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**Research Use Only. Not for
diagnostic or therapeutic use.**

EB07296 - Goat Anti-Silver homologue / Pmel 17 (C-terminus) Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: silver homologue (mouse), PMEL17, SILV, D12S53E, ME20, SI, SIL, gp100, Melanocyte protein mel 17, Pmel 17, Silver, mouse, homolog of, silver (mouse homologue) like, silver (mouse homologue)-like, silver homologue

Official Symbol: PMEL

Accession Number(s): NP_008859.1

Human GeneID(s): [6490](#)

Non-Human GeneID(s): 20431 (mouse), 362818 (rat)

Immunogen

Peptide with sequence CPIGENSPLLSGQQ, from the C Terminus of the protein sequence according to NP_008859.1.

Please note the [peptide](#) is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:16000.

Western blot: Approx 70kDa band observed in Mouse Skin lysates (calculated MW of 66.3kDa according to Mouse NP_068682.2). Recommended concentration: 1-3µg/ml. Primary incubation was 1 hour.

Immunofluorescence: Strong expression of the protein seen in the plasma membrane and cytoplasm of A431 cells. Recommended concentration: 10µg/ml.

Species Reactivity

Tested: Human, Mouse

Expected from sequence similarity: Human, Mouse

Specific Reference

This antibody (previous batch) has been successfully used in Western blot on Zebrafish:

Liu C, Yan W, Zhou B, Guo Y, Liu H, Yu H, Giesy JP, Wang J, Li G, Zhang X.

Characterization of a bystander effect induced by the endocrine-disrupting chemical 6-propyl-2-thiouracil in zebrafish embryos.

Aquat Toxicol. 2012 Aug 15;118-119:108-15.

PMID: 22542736

EB07296 (1µg/ml) staining of Mouse Skin lysate (35µg protein in RIPA buffer). Detected by chemiluminescence.

EB07296 Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing membrane and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).