# **OTOF Rabbit pAb**

Catalog No.: A20266 1 Publications



## **Basic Information**

#### **Observed MW**

Refer to figures

#### **Calculated MW**

227kDa

#### Category

Primary antibody

#### **Applications**

WB,ELISA

#### **Cross-Reactivity**

Human

## **Background**

Mutations in this gene are a cause of neurosensory nonsyndromic recessive deafness, DFNB9. The short form of the encoded protein has 3 C2 domains, a single carboxy-terminal transmembrane domain found also in the C. elegans spermatogenesis factor FER-1 and human dysferlin, while the long form has 6 C2 domains. The homology suggests that this protein may be involved in vesicle membrane fusion. Several transcript variants encoding multiple isoforms have been found for this gene.

## **Recommended Dilutions**

**WB** 1:500 - 1:2000

**ELISA** 

Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

## **Immunogen Information**

**Gene ID**9381

Swiss Prot
Q9HC10

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-300 of human OTOF (NP\_919224.1).

## **Synonyms**

AUNB1; DFNB6; DFNB9; NSRD9; FER1L2; OTOF

## **Contact**

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## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

Store at -20  $^{\circ}\text{C}.$  Avoid freeze / thaw cycles.

Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3.