



GeminiBio
CERTIFICATE OF ANALYSIS

GemCell™ Select Human Serum AB
(Part Number 100-812)

Lot Number:

H121014

Date of Manufacture: Mar 2021

Product Expiry: Mar 2026

Origin: United States

Storage Temperature: ≤ -10°C

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

Product description: *GemCell™ Select Human Serum AB is collected from healthy male donors of the AB serotype at FDA-licensed facilities located in the United States. Donor units are tested for infectious disease markers prior to processing and found to be non-reactive. Gemcell™ Select Human Serum is converted to serum from human plasma using bovine thrombin and sterile-filtered through a 0.1 µm filter prior to freeze.*

| Test | Methodology | Specification | Analysis |
|-----------------------------------|---|---------------|--------------|
| Biological Testing | | | |
| Endotoxin | USP<85>, EP 2.6.14 | < 10.0 EU/ mL | <0.750 EU/mL |
| Hemoglobin | Fleming, AF and Woolf, AJ (1985) | <20.0 mg/ dL | 6.2 mg/dL |
| Microbiological Testing | | | |
| Sterility | USP<71>, EP 2.6.1 | / | |
| Bacteria | | No Growth | No Growth |
| Fungi | | No Growth | No Growth |
| Mycoplasma | Barile, MF and Kern, J (1971), EP 2.6.7 | Not Detected | Not Detected |
| Viral Testing | | | |
| HBsAg | ABBOTT ChLIA | Non-Reactive | Non-Reactive |
| Anti-HCV | ABBOTT ChLIA | Non-Reactive | Non-Reactive |
| Anti-HIV-1/ HIV-2 | ABBOTT ChLIA | Non-Reactive | Non-Reactive |
| Syphilis | ASI RPR | Negative | Negative |
| HBV-NAT | ROCHE NAT/ PCR | Not Detected | Not Detected |
| HIV-NAT | ROCHE NAT/ PCR | Not Detected | Not Detected |
| HCV-NAT | ROCHE NAT/ PCR | Not Detected | Not Detected |
| HAV RNA | | Non-Reactive | Non-Reactive |
| Parvo B19 DNA | | Non-Reactive | Non-Reactive |
| Atypical Antibodies | | Negative | Negative |
| Viral Testing Finished Pool Level | | | |
| HBcAB | | Non-Reactive | Non-Reactive |
| HTLV-I/II | | Non-Reactive | Non-Reactive |
| West Nile Virus | | Not Detected | Not Detected |
| Chagas | | Non-Reactive | Non-Reactive |
| HEV IgG | | Non-Reactive | Non-Reactive |



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| Physical Testing | | | |
| Osmolality | USP<785>, EP 2.2.35 | 260 – 350 mOsm/ kg | 319 mOsm/kg |
| pH @ RT | USP<791> | Test and Report | 8.14 |
| Biochemistry | | | |
| Albumin | | Test and Report | 3.4 g/dL |
| ALT (SGPT) | | Test and Report | 14 U/L |
| AST (SGOT) | | Test and Report | 15 U/L |
| Bilirubin, Total | | Test and Report | 0.2 mg/dL |
| BUN | | Test and Report | 14 mg/dL |
| Calcium | | Test and Report | >15.0 mg/dL |
| Chloride | | Test and Report | 114 mmol/L |
| Cholesterol | | Test and Report | 133 mg/dL |
| Creatinine | | Test and Report | 0.72 mg/dL |
| Glucose | | Test and Report | 87 mg/dL |
| Phosphorus | | Test and Report | 3.0 mg/dL |
| Potassium | | Test and Report | 3.8 mmol/L |
| Protein, Total | Biuret | Test and Report | 5.1 g/dL |
| Sodium | | Test and Report | >165 mmol/L |
| Triglycerides | | Test and Report | 110 mg/dL |
| Uric Acid | | Test and Report | 3.0 mg/dL |

GemCell Select™ Human Serum AB is collected from a maximum of 16 healthy male donors of the AB serotype at FDA-licensed facilities in the United States. This material is defibrinated from source and the viral testing is performed on the individual donor units. All other testing is performed on the final product pool prior to release. The bovine thrombin that is used as part of the conversion to serum is sourced from controlled herds located in the United States and the source cattle are ante and post-mortem inspected by a U.S. Veterinary Service Inspector where they were deemed free of infectious and contagious diseases. All animals used in the production of the thrombin were from a natural beef program in accordance with FDA regulations. Material is derived from human blood and should be considered biohazardous. Universal precautions should be used when handling this material. Precipitate may develop in this product upon freezing and/or thawing; this occurrence does not impact culture performance.

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

Shirna L. Khan
Name

QC Associate
Title

18 May 2021
Date