

## Catalog Number: HZ-1296-GMP

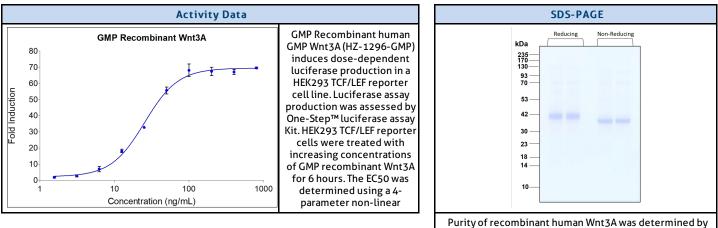
## Data Sheet



Animal Component-Free	Human cell expressed	Tag-Free	Endotoxin Free	

Product Description Animal-free Recombinant Human Wnt3A is expressed in human 293 cells with an apparent molecular mass of 38-42 kDa. This cytokine is produced in a serum-free, chemically defined media.				
Accession Number	Accession Number P56704			
Source	Human Embryonic Kidney cells (HEK293). HEK293-derived Wnt3A protein			
Adventitious Virus	Adventitious Virus Master Cell Bank tested Negative for Adventitious Viruses			

Specifications				
Test	Method	Specification		
Activity	Dose-dependent luciferase production in a Hek293 reporter cell line.	25-500 ng/mL		
Molecular Mass	SDS-PAGE	42 kDa reduced, 40 kDa non-reduced, monomer, glycosylated		
Purity	SDS-PAGE	>90%		
Endotoxin	LAL	<0.1 EU/µg		
Mycoplasma	PCR	Not Detected		



SDS- polyacrylamide gel electrophoresis. The protein was resolved in an SDS- polyacrylamide gel in reducing and non-reducing conditions and stained using Coomassie blue.

SC

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Preparation			
Shipping Temperature	ambient temperature		
Formulation	Formulation 1x PBS pH 7.4 +1% CHAPS, See Certificate of Analysis for details		
Reconstitution Briefly centrifuge the vial before opening. It is recommended to reconstitute the protein to 0.1 mg/mL in sterile 1x PBS pl containing 0.1% endotoxin-free recombinant human serum albumin (HSA). Gently swirl or tap vial to mix.			

Stability and Storage	Product Form	Temperature Conditions	Storage Time (From Date of Receipt)	
	Lyophilized	-20°C to -80°C	Until Expiry Date	
	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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