

**bs-42278R****[ Primary Antibody ]****CXCL2 Rabbit pAb****BioSS**  
**ANTIBODIES**

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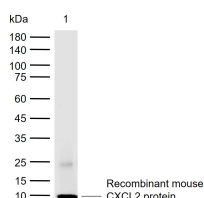
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**— DATASHEET —**

<b>Host:</b> Rabbit	<b>Isotype:</b> IgG	<b>Applications:</b> WB (1:500-2000)
<b>Clonality:</b> Polyclonal		<b>Reactivity:</b> Mouse (predicted: Human)
<b>GeneID:</b> 20310	<b>SWISS:</b> P10889	
<b>Target:</b> CXCL2		
<b>Immunogen:</b> Recombinant mouse CXCL2 protein: 28-100/100.		<b>Predicted MW.:</b> 12 kDa
<b>Purification:</b> affinity purified by Protein A		<b>Subcellular Location:</b> Secreted
<b>Concentration:</b> 1mg/ml		
<b>Storage:</b> 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.		
<b>Background:</b> GRO beta is a member of the CXC, or chemokine class. It contains the ELR domain immediately preceding the first cysteine residue near the amino terminus. Other chemokines in this group include IL8, GRO alpha/beta/gamma, mouse KC, ENA78, GCP2, PBP/CTAPIII/beta TG/NAP2. These chemokines act primarily on neutrophils as chemoattractants and activators, including neutrophil degradation with release of myeloperoxidase and other enzymes. GRO beta was originally identified as a heparin-binding protein secreted from a murine macrophage cell line in response to endotoxin stimulation. GRO beta is an approximately 8 kDa polypeptide of 73 amino acids. The precursor form of GRO beta consists of 100 amino acids. To generate the mature GRO beta, the precursor cleaves its amino terminal 27 amino acids. GRO beta shows 60% amino acid homology to human GRO alpha and GRO gamma.		

**— VALIDATION IMAGES —**

Sample: Lane 1: Recombinant mouse CXCL2 protein, N-His(bs-42278P) Primary: Anti-CXCL2 (bs-42278R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 12 kDa Observed band size: 10 kDa

**— SELECTED CITATIONS —**

- **[IF=7.3]** Wang Xinyi. et al. Identification of potential biomarkers of gout through weighted gene correlation network analysis. FRONT IMMUNOL. 2024 Mar;15: WB ;Human. 10.3389/fimmu.2024.1367019