

Product datasheet

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ARG23685 anti-Collagen IX antibody

Package: 50 μg Store at: -20°C

Summary

Product Description Rabbit Polyclonal antibody recognizes Collagen IX

Tested Reactivity Hu, Ms, Rat, Bov

Tested Application ELISA, FACS, ICC/IF, IHC-P, IP, WB

Specificity The antibody reacts with collagen type IX from Human, Mouse, Rat and Bovine. It exhibits no cross

reactivity.

Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name Collagen IX

Species Rat

Immunogen Collagen type IX from Rat chondrosarcoma.

Conjugation Un-conjugated

Alternate Names MED; EDM6; STL4; DJ149L1.1.2; Collagen alpha-1(IX) chain

Application Instructions

Application table	Application	Dilution
	ELISA	Assay-dependent
	FACS	Assay-dependent
	ICC/IF	Assay-dependent
	IHC-P	Assay-dependent
	IP	10 - 30 μg/ml
	WB	300 - 1200 ng/ml
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form	Liquid	
Purification	Purified by affinity chromatography.	
Buffer	PBS	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed before use.	

Bioinformation

Gene Symbol COL9A1

Gene Full Name collagen, type IX, alpha 1

Background This gene encodes 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 this 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. Two transcript variants that encode different isoforms have been identified for this gene. [provided by RefSeq, Jul 2008]

Function Structural component of hyaline cartilage and vitreous of the eye. [UniProt]

Calculated Mw 92 kDa

PTM 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. [UniProt]