

FBN1 Antibody (Internal region)
Peptide-affinity purified goat antibody
Catalog # AF3039a**Specification**

FBN1 Antibody (Internal region) - Product Information

Application	E
Primary Accession	P35555
Other Accession	NP_000129.2 , 2200 , 14118 (mouse) , 83727 (rat)
Predicted Host	Human, Mouse, Rat, Dog, Cow
Clonality	Goat
Concentration	Polyclonal
Isotype	0.5 mg/ml
Calculated MW	IgG
	312298

FBN1 Antibody (Internal region) - Additional Information**Gene ID** 2200**Other Names**

Fibrillin-1, FBN1, FBN

Format

0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

FBN1 Antibody (Internal region) is for research use only and not for use in diagnostic or therapeutic procedures.

FBN1 Antibody (Internal region) - Protein Information**Name** FBN1 ([HGNC:3603](#))**Synonyms** FBN**Function**

[Fibrillin-1]: Structural component of the 10-12 nm diameter microfibrils of the extracellular matrix, which conveys both structural and regulatory properties to load-bearing connective tissues (PubMed:1860873, PubMed:15062093). Fibrillin-1-containing microfibrils provide long-term force bearing structural support (PubMed:15062093).

[27026396](http://www.uniprot.org/citations/27026396)). In tissues such as the lung, blood vessels and skin, microfibrils form the periphery of the elastic fiber, acting as a scaffold for the deposition of elastin (PubMed:[27026396](http://www.uniprot.org/citations/27026396)). In addition, microfibrils can occur as elastin-independent networks in tissues such as the ciliary zonule, tendon, cornea and glomerulus where they provide tensile strength and have anchoring roles (PubMed:[27026396](http://www.uniprot.org/citations/27026396)). Fibrillin-1 also plays a key role in tissue homeostasis through specific interactions with growth factors, such as the bone morphogenetic proteins (BMPs), growth and differentiation factors (GDFs) and latent transforming growth factor-beta-binding proteins (LTBPs), cell-surface integrins and other extracellular matrix protein and proteoglycan components (PubMed:[27026396](http://www.uniprot.org/citations/27026396)). Regulates osteoblast maturation by controlling TGF- beta bioavailability and calibrating TGF-beta and BMP levels, respectively (By similarity). Negatively regulates osteoclastogenesis by binding and sequestering an osteoclast differentiation and activation factor TNFSF11 (PubMed:[24039232](http://www.uniprot.org/citations/24039232)). This leads to disruption of TNFSF11-induced Ca(2+) signaling and impairment of TNFSF11-mediated nuclear translocation and activation of transcription factor NFATC1 which regulates genes important for osteoclast differentiation and function (PubMed:[24039232](http://www.uniprot.org/citations/24039232)). Mediates cell adhesion via its binding to cell surface receptors integrins ITGAV:ITGB3 and ITGA5:ITGB1 (PubMed:[12807887](http://www.uniprot.org/citations/12807887)), (PubMed:[17158881](http://www.uniprot.org/citations/17158881)). Binds heparin and this interaction has an important role in the assembly of microfibrils (PubMed:[11461921](http://www.uniprot.org/citations/11461921)).

Cellular Location

Secreted. Note=Fibrillin-1 and Asprosin chains are still linked together during the secretion from cells, but are subsequently separated by furin (PubMed:24982166) [Asprosin]: Secreted. Note=Secreted by white adipose tissue and circulates in the plasma.

FBN1 Antibody (Internal region) - 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](#)

FBN1 Antibody (Internal region) - Images

FBN1 Antibody (Internal region) - References

Mutation of ACTA2 gene as an important cause of familial and nonfamilial nonsyndromic thoracic aortic aneurysm and/or dissection (TAAD). Morisaki H, Akutsu K, Ogino H, Kondo N, Yamanaka I, Tsutsumi Y, Yoshimuta T, Okajima T, Matsuda H, Minatoya K, Sasaki H, Tanaka H, Ishibashi-Ueda H, Morisaki T, Human mutation 2009 Jul : . PMID: 19639654