

PDGFC Blocking Peptide (N-term)

Synthetic peptide

Catalog # BP20217a

Specification

PDGFC Blocking Peptide (N-term) - Product Information

Primary Accession

[O9NRA1](#)

Other Accession

[O9EOX6](#), [O8CI19](#), [O9I946](#), [NP_057289.1](#)**PDGFC Blocking Peptide (N-term) - Additional Information****Gene ID** 56034**Other Names**

Platelet-derived growth factor C, PDGF-C, Fallotin, Spinal cord-derived growth factor, SCDGF, VEGF-E, Platelet-derived growth factor C, latent form, PDGFC latent form, Platelet-derived growth factor C, receptor-binding form, PDGFC receptor-binding form, PDGFC, SCDGF

Target/Specificity

The synthetic peptide sequence is selected from aa 88-101 of HUMAN PDGFC

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.

PDGFC Blocking Peptide (N-term) - Protein Information**Name** PDGFC**Synonyms** SCDGF**Function**

Growth factor that plays an essential role in the regulation of embryonic development, cell proliferation, cell migration, survival and chemotaxis. Potent mitogen and chemoattractant for cells of mesenchymal origin. Required for normal skeleton formation during embryonic development, especially for normal development of the craniofacial skeleton and for normal development of the palate. Required for normal skin morphogenesis during embryonic development. Plays an important role in wound healing, where it appears to be involved in three stages: inflammation, proliferation and remodeling. Plays an important role in angiogenesis and blood vessel development. Involved in fibrotic processes, in which transformation of interstitial fibroblasts into myofibroblasts plus collagen deposition occurs. The CUB domain has mitogenic activity in coronary artery smooth muscle cells, suggesting a role beyond the maintenance of the

latency of the PDGF domain. In the nucleus, PDGFC seems to have additional function.

Cellular Location

Cytoplasm, cytosol. Secreted. Nucleus. Cytoplasmic granule. Cell membrane. Note=Sumoylated form is predominant in the nucleus (PubMed:15247255). Stored in alpha granules in platelets (PubMed:15061151).

Tissue Location

Expressed in the fallopian tube, vascular smooth muscle cells in kidney, breast and colon and in visceral smooth muscle of the gastrointestinal tract. Highly expressed in retinal pigment epithelia. Expressed in medulloblastoma. In the kidney, constitutively expressed in parietal epithelial cells of Bowman's capsule, tubular epithelial cells and in arterial endothelial cells (at protein level) Highly expressed in the platelets, prostate, testis and uterus. Higher expression is observed in uterine leiomyomata. Weaker expression in the spleen, thymus, heart, pancreas, liver, ovary cells and small intestine, and negligible expression in the colon and peripheral blood leukocytes.

PDGFC Blocking Peptide (N-term) - Protocols

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

- [Blocking Peptides](#)

PDGFC Blocking Peptide (N-term) - Images

PDGFC Blocking Peptide (N-term) - Background

The protein encoded by this gene is a member of the platelet-derived growth factor family. The four members of this family are mitogenic factors for cells of mesenchymal origin and are characterized by a core motif of eight cysteines. This gene product appears to form only homodimers. It differs from the platelet-derived growth factor alpha and beta polypeptides in having an unusual N-terminal domain, the CUB domain. Alternatively spliced transcript variants have been found for this gene.

PDGFC Blocking Peptide (N-term) - References

Suo, G., et al. Biol. Reprod. 81(4):749-758(2009)
Choi, S.J., et al. Eur. J. Hum. Genet. 17(6):774-784(2009)
Crawford, Y., et al. Cancer Cell 15(1):21-34(2009)
Bruland, O., et al. BMC Cancer 9, 425 (2009) :
Zhao, J., et al. Exp. Cell Res. 314(14):2529-2543(2008)