



CERTIFICATE OF ANALYSIS

Human Albumin 25% Solution

(Catalog Number 800-120)

Lot Number: H122029

Date of Manufacture: Jun2022

Product Expiry: Nov2023

Origin: United States

Storage Temperature: $\leq 10^{\circ}\text{C}$

| Test | Methodology | Specification | Analysis |
|--------------------------------|---------------|-----------------|--------------|
| Microbiological Testing | | | |
| Sterility | USP<71> | | |
| Bacteria | | Not Detected | Not Detected |
| Fungi | | Not Detected | Not Detected |
| Viral Testing | | | |
| | 21 CFR 610.40 | | |
| HBsAg | | Non-Reactive | Non-Reactive |
| Anti-HCV | | Non-Reactive | Non-Reactive |
| Anti-HIV-1/HIV-2 | | Non-Reactive | Non-Reactive |
| Syphilis | | Negative | Negative |
| HIV-1 NAT | | Non-Reactive | Non-Reactive |
| HBV-NAT | | Non-Reactive | Non-Reactive |
| HCV-NAT | | Non-Reactive | Non-Reactive |
| Physical Testing | | | |
| Purity | | Test and Report | 100% |
| Protein | | Test and Report | 23.0g/dL |
| Endotoxin | | Test and Report | <0.757 EU/mL |
| pH | | Test and Report | 6.96 |

For Cell Culture, further Manufacturing and Research use only. Not for direct Therapeutic use. Results shown were obtained by carefully performed methods believed to be reliable. However, since some results may vary for specific tests depending upon methodology and other variables, it is suggested that tests for which results are particularly important be repeated by the user of this product.

All human plasma is collected from stringently screened donors at FDA-licensed collection centers in the United States.

VANESSA YOUNG
Name

QA ASSOCIATE II
Title

01 JUL 2022
Date