

Catalog Number: HZ-1084-GMP

Data Sheet





Animal Component-Free

Human cell expressed

Tag-Free

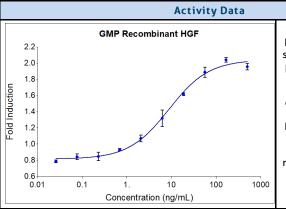
Endotoxin Free

Product Description

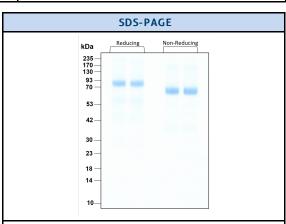
Animal-free Recombinant Human HGF is expressed in human 293 cells and has an apparent molecular mass of 70 kDa. The molecular mass of HGF is lower compared to the reported range for this cytokine expressed in NSO cells (70 to 80 kDa) which emphasizes the difference in post translational modifications. HGF is produced in a serum-free, chemically defined media. HGF is a mesanchymally derived protein mitogen for mature parenchymal hepatocyte cells and acts as a growth factor for a broad spetrum of tissues and cell types.

Alternative Names	F TCF, Hepatocyte growth factor, Hepatopoeitin A, HGF, HGFB, HPTA, Scatter factor, SF	
Accession Number	P14210	
Source	Source Human Embryonic Kidney cells (HEK293). HEK293-derived HGF protein	
Adventitious Virus Master Cell Bank tested Negative for Adventitious Viruses		

Specifications					
Test	Method	Specification			
Activity	Dose-dependent stimulation of the proliferation of the monkey epithelial cell line 4MBr-5	5-25 ng/mL			
Molecular Mass	SDS-PAGE	85 kDa reduced, 70 kDa non-reduced, single chain, glycosylated			
Purity	SDS-PAGE	>95%			
Endotoxin	LAL	<0.1 EU/ μ g			
Mycoplasma	PCR	Not Detected			



GMP-grade recombinant human HGF (HZ-1084-GMP) stimulates dose-dependent proliferation of the 4MBr-5 Monkey epithelial cell line. Viable cell number was quantitiatively assessed by PrestoBlue Cell Viability Reagent. 4MBr-5 cells were treated with increasing concentrations of recombinant human HGF for 120 hours. The EC50 was determined using a 4-parameter non-linear



Purity of GMP-grade recombinant human HGF was determined by SDS- polyacrylamide gel electrophoresis. The protein was resolved in an SDS- polyacrylamide gel in reducing and non-reducing conditions and stained using Coomassie blue.

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	Preparation				
Shipping Temperature ambient temperature					
Formulation 1x PBS, See Certificate of Analysis for details					
Reconstitution Briefly centrifuge the vial before opening. It is recommended to reconstitute the protein to 0.2 mg/mL in sterile 1x PE containing 0.1% endotoxin-free recombinant human serum albumin (HSA). Gently swirl or tap vial to mix.					

	Product Form	Temperature Conditions	Storage Time (From Date of Receipt)
	Lyophilized	-20°C to -80°C	Until Expiry Date
Stability and Storage	Lyophilized	Room Temperature	2 weeks
	Reconstituted as per CofA	-20°C to -80°C	6 months
	Reconstituted as per CofA	4°C	1 week
		Avoid repeated freeze-thaw cycles.	

Proteintech GMP Quality Policy HumanKine® GMP Proteins

Invitro recombinant protein production can be prone to variability due to various reasons ranging from quality of raw materials to inconsistency in the process. Therefore, HumanKine®, a proteintech brand's GMP proteins are produced and tested under an ISO 13485 certified quality management system in a clean room facility. Proteintech manufactures the GMP HumanKine® products according to the applicable sections in the following documents: USP Chapter 1043 (Ancillary Materials for Cell, Gene, and Tissue-Engineered Products, USP Chapter 92 (Growth Factors and Cytokines Used in Cell Therapy Manufacturing), WHO TRS, No. 822, 1992 Annex 1 (Good Manufacturing Practices for Biological Products), Ph. Eur. General Chapter 5.2.12, and EudraLex – Volume 4 – Part IV (Guidelines on GMP specific to ATMPs). Proteintech strives to achieve the utmost quality GMP raw material ensuring all applicable guidelines are taken into consideration.

The QMS is built to provide our customers with consistent and pure product delivered by documented processes, qualified personnel, validated processes, qualified equipment, qualified vendors, and a stringent final product release process. Although the final product release process is important, Proteintech performs in-process QC steps after each major manufacturing stage. Production records and facilities may be available for an inspection by approved personnel.

Our GMP policy covers all the aspects of production; from raw materials, facility, equipment, and Instruments to training and personal hygiene of staff. It also guarantees that the process is explicit, validated and well documented for transparency and traceability.

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