CORNING

Certificate of Analysis

Product:	Fetal Bovine Serum		Description:		Regular, USDA Approved Origin			
Catalog Number:	35-010-CF, 35-010-CV		Lot Number:		05423001			
Date of Manufacture:	24 MAR 2023		Date of Expiration:		MAR 2028			
Country of Raw Material Origin:			Country of Final Processing:		United States			
Sterility Assurance Level:	10 ⁻³ (Triple Filtered)	0.1 μm Storage:		ge:	-40°C to -10°C			
Test Endotoxin Hemoglobin		Method USP <85> & EP 2.6. USP <90>	14	Specification ≤ 20 ≤ 30	Results <0.1 9.72	Units EU/mL mg/dL		
Mycoplasma		Barile and Kern &	a Ctain	Not Detected	Not Detected	N/A		
Osmolality		Hoechst Fluorochron USP <785> & EP 2.2		260 – 350	307	mOsm/kg H₂O		
рН		USP <791> & EP 2.2		6.5 – 8.5	7.23	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	3.5	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				N1/A		
Bluetongue Virus				Not Detected	Not Detected	N/A		
Bovine Adenovirus 1 & 5 Bovine Parvovirus				Not Detected	Not Detected	N/A		
				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	272	IU/L		
Blood Urea Nitrogen (BUN)				As Reported	16	mg/dL		
Cholesterol				As Reported	31	mg/dL		
Creatinine				As Reported	2.9	mg/dL		
Glucose				As Reported	81	mg/dL		
Glutamic Oxaloacetic Transar				As Reported	25	IU/L		
Glutamic Pyruvic Transamina				As Reported	5	IU/L		
High Density Lipoprotein Chol	lesterol (HDL)			As Reported	10	mg/dL		
Immunoglobulin G (IgG)		ELISA		As Reported	176.3	µg/mL		
Low Density Lipoprotein Chole	esterol (LDL)			As Reported	8	mg/dL		
Total Bilirubin				As Reported	0.2	mg/dL		
Triglyceride				As Reported	64	mg/dL		
Uric Acid				As Reported	2.4	mg/dL		

Biochemical Testing - Electrophoretic Profile	Cellulose Acetate Electrophoresis	Normal	Normal	N/A
Albumin		As Reported	2.1	g/dL
Alpha		As Reported	1.2	g/dL
Beta		As Reported	0.3	g/dL
Gamma		As Reported	0	g/dL
Biochemical Testing - Trace Metals				
Calcium		As Reported	13.1	mg/dL
Chloride		As Reported	98	mmol/L
Iron		As Reported	171	µg/dL
Magnesium		As Reported	3.0	mg/dL
Phosphorus		As Reported	9.0	mg/dL
Potassium		As Reported	11.6	mmol/L
Sodium		As Reported	136	mmol/L
Biochemical Testing - Hormones				
Cortisol		As Reported	0.310	µg/dL
Insulin		As Reported	11.33	µIU/mL
Progesterone		As Reported	<0.1	ng/mL
Testosterone		As Reported	0.02	ng/mL
Т3		As Reported	107.0	ng/dL
T4		As Reported	16.0	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.



