bs-1023R

- DATASHEET -

[Primary Antibody]

G-CSF/CSF3 Rabbit pAb



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Applications: IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500)

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

Predicted MW.: ^{19 kDa}

Subcellular Location: Secreted

Clonality: Polyclonal

GeneID: 1440

Host: Rabbit

SWISS: P09919

Isotype: IgG

Target: G-CSF/CSF3

Immunogen: KLH conjugated synthetic peptide derived from human CSF3: 155-198/207.

Purification: affinity purified by Protein A

Concentration: 1mg/ml

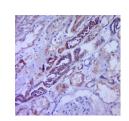
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: Granulocyte-colony stimulating factor (G-CSF) is a growth factor and an essential cytokine belonging to the CSF family of hormonelike glycoproteins that regulate haematopoietic cell proliferation and differentiation. G-CSF was isolated initially as a factor supporting the growth of colonies of granulocytes in soft agar cultures. Cells of the monocyte/macrophage lineage are among the most prominent sources of G-CSF, but this factor can also be produced by normal cells of mesodermal origin, including vascular endothelial cells, fibroblasts, and mesothelial cells. Production of G-CSF can be induced in vitro in these cells by a wide variety of stimulatory agents, including LPS, TNF, IL-1, IL-3, I L-4, and IFN-Gamma. G-CSF is likely to play a role in the basal regulation of neutrophil production, and also functions as a primary regulatory factor controlling the neutrophil response to inflammatory stimuli. Furthermore, G-CSF exhibits other biological activities besides the proliferative effects, since G-CSF appears to modulate the distribution of neutrophils and progenitor cells within the body.

– VALIDATION IMAGES



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (G-CSF) Polyclonal Antibody, Unconjugated (bs-1023R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human kidney tissue); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (CSF3) Polyclonal Antibody, Unconjugated (bs-1023R) at 1:400 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.

- SELECTED CITATIONS -

• [IF=7.3] Kang Zhi-Ying. et al. Heterogeneity of Immune Cells and Their Communications Unveiled by Transcriptome

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- [IF=5.085] Yi Yang. et al. γδ T/Interleukin-17A Contributes to the Effect of Maresin Conjugates in Tissue Regeneration 1 on Lipopolysaccharide-Induced Cardiac Injury. Front Immunol. 2021; 12: 674542 WB ;MOUSE. 33981320
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- [IF=5.177] Wen-Chieh Liao. et al. CHPF promotes malignancy of breast cancer cells by modifying syndecan-4 and the tumor microenvironment. Am J Cancer Res. 2021; 11(3): 812–826 WB,IHC ;Human. 33791155