## CORNING

## **Certificate of Analysis**

Product: Fetal Bovine		e Serum De		iption:	Regular, USDA Approved Origin		
Catalog Number:	35-010-CF,	35-010-CV	Lot Number:		20722002		
Date of Manufacture:	27 JUL 202	2	Date of	of Expiration:	JUL 2027		
Country of Raw Material Origin:	Mexico		Country of Final Processing:		United States		
Sterility Assurance Level:	10 <sup>-3</sup> (Triple Filtered)	0.1 µm	μm <b>Storage</b> :		-40°C to -10°C		
Test		Method		Specification	Results	Units	
Endotoxin		USP <85> & EP 2.6.	14	≤ 20	<0.1	EU/mL	
Hemoglobin		USP <90>		≤ 30	9.52	mg/dL	
Mycoplasma		Barile and Kern & Hoechst Fluorochror	ne Stain	Not Detected	Not Detected	N/A	
Osmolality		USP <785> & EP 2.2		260 - 350	298	mOsm/kg H₂O	
pH		USP <791> & EP 2.2	2.3	6.5 – 8.5	7.26	N/A	
Sterility (Bacteria & Fungi)		USP <71> & EP 2.6.	1	No Growth	No Growth	N/A	
Total Protein	Total Protein		USP <1057> & EP 2.5.33		3.7	g/dL	
Gamma-Glutamyl Transferase (GGT)		Chemistry Analyzer		≤10	5	IU/L	
Cell Growth Performance Testing		MTT Assay EP 2.7.1		Passed	Passed	N/A	
Identification (Species ID)				Bovine	Bovine	N/A	
Virus Testing		9 CFR Part 113.53c				N1/A	
Bluetongue Virus				Not Detected	Not Detected	N/A	
Bovine Adenovirus 1 & 5				Not Detected	Not Detected	N/A	
Bovine Parvovirus Bovine Respiratory Syncytial Virus				Not Detected	Not Detected	N/A	
Bovine Respiratory Syncytial Virus Bovine Viral Diarrhea Virus				Not Detected	Not Detected Not Detected	N/A N/A	
Rabies Virus				As Reported Not Detected	Not Detected	N/A 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	271	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	87	mg/dL	
Glutamic Oxaloacetic Transaminase (AST)				As Reported	28	IU/L	
Glutamic Pyruvic Transaminase (ALT)				As Reported	6	IU/L	
High Density Lipoprotein Cholesterol (HDL)				As Reported	9	mg/dL	
Immunoglobulin G (IgG)		ELISA		As Reported	197.3	µg/mL	
Low Density Lipoprotein Cholesterol (LDL)				As Reported	9	mg/dL	
Total Bilirubin				As Reported	0.2	mg/dL	
Triglyceride				As Reported	64	mg/dL	
Uric Acid		Cellulose Acetate		As Reported	2.7	mg/dL	
Biochemical Testing - Electrophore	tic Profile	Electrophoresis		Normal	Normal	N/A	
Albumin				As Reported	2.1	g/dL	

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Alpha	As Reported	1.2	g/dL
Beta	As Reported	0.3	g/dL
Gamma	As Reported	0.1	g/dL
Biochemical Testing - Trace Metals			
Calcium	As Reported	13.5	mg/dL
Chloride	As Reported	98	mmol/L
Iron	As Reported	175	µg/dL
Magnesium	As Reported	3.1	mg/dL
Phosphorus	As Reported	9.4	mg/dL
Potassium	As Reported	>10.0	mmol/L
Sodium	As Reported	137	mmol/L
Biochemical Testing - Hormones			
Cortisol	As Reported	0.328	µg/dL
Insulin	As Reported	8.09	µIU/mL
Progesterone	As Reported	<0.1	ng/mL
Testosterone	As Reported	<0.01	ng/mL
Т3	As Reported	119	ng/dL
T4	As Reported	18.0	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.

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



