

Product datasheet

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ARG62490 anti-Fibrillin 1 antibody [11C1.3]

Package: 100 μl Store at: -20°C

Summary

Product Description Mouse Monoclonal antibody [11C1.3] recognizes Fibrillin 1

Tested Reactivity Hu

Tested Application EM, FACS, ICC/IF, IHC-Fr, IHC-P, WB

Host Mouse

Clonality Monoclonal

11C1.3 Clone

Isotype IgG1, kappa

Target Name Fibrillin 1

Species Bovine

Immunogen Microfibrils from zonular apparatus of bovine eye.

Epitope Amino acids 451 - 909.

Conjugation Un-conjugated

Alternate Names ECTOL1; MFS1; WMS; SGS; SSKS; MASS; GPHYSD2; WMS2; ACMICD; OCTD; Fibrillin-1; FBN

Application Instructions

Application Note WB: 1 - 2 μg/ml

> FACS: 2µg for 106 cells ICC/IF: 5 μg/ml IHC: 1/10 - 1/500

* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations

should be determined by the scientist.

Properties

Form Liquid

Purification Protein G purified

Buffer 1X PBS buffer with < 0.1% sodium azide.

Preservative < 0.1% sodium azide.

Concentration 2 mg/ml

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.

Note For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Database links GeneID: 2200 Human

Swiss-port # P35555 Human

Gene Symbol FBN1

Gene Full Name fibrillin 1

Background This gene encodes a member of the fibrillin family. The encoded protein is a large, extracellular matrix

glycoprotein that serve as a structural component of 10-12 nm calcium-binding microfibrils. These microfibrils provide force bearing structural support in elastic and nonelastic connective tissue throughout the body. Mutations in this gene are associated with Marfan syndrome, isolated ectopia lentis, autosomal dominant Weill-Marchesani syndrome, MASS syndrome, and Shprintzen-Goldberg

craniosynostosis syndrome. [provided by RefSeq, Jul 2008]

Function Fibrillins are structural components of 10-12 nm extracellular calcium-binding microfibrils, which occur

either in association with elastin or in elastin-free bundles. Fibrillin-1-containing microfibrils provide long-term force bearing structural support. Regulates osteoblast maturation by controlling TGF-beta bioavailability and calibrating TGF-beta and BMP levels, respectively (By similarity). [UniProt]

Research Area Signaling Transduction antibody

Calculated Mw 312 kDa

PTM Fibrillin-1: Cleavage of N- and C-terminus by furin is required for incorporation into the extracellular

matrix and assembly into microfibrils (PubMed:27026396). The C-terminus, which corresponds to the Asprosin chain, was initially thought to constitute a propeptide (PubMed:24982166). Fibrillin-1 and Asprosin chains are still linked together during the secretion from cells, but are subsequently separated

by furin, an essential step for incorporation of Fibrillin-1 into the nascent microfibrils

(PubMed:24982166).

Fibrillin-1: Forms intermolecular disulfide bonds either with other fibrillin-1 molecules or with other

components of the microfibrils.

Cellular Localization Extracellular microfibrils