CORNING

Certificate of Analysis

Product: Fetal Bovine Serum Description: Regular,

USDA Approved Origin

Catalog Number: 35-010-CF, 35-010-CV **Lot Number:** 17323001

Date of Manufacture: 09 JUN 2023 Date of Expiration: JUN 2028

Country of Raw Material

Origin:

Mexico

Country of Final Processing:

Storage:

United States

-40°C to -10°C

Sterility Assurance 10⁻³ (Triple 0.1 µm

Level: Filtered)

Test	Method	Specification	Results	Units		
Endotoxin	USP <85> & EP 2.6.14	≤ 20	<0.1	EU/mL		
Hemoglobin	USP <90>	≤ 30	10.22	mg/dL		
Mycoplasma	Barile and Kern & Hoechst Fluorochrome Stain	Not Detected	Not Detected	N/A		
Osmolality	USP <785> & EP 2.2.35	260 – 350	312	mOsm/kg H₂O		
pH	USP <791> & EP 2.2.3	6.5 – 8.5	7.21	N/A		
Sterility (Bacteria & Fungi)	USP <71> & EP 2.6.1	No Growth	No Growth	N/A		
Total Protein	USP <1057> & EP 2.5.33	3.0 – 4.5	3.6	g/dL		
Gamma-Glutamyl Transferase (GGT)	Chemistry Analyzer	≤10	5	IU/L		
Cell Growth Performance Testing	MTT Assay	Passed	Passed	N/A		
Identification (Species ID)	EP 2.7.1	Bovine	Bovine	N/A		
USDA Safety Testing		Pass	Pass	N/A		
Virus Testing	9 CFR Part 113.53c					
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		As Reported	Not Detected	N/A		
Rabies Virus		Not Detected	Not Detected	N/A		
Reovirus		Not Detected	Not Detected	N/A		
Cytopathogenic Agents (IBR)		Not Detected	Not Detected	N/A		
Hemadsorbing Agents (PI3)		Not Detected	Not Detected	N/A		
Biochemical Testing - Proteins & Organic Compounds						
Alkaline Phosphatase		As Reported	261	IU/L		
Blood Urea Nitrogen (BUN)		As Reported	15	mg/dL		
Cholesterol		As Reported	31	mg/dL		
Creatinine		As Reported	3.0	mg/dL		
Glucose		As Reported	85	mg/dL		
Glutamic Oxaloacetic Transaminase (AST)		As Reported	34	IU/L		
Glutamic Pyruvic Transaminase (ALT)		As Reported	5	IU/L		
High Density Lipoprotein Cholesterol (HDL)		As Reported	9	mg/dL		
Immunoglobulin G (IgG)	ELISA	As Reported	159.3	μg/mL		
Low Density Lipoprotein Cholesterol (LDL)		As Reported	8	mg/dL		
Total Bilirubin		As Reported	0.2	mg/dL		
Triglyceride		As Reported	68	mg/dL		
Uric Acid		As Reported	2.4	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	0.3	g/dL
Gamma		As Reported	0.1	g/dL
Biochemical Testing - Trace Metals				
Calcium		As Reported	13.6	mg/dL
Chloride		As Reported	100	mmol/L
Iron		As Reported	166	μg/dL
Magnesium		As Reported	3.2	mg/dL
Phosphorus		As Reported	9.5	mg/dL
Potassium		As Reported	12.2	mmol/L
Sodium		As Reported	138	mmol/L
Biochemical Testing - Hormones				
Cortisol		As Reported	0.274	μg/dL
Insulin		As Reported	8.86	μIU/mL
Progesterone		As Reported	<0.1	ng/mL
Testosterone		As Reported	<0.01	ng/mL
T3		As Reported	121.0	ng/dL
T4		As Reported	19.0	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: BT 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.

Origin and BSE/TSE Statements:

The aforementioned lot was manufactured from fetal bovine blood collected from government approved harvest facilities located in Mexico, 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. This material was imported into the United States under a USDA/APHIS issued import permit.

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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 25-Jul-2023 25-Ju1-2023 Written By/Date: Reviewed By/Date: aniel Au Signer Name: La Her Signer Name: Daniel Hu Signing Reason: I approve this document Signing Reason: I have reviewed this document Signing Time: 25-Jul-2023 | 8:25:39 PM EDT Signing Time: 25-Jul-2023 | 8:28:46 PM EDT -B4EFF4CB53B04B75975F5AF91FB7EC70 BC8EFB6D21DA4812808AFCB60DF446E4



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