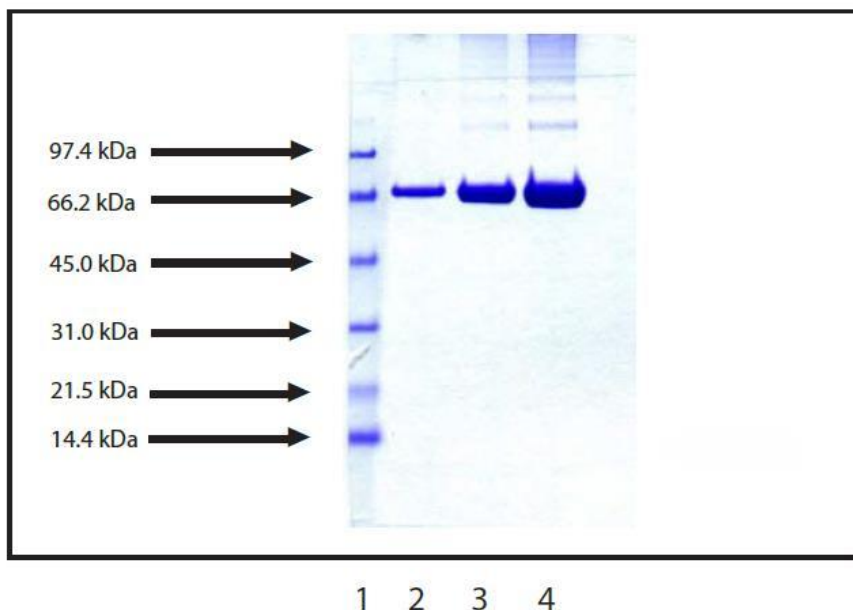


Gel Scan of Representative Lot

Transferrin, Human Plasma (SIDERO), Partially Iron Saturated

No.np001011



SDS-PAGE:

4-12% Bis-Tris NuPAGE gel

1. LMW Standard
2. ransferrin-SIDERO-TC- 5 µg(non-Reduced/no heat)
3. ransferrin-SIDERO-TC- 10 µg(non-Reduced/no heat)
4. ransferrin-SIDERO-TC- 20 µg(non-Reduced/no heat)

Molecular Weight: 80,000

Sterility:

Endotoxin by LAL Assay ≤ 1.0 EU/mg

Sterile filtered using 0.1 micron filter

Considerations:

Storage Conditions: $\leq -20^{\circ}\text{C}$; Stability > 1 year

Buffer: Lyophilized from de-ionized water; pH 3% solution: 6.0-8.0; Heat treated at 60°C for 10 hours.

Iron Content: 300-600 ppm

Protein Determination: Biuret Method

Physical Specifications: Lyophilized; Purity: $\geq 98\%$ by SDS-PAGE

Quality Assurance: Manufactured under ISO9001:2015

Product Datasheet

Product: Transferrin, Human Plasma (SIDERO), Partially Iron Saturated

Number: np001011

Description: Lyophilized

Buffer: from Sterile deionized water, pH 6.0-8.0, heat treated at 60°C for 10 hours

Endotoxin Level: <=1.0 Eu per mg, sterile filtered using 0.1 micron filter

Purity: >=98% by SDS-PAGE

Iron Content: >=300-600 ppm

Molecular Weight: MW 80,000

Protein Determination: Biuret Method

Storage: <= -20°C

Stability: >1 year

Available Packaging: 100mg & per vial; Larger aliquots upon request

Source: Human Plasma

Testing: Shown to be non reactive for HBsAg, anti-HCV, anti-HBc, and negative for anti-HIV 1 & 2 by FDA approved tests.

Certificate Of Origin

Product Name: Transferrin, Human Plasma (SIDERO), Partially Iron Saturated

Catalog Number: np001011

Lot Number: Information applies to all lots produced.

Manufacturing Information: The methods and controls used in this facility for manufacturing, processing, packaging and holding of our products are in accordance with ISO9001:2015 Quality Management System.

Raw Material Information: The product contains no materials directly of animal origin and has not been in contact with products of animal origin in the course of its purification.

Country of Origin (manufacturing): United States

Biological Source Material (COO): United States

Donor Consent: The raw material used in the purification of this product has been obtained from an FDA licensed collection center. Further the donor or the donor's next of kin has given by signature full informed consent for the material to be used in commercial scientific research, including the sharing, transferring, or selling of samples to third parties involved in this research. The raw material was obtained in an ethical manner in full compliance with national legislation, and the donor identity is held as confidential information by the licensed collection center.