

ADAM17 Antibody (C-term)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP1492b

Specification

ADAM17 Antibody (C-term) - Product Information

Application WB,E
Primary Accession P78536
Reactivity Human
Host Rabbit
Clonality Polyclonal
Isotype Rabbit IgG
Antigen Region 794-823

ADAM17 Antibody (C-term) - Additional Information

Gene ID 6868

Other Names

Disintegrin and metalloproteinase domain-containing protein 17, ADAM 17, Snake venom-like protease, TNF-alpha convertase, TNF-alpha-converting enzyme, CD156b, ADAM17, CSVP, TACE

Target/Specificity

This ADAM17 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 794-823 amino acids from the C-terminal region of human ADAM17.

Dilution

WB~~1:1000

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

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

ADAM17 Antibody (C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

ADAM17 Antibody (C-term) - Protein Information

Name ADAM17

Synonyms CSVP, TACE

Function Cleaves the membrane-bound precursor of TNF-alpha to its mature soluble form



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(PubMed:<u>9034191</u>). Responsible for the proteolytical release of soluble JAM3 from endothelial cells surface (PubMed:<u>20592283</u>). Responsible for the proteolytic release of several other cell-surface proteins, including p75 TNF-receptor, interleukin 1 receptor type II, p55 TNF-receptor, transforming growth factor-alpha, L-selectin, growth hormone receptor, MUC1 and the amyloid precursor protein (PubMed:<u>12441351</u>). Acts as an activator of Notch pathway by mediating cleavage of Notch, generating the membrane-associated intermediate fragment called Notch extracellular truncation (NEXT) (PubMed:<u>24226769</u>). Plays a role in the proteolytic processing of ACE2 (PubMed:<u>24227843</u>). Plays a role in hemostasis through shedding of GP1BA, the platelet glycoprotein Ib alpha chain (By similarity). Mediates the proteolytic cleavage of LAG3, leading to release the secreted form of LAG3 (By similarity). Mediates the proteolytic cleavage of IL6R, leading to the release of secreted form of IL6R (PubMed:<u>26876177</u>, PubMed:<u>28060820</u>). Mediates the proteolytic cleavage and shedding of FCGR3A upon NK cell stimulation, a mechanism that allows for increased NK cell motility and detachment from opsonized target cells.

Cellular Location

Membrane; Single-pass type I membrane protein.

Tissue Location

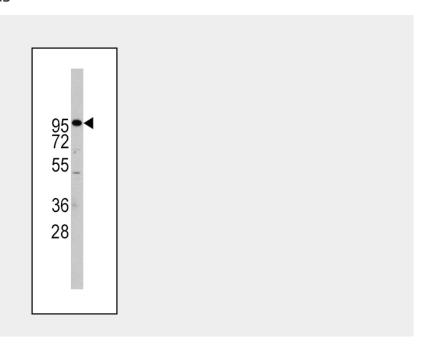
Ubiquitously expressed. Expressed at highest levels in adult heart, placenta, skeletal muscle, pancreas, spleen, thymus, prostate, testes, ovary and small intestine, and in fetal brain, lung, liver and kidney. Expressed in natural killer cells (at protein level) (PubMed:24337742).

ADAM17 Antibody (C-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

ADAM17 Antibody (C-term) - Images





Western blot analysis of ADAM17 Antibody (N-term) in Jurkat cell line lysates (35ug/lane).

ADAM17 Antibody (C-term) - Background

ADAM17(arrow) was detected using the purified Pab.

ADAM17 is a member of the ADAM (a disintegrin and metalloprotease domain) family. Members of this family are membrane-anchored proteins structurally related to snake venom disintegrins, and have been implicated in a variety of biologic processes involving cell-cell and cell-matrix interactions, including fertilization, muscle development, and neurogenesis. ADAM17 functions as a tumor necrosis factor-alpha converting enzyme; binds mitotic arrest deficient 2 protein; and also plays a prominent role in the activation of the Notch signaling pathway.

ADAM17 Antibody (C-term) - References

Tellier, E., J. Cell. Physiol. 214 (3), 687-693 (2008) Takamune, Y., Biochem. Biophys. Res. Commun. 365 (2), 393-398 (2008)