

Collagen IX Polyclonal Antibody
Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP55360**Specification****Collagen IX Polyclonal Antibody - Product Information**

Application	IHC-P, IHC-F, IF, ICC, E
Primary Accession	P20849
Reactivity	Rat
Host	Rabbit
Clonality	Polyclonal
Calculated MW	89 KDa
Physical State	Liquid
Immunogen	KLH conjugated synthetic peptide derived from human Collagen IX
Epitope Specificity	801-921/921
Isotype	IgG
Purity	
affinity purified by Protein A	
Buffer	0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
SUBCELLULAR LOCATION	Secreted, extracellular space, extracellular matrix (By similarity).
SIMILARITY	Belongs to the fibril-associated collagens with interrupted helices (FACIT) family. Contains 10 collagen-like domains. Contains 1 laminin G-like domain. Heterotrimer of an alpha 1(IX), an alpha 2(IX) and an alpha 3(IX) chain.
SUBUNIT	Covalently linked to the telopeptides of type II collagen by lysine-derived cross-links. Prolines at the third position of the tripeptide repeating unit (G-X-Y) are hydroxylated in some or all of the chains.
Post-translational modifications	Multiple epiphyseal dysplasia 6 (EDM6) [MIM:614135]: A generalized skeletal dysplasia associated with significant morbidity. Joint pain, joint deformity, waddling gait, and short stature are the main clinical signs and symptoms. Radiological examination of the skeleton shows delayed, irregular mineralization of the epiphyseal ossification centers and of the centers of the carpal and tarsal bones. Multiple epiphyseal dysplasia is broadly categorized into the more severe Fairbank and the milder Ribbing types. The Fairbank type is characterized by shortness of stature, short and stubby fingers, small epiphyses in several joints, including the
DISEASE	

knee, ankle, hand, and hip. The Ribbing type is confined predominantly to the hip joints and is characterized by hands that are normal and stature that is normal or near-normal. **Note=**The disease is caused by mutations affecting the gene represented in this entry.

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

Important Note

Background Descriptions

Type IX collagen proteoglycan is a major component of hyaline cartilages where it is located on the surface of the collagen fibrils so that a collagenous domain of the molecule (called COL 3) and a non-collagenous domain (called NC4) project at periodic distances away from the surface of the fibrils.

Collagen IX Polyclonal Antibody - Additional Information

Gene ID 1297

Other Names

Collagen alpha-1(IX) chain, COL9A1

Target/Specificity

Cytoplasmic

Dilution

IHC-P ~ ~ N/A
IHC-F ~ ~ N/A
IF ~ ~ 1:50 ~ 200
ICC ~ ~ N/A
E ~ ~ N/A

Format

0.01M TBS(pH7.4), 0.09% (W/V) sodium azide and 50% Glyce

Storage

Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.

Collagen IX Polyclonal Antibody - Protein Information

Name COL9A1

Function

Structural component of hyaline cartilage and vitreous of the eye.

Cellular Location

Secreted, extracellular space, extracellular matrix

Collagen IX Polyclonal Antibody - Protocols

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

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

Collagen IX Polyclonal Antibody - Images