**Product** Data Sheet

## IL-8/CXCL8 Protein, Cynomolgus (HEK293, C-His)

Cat. No.: HY-P701079

Synonyms: Interleukin-8; IL-8; GCP-1; MDNCF; MONAP; NAP-1; GCP/IL-8 protein IV; IL8/NAP1 form I; CXCL8;

LECT; LUCT; LYNAP; NAF

Species: Cynomolgus Source: **HEK293** 

Accession: A0A2K5TUL7 (A23-P100)

Gene ID: 102127272 Molecular Weight: 12-14 kDa

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TROTERTIES			
Biological Activity	Immobilized Cynomolgus IL-8, His Tag at $0.5\mu g/ml$ ( $100\mu l/well$ ) on the plate. Dose response curve for Anti-IL-8 Antibody, hFc Tag with the EC <sub>50</sub> of 13.0ng/ml determined by ELISA.		
Appearance	Solution.		
Formulation	Supplied as a 0.22μm filtered solution of PBS, pH 7.4.		
Endotoxin Level	<1 EU/μg, determined by LAL method.		
Reconsititution	N/A.		
Storage & Stability	Stored at -80°C for 1 year. It is stable at -20°C for 3 months after opening. It is recommended to freeze aliquots at -80°C for extended storage. Avoid repeated freeze-thaw cycles.		
Shipping	Shipping with dry ice.		

## **DESCRIPTION**

## Background

IL-8/CXCL8 protein serves as a pivotal chemotactic factor, playing a central role in mediating inflammatory responses by attracting neutrophils, basophils, and T-cells to effectively clear pathogens and protect the host from infections. It also contributes significantly to neutrophil activation. Released in response to inflammatory stimuli, IL-8/CXCL8 exerts its effects by binding to G-protein-coupled receptors CXCR1 and CXCR2, primarily found in neutrophils, monocytes, and endothelial cells. The G-protein heterotrimer (alpha, beta, gamma subunits) constitutively binds to CXCR1/CXCR2 receptors, and activation by IL-8 leads to the release of beta and gamma subunits from Galpha (GNAI2 in neutrophils) and subsequent activation of downstream signaling pathways, including PI3K and MAPK pathways. IL-8/CXCL8 forms homodimers, and this dimerization is disrupted by tick evasin-3. Furthermore, IL-8/CXCL8 interacts with TNFAIP6 via its Link domain, and this interaction interferes with chemokine binding to glycosaminoglycans, suggesting a regulatory role in modulating chemokine activity within the inflammatory microenvironment.

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