

COL9A1 Antibody (Center) Blocking Peptide
Synthetic peptide
Catalog # BP2762c**Specification**

COL9A1 Antibody (Center) Blocking Peptide - Product Information

Primary Accession [P20849](#)

COL9A1 Antibody (Center) Blocking Peptide - Additional Information

Gene ID 1297

Other Names

Collagen alpha-1(IX) chain, COL9A1

Target/Specificity

The synthetic peptide sequence used to generate the antibody [AP2762c](/products/AP2762c) was selected from the Center region of human COL9A1. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

COL9A1 Antibody (Center) Blocking Peptide - Protein Information

Name COL9A1

Function

Structural component of hyaline cartilage and vitreous of the eye.

Cellular Location

Secreted, extracellular space, extracellular matrix

COL9A1 Antibody (Center) Blocking Peptide - Protocols

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

- [Blocking Peptides](#)

COL9A1 Antibody (Center) Blocking Peptide - Images**COL9A1 Antibody (Center) Blocking Peptide - Background**

COL9A1 is one of the three alpha chains of type IX collagen, which is a minor (5-20%) collagen component of hyaline cartilage. Type IX collagen is usually found in tissues containing type II collagen, a fibrillar collagen. Studies in knockout mice have shown that synthesis of the alpha 1 chain is essential for assembly of type IX collagen molecules, a heterotrimeric molecule, and that lack of type IX collagen is associated with early onset osteoarthritis. Mutations in the COL9A1 gene are associated with osteoarthritis in humans, with multiple epiphyseal dysplasia, 6, a form of chondrodysplasia, and with Stickler syndrome, a disease characterized by ophthalmic, orofacial, articular, and auditory defects.

COL9A1 Antibody (Center) Blocking Peptide - References

Fresquet, M., J. Biol. Chem. 282 (48), 34634-34643 (2007) Liu, L.Y., Yi Chuan 29 (4), 427-432 (2007) Van Camp, G., Am. J. Hum. Genet. 79 (3), 449-457 (2006) Sivakumaran, T.A., J. Assoc. Res. Otolaryngol. 7 (2), 160-172 (2006)