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# MMP17 Antibody (Center)

**Purified Rabbit Polyclonal Antibody (Pab)**

**Catalog # AP6201a**

## Specification

### MMP17 Antibody (Center) - Product Information

| | |
|---|---|
| Application | **IHC-P, WB,E** |
| Primary Accession | Q9ULZ9 |
| Other Accession | Q9R0S3, NP_057239 |
| Reactivity | **Human** |
| Predicted | **Mouse** |
| Host | **Rabbit** |
| Clonality | **Polyclonal** |
| Isotype | **Rabbit IgG** |
| Calculated MW | **66653** |
| Antigen Region | **391-420** |

### MMP17 Antibody (Center) - Additional Information

**Gene ID** 4326

**Other Names**
Matrix metalloproteinase-17, MMP-17, 3424-, Membrane-type matrix metalloproteinase 4, MT-MMP 4, MTMMP4, Membrane-type-4 matrix metalloproteinase, MT4-MMP, MT4MMP, MMP17, MT4MMP

**Target/Specificity**
This MMP17 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 391-420 amino acids from the Central region of human MMP17.

**Dilution**
IHC-P~~1:50~100
WB~~1:1000
E~~Use at an assay dependent concentration.

**Format**
Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.

**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**
MMP17 Antibody (Center) is for research use only and not for use in diagnostic or therapeutic procedures.

### MMP17 Antibody (Center) - Protein Information

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**Name** MMP17

**Synonyms** MT4MMP

**Function** Endopeptidase that degrades various components of the extracellular matrix, such as fibrin. May be involved in the activation of membrane-bound precursors of growth factors or inflammatory mediators, such as tumor necrosis factor-alpha. May also be involved in tumoral process. Cleaves pro-TNF-alpha at the '74-Ala-|-Gln-75' site. Not obvious if able to proteolytically activate progelatinase A. Does not hydrolyze collagen types I, II, III, IV and V, gelatin, fibronectin, laminin, decorin nor alpha1-antitrypsin.

**Cellular Location**
[Isoform Long]: Cell membrane; Lipid-anchor, GPI- anchor; Extracellular side. Secreted, extracellular space, extracellular matrix

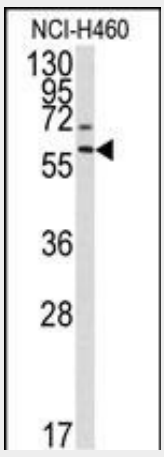
**Tissue Location**
Expressed in brain, leukocytes, colon, ovary testis and breast cancer. Expressed also in many transformed and non- transformed cell types

## MMP17 Antibody (Center) - 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

## MMP17 Antibody (Center) - Images




Western blot analysis of anti-MMP17 Antibody (Center) (Cat.#AP6201a) in NCI-H460 cell line lysates (35ug/lane). MMP17(arrow) was detected using the purified Pab.

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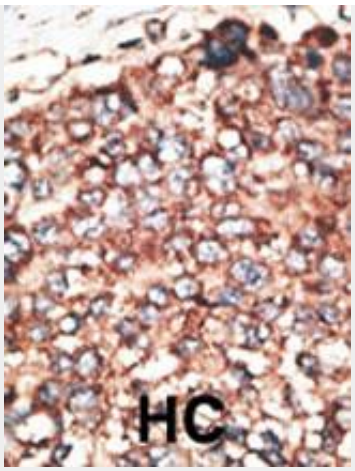


Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated. BC = breast carcinoma; HC = hepatocarcinoma.

**MMP17 Antibody (Center) - Background**

Proteins of the matrix metalloproteinase (MMP) family are involved in the breakdown of extracellular matrix in normal physiological processes, such as embryonic development, reproduction, and tissue remodeling, as well as in disease processes, such as arthritis and metastasis. Most MMPs are secreted as inactive proproteins which are activated when cleaved by extracellular proteinases. MMP17 is considered a member of the membrane-type MMP (MT-MMP) subfamily. However, this protein is unique among the MT-MMPs in that it is a GPI-anchored protein rather than a transmembrane protein. The protein activates MMP-2 by cleavage.

**MMP17 Antibody (Center) - References**

Jung, M., et al., Prostate 55(2):89-98 (2003).
Itoh, Y., et al., J. Biol. Chem. 274(48):34260-34266 (1999).
Wang, Y., et al., J. Biol. Chem. 274(46):33043-33049 (1999).
Kajita, M., et al., FEBS Lett. 457(3):353-356 (1999).
Nagase, H., et al., J. Biol. Chem. 274(31):21491-21494 (1999).