## **COL1A1 Propeptide Antibody**

**Tested Species Reactivity** Human (Hu) Mouse (Ms) Rat (Rt)

Tested Applications	Dilution *
Western Blot (WB)	1:1000
Immunofluorescence (IF)	1:100
Immunocytochemistry (ICC)	1:100
Immunohistochemistry (IHC)	1:100

Suggested working dilutions are given as a guide only. It is recommended that the user titrates the product for use in their own experiment using appropriate negative and positive controls

Lot Number: RJ2281701

**Product Data Sheet** 

Details	
Catalog Number:	PA5-35379
Size:	100 μL
Class:	Polyclonal
Type:	Antibody
Clone:	
Host / Isotype:	Rabbit / IgG
Immunogen:	Synthetic peptide corresponding to amino acid residues specific to human collagen 1, alpha 1 propeptide conjugated to KLH

Form Information	
Form:	Liquid
Concentration:	Lot-specific
Purification:	Affinity chromatography
Storage Buffer:	10mM HEPES, pH 7.5, with 150mM NaCl, 100µg/ml BSA, 50% glycerol
Preservative:	no preservative
<b>Storage Conditions:</b>	-20°C

## **Product Specific Information**

This antibody is predicted to react with most mammals, birds, amphibians and fish based on 100% sequence homology.

This antibody contains enough material to conduct 10 mini-Western blots. For Research Use Only. Not for use in diagnostic procedures. Not for resale without express authorization.

## **General Information**

Type I collagen is a triple helix comprised of two alpha-1 chains and one alpha-2 chain. Type I collagen is a member of group I collagen (fibril-forming collagen) found in most connective tissues, and is abundant in bone, cornea, dermis and tendon. Mutations in the COL1A1 gene encoding the alpha-1 chain are associated with osteogenesis imperfecta types I-IV, Ehlers-Danlos syndrome type VIIA, Ehlers-Danlos syndrome Classical type, Caffey Disease and idiopathic osteoporosis. Mutations in the COL1A2 gene encoding the alpha-2 chain are associated with osteogenesis imperfecta types I-IV, Ehlers-Danlos syndrome type VIIB, recessive Ehlers-Danlos syndrome Classical type, idiopathic osteoporosis, and atypical Marfan syndrome. Symptoms associated with mutations in this gene, however, tend to be less severe than mutations in the COL1A1 gene, reflecting the different role of alpha-2 chains in matrix integrity.

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