

## Anti-ADAM9 antibody



**Description** Unconjugated Rabbit polyclonal to ADAM9

Model STJ190515

**Host** Rabbit

**Reactivity** Human

**Applications** ELISA, WB

Immunogen Synthesized peptide derived from human ADAM9 protein.

**Immunogen Region** 120-200aa

**Gene ID** <u>8754</u>

Gene Symbol ADAM9

**Dilution range** WB 1:500-2000 ELISA 1:5000-20000

**Specificity** ADAM9 Polyclonal Antibody detects endogenous levels of protein.

**Tissue Specificity** Widely expressed. Expressed in chondrocytes. Isoform 2 is highly expressed

in liver and heart.

**Purification** ADAM9 antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

**Note** For Research Use Only (RUO).

**Protein Name** Disintegrin and metalloproteinase domain-containing protein 9 ADAM 9

Cellular disintegrin-related protein Meltrin-gamma

Metalloprotease/disintegrin/cysteine-rich protein 9 Myeloma cell

metalloproteinase

Molecular Weight 90 kDa

**Clonality** Polyclonal

**Conjugation** Unconjugated

**Isotype** IgG

**Formulation** Liquid form in PBS containing 50% glycerol, and 0.02% sodium azide.

**Concentration** 1 mg/ml

**Storage Instruction** Store at -20°C, and avoid repeat freeze-thaw cycles.

Database Links HGNC:2160MIM:602713

Alternative Names Disintegrin and metalloproteinase domain-containing protein 9 ADAM 9

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metalloproteinase

**Function** Cleaves and releases a number of molecules with important roles in

tumorigenesis and angiogenesis, such as TEK, KDR, EPHB4, CD40, VCAM1 and CDH5. May mediate cell-cell, cell-matrix interactions and regulate the motility of cells via interactions with integrins. Isoform 2: May act as alpha-

secretase for amyloid precursor protein (APP).

**Cellular Localization** Isoform 1: Cell membrane Isoform 2: Secreted

**Post-translational** Proteolytically cleaved in the trans-Golgi network before it reaches the plasma

Modifications membrane to generate a mature protein. The removal of the pro-domain

occurs via cleavage at two different sites. Processed most likely by a proprotein convertase such as furin, at the boundary between the pro-domain and the catalytic domain. An additional upstream cleavage pro-protein convertase site (Arg-56/Glu-57) has an important role in the activation of ADAM9. Phosphorylation is induced in vitro by phorbol-12-myristate-13-acetate

(PMA).

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