

TMEM132E Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP11487b

Product Information

Application WB, FC, E **Primary Accession** QGIEE7

Other Accession Q6IEE6, NP_997196.1
Reactivity Human, Mouse

Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG
Clone Names RB29313
Calculated MW 116150
Antigen Region 704-733

Additional Information

Gene ID 124842

Other Names Transmembrane protein 132E, TMEM132E

Target/Specificity This TMEM132E antibody is generated from rabbits immunized with a KLH

conjugated synthetic peptide between 704-733 amino acids from the

C-terminal region of human TMEM132E.

Dilution WB~~1:1000 FC~~1:10~50 E~~Use at an assay dependent concentration.

Format Purified polyclonal antibody supplied in PBS with 0.05% (V/V) Proclin 300. This

antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation

followed by dialysis against PBS.

Storage Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions TMEM132E Antibody (C-term) is for research use only and not for use in

diagnostic or therapeutic procedures.

Protein Information

Name TMEM132E (<u>HGNC:26991</u>)

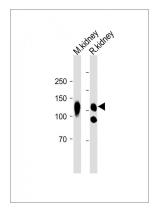
Function Required for normal inner ear hair cell function and hearing.

Cellular Location Membrane; Single-pass type I membrane protein

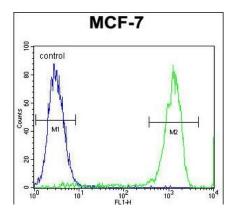
References

Kehrer-Sawatzki, H., et al. J. Med. Genet. 40 (10), E116 (2003):

Images



All lanes: Anti-TMEM132E Antibody (C-term) at 1:1000 dilution Lane 1: Mouse kidney lysate Lane 2: Rat kidney lysate Lysates/proteins at 20 µg per lane. Secondary: Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated (ASP1615) at 1/15000 dilution. Observed band size: 116 KDa Blocking/Dilution buffer: 5% NFDM/TBST.



TMEM132E Antibody (C-term) (Cat. #AP11487b) flow cytometric analysis of MCF-7 cells (right histogram) compared to a negative control cell (left histogram).FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

Citations

• Whole-exome sequencing identifies a variant in TMEM132E causing autosomal-recessive nonsyndromic hearing loss DFNB99.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.