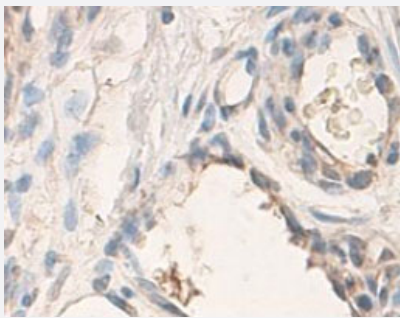


# ADAMTS3 polyclonal antibody

Catalog # PAB18818      Size 100 ug

## Applications



### Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of formalin-fixed paraffin-embedded human skin tissue with ADAMTS3 polyclonal antibody (Cat # PAB18818) at 1 : 100 dilution.

## Specification

<b>Product Description</b>	Rabbit polyclonal antibody raised against synthetic peptide of ADAMTS3.
<b>Immunogen</b>	A synthetic peptide corresponding to 16 amino acids at C-terminus of human ADAMTS3.
<b>Host</b>	Rabbit
<b>Reactivity</b>	Human
<b>Form</b>	Liquid
<b>Purification</b>	Peptide affinity purification
<b>Recommend Usage</b>	ELISA (1:5000-1:20000) Immunohistochemistry (1:100-1:500) The optimal working dilution should be determined by the end user.
<b>Storage Buffer</b>	In buffer containing 0.02% sodium azide
<b>Storage Instruction</b>	Store at 4°C for three months. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
<b>Note</b>	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

## Applications

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining of formalin-fixed paraffin-embedded human skin tissue with ADAMTS3 polyclonal antibody (Cat # PAB18818) at 1 : 100 dilution.

- Enzyme-linked Immunoabsorbent Assay

## Gene Info — ADAMTS3

Entrez GeneID [9508](#)

Gene Name ADAMTS3

Gene Alias ADAMTS-4, KIAA0366

Gene Description ADAM metalloproteinase with thrombospondin type 1 motif, 3

Omim ID [605011](#)

Gene Ontology [Hyperlink](#)

**Gene Summary** This gene encodes a member of the ADAMTS (a disintegrin and metalloproteinase with thrombospondin motifs) protein family. Members of the family share several distinct protein modules, including a propeptide region, a metalloproteinase domain, a disintegrin-like domain, and a thrombospondin type 1 (TS) motif. Individual members of this family differ in the number of C-terminal TS motifs, and some have unique C-terminal domains. The protein encoded by this gene is the major procollagen II N-proteinase. A deficiency of this protein may be responsible for dermatosparaxis, a genetic defect of connective tissues. [provided by RefSeq]

**Other Designations** a disintegrin-like and metalloproteinase (reprolysin type) with thrombospondin type 1 motif, 3|zinc metalloendopeptidase

## Disease

- [Genetic Predisposition to Disease](#)
- [Osteoarthritis](#)