



CERTIFICATE OF ANALYSIS

Human Albumin 25% Solution

(Catalog Number 800-120)

Lot Number: H423040

Date of Manufacture: Nov2023

Product Expiry: Jun2027

Origin: United States

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

| Test | Methodology | Specification | Analysis |
|--------------------------------|---------------|-----------------|--------------|
| Microbiological Testing | | | |
| Sterility | USP<71> | | |
| Bacteria | | No Growth | No Growth |
| Fungi | | No Growth | No Growth |
| Viral Testing | | | |
| HBsAg | 21 CFR 610.40 | 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 | 98% |
| Protein | | Test and Report | 25.0 g/dL |
| Endotoxin | USP<85> | Test and Report | <1.25 EU/mL |
| pH | USP<791> | Test and Report | 7.01 |

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.

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Name

QA ASSOCIATE II
Title

12DEC2023
Date