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## TIE2 Rabbit pAb

Catalog Number: bs-1300R

Target Protein: TIE2
Concentration: 1mg/ml

Form: Liquid Host: Rabbit

Clonality: Polyclonal

Isotype: IgG

Applications: Flow-Cyt (3ug/Test), ICC/IF (1:100)

Reactivity: Human, Rat (predicted:Mouse, Pig, Cow, Horse)

Predicted MW: 124 kDa Entrez Gene: 7010 Swiss Prot: Q02763

Source: KLH conjugated synthetic peptide derived from human Tie2: 401-550/1120.

Purification: affinity purified by Protein A

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

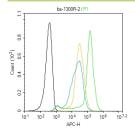
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: The TEK receptor tyrosine kinase is expressed almost exclusively in endothelial cells in mice,

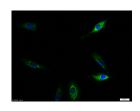
rats, and humans. This receptor possesses a unique extracellular domain containing 2 immunoglobulin-like loops separated by 3 epidermal growth factor-like repeats that are connected to 3 fibronectin type III-like repeats. The ligand for the receptor is angiopoietin-1. Defects in TEK are associated with inherited venous malformations; the TEK signaling pathway appears to be critical for endothelial cell-smooth muscle cell communication in

venous morphogenesis. TEK is closely related to the TIE receptor tyrosine kinase.

## **VALIDATION IMAGES**



Blank control (Black line): HUVEC (Black). Primary Antibody (green line): Rabbit Anti-TIE2 antibody (bs-1300R) Dilution:  $1\mu g/10^6$  cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-AF647 Dilution:  $1\mu g/test$ . Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 0.1% PBST for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.



Tissue/cell:A549 cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Antibody incubation with (TIE2) polyclonal Antibody, Unconjugated (bs-1300R) 1:100, 90 minutes at 37°C; followed by a FITC conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.

## PRODUCT SPECIFIC PUBLICATIONS

[IF=9.77] Ruijun He . et al. HIF1A Alleviates compression-induced apoptosis of nucleus pulposus derived stem cells via upregulating autophagy. Autophagy. 2021 Jan 18;1-23 IF; Rat . 33455530

[IF=6.6] Ling Tang. et al. Construction of ROS-Responsive Hyaluronic Acid Modified Paclitaxel and Diosgenin Liposomes and Study on Synergistic Enhancement of Anti-Ovarian Cancer Efficacy. INT J NANOMED. 2024 六月 04 IHC; Mouse . 38859958

[IF=6.7] Lijuan Shi. et al. Vascularized characteristics and functional regeneration of three-dimensional cell reconstruction of oral mucosa equivalents based on vascular homeostasis phenotypic modification. J TISSUE ENG. ;(): WB; Human . 39301507

[IF=6.208] Haiyan Zhou. et al. Glycation of Tie-2 Inhibits Angiopoietin-1 Signaling Activation and Angiopoietin-1-Induced Angiogenesis.

INT J MOL SCI. 2022 Jan;23(13):7137 IHC; Human . 35806141

[IF=5.962] Akatsu Y et al. Fibroblast growth factor signals regulate transforming growth factor-β-induced endothelial-to-myofibroblast transition of tumor endothelial cells via Elk1. Mol Oncol. 2019 May 15. ICC; Mouse . 31094056