CORNING

Certificate of Analysis

Product:	oduct: Fetal Bovine S		Im Description:		Regular, USDA Approved Origin	
Catalog Number:	35-010-CF,	35-010-CV	Lot Number:		27021001	
Date of Manufacture:	27 SEP 202	21	Date of Expiration:		SEP 2026	
Country of Raw Material Origin:	Mexico		Country of Final Processing:		United States	
Sterility Assurance Level:	10 ⁻³ (Triple Filtered)	0.1 µm Storaç		ge:	-40°C to -10°C	
Test Endotoxin		Method USP <85> & EP 2.6.	4.4	Specification ≤ 20	Results 8.65	Units EU/mL
Hemoglobin		USP <90> & EF 2.0.	14	≤ 20 ≤ 30	12.88	mg/dL
-		Barile and Kern &				-
Mycoplasma		Hoechst Fluorochror		Not Detected	Not Detected	N/A
Osmolality		USP <785> & EP 2.2		260 – 350	299	mOsm/kg H₂O
pH Storility (Ractoria & Eungi)		USP <791> & EP 2.2		6.5 – 8.5 No Growth	7.24	N/A
Total Protein	Sterility (Bacteria & Fungi)		USP <71> & EP 2.6.1 USP <1057> & EP 2.5.33		No Growth 3.6	N/A g/dL
Gamma-Glutamyl Transferase (GG	T)	Chemistry Analyzer		3.0 – 4.5 ≤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
Virus Testing		9 CFR Part 113.53c		20000		
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 & O	rganic Compour	nds				
Alkaline Phosphatase				As Reported	269	IU/L
Blood Urea Nitrogen (BUN)				As Reported	16	mg/dL
Cholesterol				As Reported	30	mg/dL
Creatinine Glucose				As Reported	2.9	mg/dL
_	minaga (AST)			As Reported As Reported	95 34	mg/dL IU/L
Glutamic Oxaloacetic Transaminase (AST) Glutamic Pyruvic Transaminase (ALT)		ELISA		As Reported	54 6	IU/L
High Density Lipoprotein Cholesterol (HDL)				As Reported	9	mg/dL
Immunoglobulin G (IgG)				As Reported	9 240.3	µg/mL
Low Density Lipoprotein Cholesterol (LDL)				As Reported	8	mg/dL
Total Bilirubin				As Reported	0.2	mg/dL
Triglyceride				As Reported	64	mg/dL
Uric Acid				As Reported	2.8	mg/dL
Biochemical Testing - Electrophoretic Profile		Cellulose Acetate Electrophoresis		Normal	Normal	N/A
Albumin		,		As Reported	2.0	g/dL

Mediatech, Inc. DBA J R Scientific 1242 Commerce Ave. Woodland, CA 95776 1-800-492-1110

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	13.3	mg/dL
Chloride	As Reported	99	mmol/L
Iron	As Reported	171	µg/dL
Magnesium	As Reported	3.1	mg/dL
Phosphorus	As Reported	9.2	mg/dL
Potassium	As Reported	>10.0	mmol/L
Sodium	As Reported	136	mmol/L
Biochemical Testing - Hormones			
Cortisol	As Reported	0.454	µg/dL
Insulin	As Reported	6.65	µIU/mL
Progesterone	As Reported	<0.1	ng/mL
Testosterone	As Reported	<0.01	ng/mL
Т3	As Reported	125	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: 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.

Written By/Date:	Jo fler	08 NOV 2021	Reviewed By/Date: Minhall Muniec	10 NOV 2021
		TRACENSE	TY PROVIN	

THURSDAY.