

Human/Mouse VEGF-B_{167/186} **Antibody**

Monoclonal Mouse IgG₁ Clone # 58013 Catalog Number: MAB751

DESCRIPTION		
Species Reactivity	Human/Mouse	
Specificity	Detects human and mouse VEGF-B in direct ELISAs and Western blots. Recognizes both the VEGF-B ₁₆₇ and VEGF-B ₁₈₆ isoforms. In Western blots, no cross-reactivity with recombinant human (rh) VEGF, rhVEGF-C, rhVEGF-D, rhCTGF, rhPIGF, rhLDGF, or rhPDGF is observed.	
Source	Monoclonal Mouse IgG ₁ Clone # 58013	
Purification	Protein A or G purified from hybridoma culture supernatant	
Immunogen	E. coli-derived recombinant human VEGF-B ₁₆₇ Pro22-Arg188 Accession # AAB06274	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS.	
APPLICATIONS		
Please Note: Optimal dilution	ions should be determined by each laboratory for each application.	General Protocols are available in the Technical Information section on our website.
	Recommended Concentration	Sample
Western Blot	· ·	Recombinant Human VEGF-B ₁₆₇ (Catalog # 751-VE) Recombinant Mouse VEGF-B ₁₆₇ (Catalog # 2595-VE) Recombinant Mouse VEGF-B ₁₈₆ (Catalog # 767-VE)
Immunohistochemis	stry 8-25 μg/mL	Immersion-fixed paraffin-embedded sections of human lung and breast
PREPARATION AND S	STORAGE	
Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C	
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 6 months, -20 to -70 °C under sterile conditions after reconstitution.	

BACKGROUND

Vascular endothelial growth factor B (VEGF-B), also known as vascular endothelial growth factor-related factor (VRF), is a member of the VEGF family of growth factors that share structural and functional similarity (1, 2). Five mammalian members, including VEGF-A, -B, -C, -D and PIGF, have been identified. VEGF family members are disulfide-linked dimeric proteins that are important regulators of physiological and pathological vasculogenesis, angiogenesis and lymphangiogenesis. VEGF-B is expressed in most tissues, especially in heart, skeletal muscle and pancreas. In many tissues, VEGF-B is co-expressed and can heterodimerize with VEGF (3). By alternative splicing, two isoforms of mature VEGF-B containing 167 or 186 amino acid (aa) exist (3, 4). The two VEGF-B isoforms have identical amino-terminal cysteine-knot VEGF homology domains but the carboxyl end of VEGF-B167 differs from that of VEGF-B186 by the presence of a highly basic cysteine-rich heparin binding domain. Whereas VEGF-B186 is a secreted diffusible protein, VEGF-B167 is sequestered into the cell matrix after secretion. Both VEGF-B isoforms bind VEGF receptor 1 (VEGF R1), but not VEGF R2 or VEGF R3 (5). On endothelial cells, ligation of VEGF-B has been shown to regulate the expression and activity of urokinase type plasminogen activator and plasminogen activator inhibitor 1. VEGF-B167 and a proteolytically processed form of VEGF-B186 (VEGF-B127) also bind neuropilin-1 (NP-1), a type I transmembrane receptor for semaphorins/collapsins, ligands involved in neuron guidance (6). Besides VEGF-B, NP-1 has been shown to bind PLGF-2, VEGF165 and VEGF R1 (6, 7). The many interactions of NP-1 with VEGF ligands and receptor suggests that NP-1 may function as a regulator of angiogenesis (7).

References:

- 1. Li, X. and U. Eriksson (2001) Int. J. Biochem Cell Biol. 33:421.
- 2. Olofsson, B. et al. (1999) Curr. Opin. Biotechnol. 10:528.
- 3. Olofsson, B. et al. (1996) Proc. Nat. Acad. Sci. USA 93:2576.
- 4. Grimmond, S. et al. (1996) Benome Res. 6:124.
- Olofsson, B. et al. (1998) Proc. Nat. Acad. Sci. USA 95:11709.
- Makinen, T. et al. (1999) J. Biol. Chem. 274:21217.
- 7. Fuh, G. et al. (2000) J. Biol. Chem. 275:26690

