

# MERTK Rabbit pAb

MERTK Rabbit pAb Catalog # AP94184

#### **Product Information**

**Application** WB, E Reactivity Mouse Host Rabbit Clonality Polyclonal **Calculated MW** 108 KDa **Physical State** Liquid

**Immunogen** Recombinant mouse MERTK protein

**Epitope Specificity** 23-497/994

Isotype IgG

**Purity** affinity purified by Protein A

**Buffer** SUBCELLULAR LOCATION

**SIMILARITY** 

**SUBUNIT** 

Post-translational modifications

**DISEASE** 

**Important Note** 

**Background Descriptions** 

0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

Membrane; Single-pass type I membrane protein.

Belongs to the protein kinase superfamily. Tyr protein kinase family, AXL/UFO subfamily. Contains 2 fibronectin type-III domains. Contains 2 Ig-like C2-type

(immunoglobulin-like) domains. Contains 1 protein kinase domain. Interacts (upon activation) with TNK2; stimulates TNK2 autophosphorylation.

Interacts (via N-terminus) with extracellular ligands LGALS3, TUB, TULP1 and GAS6. Interacts with VAV1 in a phosphotyrosine-independent manner. Autophosphorylated on Tyr-744, Tyr-748 and Tyr-749 in the activation loop

allowing full activity. Autophosphorylated on Tyr-867 leading to recruitment of downstream partners of the signaling cascade such as PLCG2.

Defects in MERTK are the cause of retinitis pigmentosa type 38 (RP38) [MIM:613862]. RP38 is a retinal dystrophy belonging to the group of pigmentary retinopathies. Retinitis pigmentosa is characterized by retinal pigment deposits visible on fundus examination and primary loss of rod photoreceptor cells followed by secondary loss of cone photoreceptors. Patients typically have night vision blindness and loss of midperipheral visual field. As their condition progresses, they lose their far peripheral visual field

and eventually central vision as well.

This product as supplied is intended for research use only, not for use in

human, therapeutic or diagnostic applications.

The Major Facilitator Superfamily (MFS) is a large and diverse group of secondary transporters that includes uniporters, symporters, and antiporters.

MFS proteins facilitate the transport across cytoplasmic or internal

membranes of a variety of substrates including ions, sugar phosphates, drugs, neurotransmitters, nucleosides, amino acids, and peptides. They do so using the electrochemical potential of the transported substrates. Uniporters transport a single substrate, while symporters and antiporters transport two substrates in the same or in opposite directions, respectively, across membranes. Peptide-transporters 2 [solute carrier family 15 (H+/peptide transporter), member 2; SLC15A2; PEPT2; Oligopeptide transporter, kidney isoform; Kidney H(+)/peptide cotransporter; ].

#### **Additional Information**

Target/Specificity Expressed predominantly in the hematopoietic lineages: macrophages, NK

cells, NKT cells, dendritic cells and platelets.

**Dilution** WB=1:500-2000,ELISA=1:5000-10000

Format 0.01M TBS(pH7.4) with 1% BSA, 0.09% (W/V) sodium azide and 50% Glyce

**Storage** Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When

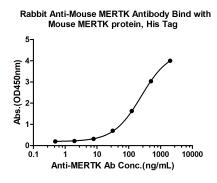
reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody

is stable for at least two weeks at 2-4 °C.

## **Background**

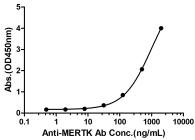
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### **Images**

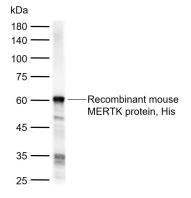


Measured by its binding ability in a indirect ELISA. Immobilized Mouse MERTK protein, His Tag (Cat. bs-41398P) at 2  $\mu$ g/mL (100  $\mu$ L/well) can bind Rabbit Anti-Mouse MERTK Antibody, the EC50 is 246.3 ng/mL.





Measured by its binding ability in a indirect ELISA. Immobilized Mouse MERTK protein, His Tag (Cat. bs-41398P) at 2  $\mu$ g/mL (100  $\mu$ L/well) can bind Rabbit Anti-Mouse MERTK Antibody, the EC50 is 992.7 ng/mL.



Sample: Lane 1: Recombinant mouse MERTK protein, His Primary: Anti-MERTK (AP94184) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: kDa Observed band size: 60 kDa

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