bs-4886R

DATACHEET

[Primary Antibody]

TREM1 Rabbit pAb



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- DATASHEET	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		Flow-Cyt (lug/Test)
GenelD: 58217	SWISS: Q9JKE2	Reactivity: Mouse, Rat
Target: TREM1		
Immunogen: KLH conjugated sy 65-150/230.	nthetic peptide derived from mouse TREM1:	
Purification: affinity purified by Protein A		Predicted MW.: ^{23 kDa}
Concentration: 1mg/ml		Subcellular Location: Secreted ,Cell membrane
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.		Location: Secreted, Cell membrane
is expressed on my monocyte-mediate bacterial and fung inflammatory cher surface expression transcript variants	a receptor belonging to the Ig superfamily t veloid cells. This protein amplifies neutrophi ed inflammatory responses triggered by al infections by stimulating release of pro- nokines and cytokines, as well as increased of cell activation markers. Alternatively spli- encoding different isoforms have been note d by RefSeq, Jun 2011].	land

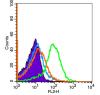
- VALIDATION IMAGES -



Sample: Liver (Rat) Lysate at 40 ug Primary: Anti-TREM1 (bs-4886R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 23 kD Observed band size: 25 kD



Sample: kidney (Mouse) Lysate at 40 ug Primary: Anti-TREM1 (bs-4886R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 23 kD Observed band size: 25 kD



Blank control (black line): Mouse spleen(Black). Primary Antibody (green line): Rabbit Anti-TREM1 antibody (bs-4886R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-PE Dilution: 1µg /test. Protocol The cells were fixed with 4% paraformaldehyde for 10 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 10,000 events was performed.

- SELECTED CITATIONS -

- [IF=38.079] Liu, Xuanyu. et al. Single-cell RNA sequencing of subcutaneous adipose tissues identifies therapeutic targets for cancer-associated lymphedema. CELL DISCOV. 2022 Jun;8(1):1-20 IF ;MOUSe. 35725971
- [IF=10.849] Liu, Xuanyu. et al. Single-cell RNA sequencing identifies an Il1rn+/Trem1+ macrophage subpopulation as a cellular target for mitigating the progression of thoracic aortic aneurysm and dissection. Cell Discov. 2022 Feb;8(1):1-21 IF

;Mouse. 35132073

- [IF=8.484] Amir Boufenzer. et al. Potentiation of NETs release is novel characteristic of TREM-1 activation and the pharmacological inhibition of TREM-1 could prevent from the deleterious consequences of NETs release in sepsis. Cell Mol Immunol. 2021 Jan;18(2):452-460 WB ;Human. 33420354
- [IF=3.754] Zhao, et al. Vitamin D suppresses macrophage infiltration by down-regulation of TREM-1 in diabetic nephropathy rats. (2018) Molecular and Cellular Endocrinology. :. WB, ICC ; Mouse. 29331667
- [IF=3.923] Zhang X et al. Active vitamin D regulates macrophage M1/M2 phenotypes via the STAT-1-TREM-1 pathway in diabetic nephropathy.(2018) J. Cell. Physiol. WB,IHC,IF,ICC ;Human,Rat&Mouse. 30478987