## **CORNING**

Test

## **Certificate of Analysis**

Product: Fetal Bovine Serum Description: Premium,

United States Origin

**United States** 

Units

Results

**Specification** 

As Reported

As Reported

As Reported

As Reported

As Reported

As Reported

8

4

0.3

74

2.6

56.3

Catalog Number: 35-015-CF, 35-015-CV Lot Number: 11122001

Date of Manufacture: 18 APR 2022 Date of Expiration: APR 2027

Country of Raw Material Lipited States Country of Final

Origin: United States Processing:

Sterility Assurance 10<sup>-3</sup> (Triple 0.1 μm Level: Storage: -40°C to -10°C

Method

**ELISA** 

Endotoxin USP <85> & EP 2.6.14 ≤ 5.0 < 0.1 EU/mL Hemoglobin USP <90> ≤ 25 mg/dL 6.66 Barile and Kern & Mycoplasma Not Detected Not Detected N/A Hoechst Fluorochrome Stain Osmolality USP <785> & EP 2.2.35 260 - 350309 mOsm/kg H<sub>2</sub>O рΗ USP <791> & EP 2.2.3 6.5 - 8.57.16 N/A Sterility (Bacteria & Fungi) USP <71> & EP 2.6.1 No Growth No Growth N/A USP <1057> & EP 2.5.33 **Total Protein** 3.0 - 4.5g/dL 3.5 IU/L Gamma-Glutamyl Transferase (GGT) Chemistry Analyzer ≤ 10 Cell Growth Performance Testing Passed MTT Assay Passed N/A Identification (Species ID) EP 2.7.1 Bovine Bovine N/A 9 CFR Part 113.53c Virus Testing Bluetongue Virus Not Detected Not Detected N/A Bovine Adenovirus 1 & 5 Not Detected Not Detected N/A **Bovine Parvovirus** Not Detected Not Detected N/A Bovine Respiratory Syncytial Virus Not Detected Not Detected N/A Bovine Viral Diarrhea Virus 0.71 TCID<sub>50</sub>/mL As Reported Rabies Virus Not Detected Not Detected N/A Reovirus Not Detected Not Detected N/A Cytopathogenic Agents (IBR) Not Detected N/A Not Detected Hemadsorbing Agents (PI3) Not Detected Not Detected N/A **BVDV Serum Neutralization** Alpha Antibody Test **BVDV Antibody Type 1** As Reported No Neutralization N/A BVDV Antibody Type 2 As Reported N/A No Neutralization Biochemical Testing - Proteins & Organic Compounds Alkaline Phosphatase IU/L As Reported 257 Blood Urea Nitrogen (BUN) As Reported 14 mg/dL Cholesterol As Reported 27 mg/dL Creatinine As Reported mg/dL 2.7 mg/dL Glucose As Reported 100 Glutamic Oxaloacetic Transaminase (AST) As Reported IU/L 45 Glutamic Pyruvic Transaminase (ALT) As Reported IU/L 5

High Density Lipoprotein Cholesterol (HDL)

Low Density Lipoprotein Cholesterol (LDL)

Immunoglobulin G (IgG)

Total Bilirubin

Triglyceride

Uric Acid

mg/dL

µg/mL

mg/dL

mg/dL

mg/dL

mg/dL

Biochemical Testing - Electrophoretic Profile	Cellulose Acetate Electrophoresis	Normal	Normal	N/A
Albumin		As Reported	2.1	g/dL
Alpha		As Reported	1.1	g/dL
Beta		As Reported	0.2	g/dL
Gamma		As Reported	0.1	g/dL
Biochemical Testing - Trace Metals				
Calcium		As Reported	13.3	mg/dL
Chloride		As Reported	97	mmol/L
Iron		As Reported	192	μg/dL
Magnesium		As Reported	3.1	mg/dL
Phosphorus		As Reported	10.2	mg/dL
Potassium		As Reported	>10.0	mmol/L
Sodium		As Reported	136	mmol/L
Biochemical Testing - Hormones				
Cortisol		As Reported	<0.20	μg/dL
Insulin		As Reported	7.78	μIU/mL
Progesterone		As Reported	<0.1	ng/mL
Testosterone		As Reported	<0.01	ng/mL
T3		As Reported	129	ng/dL
T4		As Reported	13.0	μg/dL

## Notes:

For additional information on this Mediatech product, please contact our Technical Services department at (800) 492-1110 or via email at scientificsupport@corning.com.

This product was tested in accordance with Mediatech, Inc. specifications and procedures current as of date of manufacture. Testing procedures are maintained in compliance with the current versions of the USP and/or EP, where applicable.

Cell lines used in MTT Assay: 3T3, L929, MDCK and Vero. Cell lines used in 9CFR Part 113.53c virus testing: MDBK and Vero.

For research or further manufacturing uses only. Not for use as an excipient. Not for therapeutic or diagnostic uses. Not for human or animal consumption. Utilization of this product apart from the labeled intended use may be a violation of local and/or Federal Law.

Product meets European Union requirements for the production of technical blood products.

This product has been granted a Certificate of Suitability, R0-CEP 2018-271, by the European Directorate for the Quality of Medicines (EDQM) certifying that the product meets the criteria described in the current version of the monograph no. 1483 of the European Pharmacopoeia "Product with risk of transmitting agent of animal spongiform encephalopathies"

## Origin and BSE/TSE Statements:

The aforementioned lot was manufactured from fetal bovine blood collected from government approved harvest facilities located in the United States, a country recognized by the United States Department of Agriculture - Animal and Plant Health Inspection Service (USDA-APHIS) and the World Organization for Animal Health (OIE) as having a negligible risk of Bovine Spongiform Encephalopathy (BSE), a bovine-specific Transmissible Spongiform Encephalopathy (TSE). All fetal bovine serum is derived from fetuses collected from healthy bovine cows and heifers that have passed ante and post mortem inspection and were found free of contagious diseases.

Lot Number: 11122001

The following signatures indicate the above material has met all quality specifications and has been reviewed by a Quality representative.

DocuSigned by: DocuSigned by: La Her 03-Jun-2022 Michael Maurer 03-Jun-2022 Written By/Date: Reviewed By/Date: Signer Name: La Her Signer Name: Michael Maurer Signing Reason: I am the author of this document Signing Reason: I have reviewed this document Signing Time: 03-Jun-2022 | 10:09:47 AM EDT Signing Time: 03-Jun-2022 | 11:14:02 AM EDT -B4EFF4CB53B04B75975F5AF91FB7EC70 10857DA1734D435FAB688CBE4BA09317



Lot Number: 11122001