

CERTIFICATE OF ANALYSIS

Human holo-Transferrin

Powder

(Catalog Number 800-131P)

Lot Number: H425030

Origin: United States Date of Manufacture: May2025

Storage Temperature: 2°C – 8°C Product Expiry: Feb2028

For Cell culture, Further Manufacturing or Research use Only. Not for Direct Therapeutic Use.

Test	Methodology	Specification	Analysis	
Biological Testing			4.48	
Bioburden	BBI SMP 1097	<10 cfu/mL	Pass	
Endotoxin	LAL	≤1 EU/mg	<0.05 EU/mg	
Microbiological Testing				
Mycoplasma		Not Detected	Not Detected	
Viral Testing				
HBsAg		Non-Reactive	Non-Reactive	
Anti-HCV		Non-Reactive	Non-Reactive	
Anti-HIV-1/HIV-2		Non-Reactive	Non-Reactive	
ALT		Negative	Negative	
Syphilis		Negative	Negative	
HBV-NAT		Non-Reactive	Non-Reactive	
HIV-NAT		Non-Reactive	Non-Reactive	
HAV-NAT		Non-Reactive	Non-Reactive	
HCV-NAT		Non-Reactive	Non-Reactive	
Parvo B19-NAT		Non-Reactive	Non-Reactive	
Physical Testing				
Appearance		Report	Salmon Pink to Reddish	
			Powder	
oH @ RT, 3% Solution		6.5 – 8.0	7.49	
Moisture		≤5.0%	1.22%	
Solubility		Readily Soluble at 1% in	Pass	
		De-ionized water		
Optical Density		Report	12.58	
Biochemical Testing		·		
Purity	Cellulose Acetate Electrophoresis	≥98% of total protein	Pass	
Protein Analysis	•	>98%	101.37%	
SDS-Polyacrylamide Gel		Conforms	Conforms	
Iron Estimated by ICP		1200-1700 ppm	1214 ppm	



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Results shown were obtained by carefully performed methods believed to be reliable. However, since some results may vary for specific tests depending on the methodology and other variables, it is suggested that tests for which results are particularly important to be repeated by the user of this product. All human blood products are collected from stringently screened donors at FDA-licensed collection centers in the United States. Single homogenous batch, heat treated at $62^{\circ}\text{C} \pm 2^{\circ}\text{C}$ for 10 hours and lyophilized from approximately 0.02M NH₄HCO₃. May contain traces of buffer salts. Dispensed by dry weight. Store at $2^{\circ}\text{C} - 8^{\circ}\text{C}$. Allow temperature to equilibrate to room temperature prior to use.

The testing that has been performed as part of this lot release has been reviewed by Quality Assurance personnel and it has been confirmed that the testing meets the specifications presented on this Certificate of Analysis.

Kare	en Zuniga		
Name	J		99
QA	Associate	11	
Title			
07m	PM 2025		
Date			