease:Defects in MARVELD2 are the cause of non-syndromic sensorineural deafness autosomal recessive type 49 (DFNB49) [MIM:610153]. DFNB49 is a form of sensorineural hearing loss. Sensorineural deafness results from damage to the neural receptors of the inner ear, the nerve pathways to the brain, or the area of the brain that receives sound information.,function:Plays a role in the formation of the epithelial barriers. The separation of the endolymphatic and perilymphatic spaces of the organ of Corti from one another by epithelial barriers is required for normal hearing.,PTM:Phosphorylated upon DNA damage, probably by ATM or ATR.,similarity:Contains 1 MARVEL domain.,subcellular location:Found at tricellular contacts., |

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Background

The protein encoded by this gene is a membrane protein found at the tight junctions between epithelial cells. The encoded protein helps establish epithelial barriers such as those in the organ of Corti, where these barriers are required for normal hearing. Defects in this gene are a cause of deafness autosomal recessive type 49 (DFNB49). Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2011],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images