

## **CERTIFICATE OF ANALYSIS**

## Chrysalis<sup>™</sup> Insect Cell Qualified Fetal Bovine Serum

(Catalog Number 100-135)

Lot Number:

A32H00J

Date of Manufacture: Sep 2018

Origin:

**United States** 

**Product Expiry:** Sep 2023

For Cell culture, Further Manufacturing or Research

use Only. Not for Direct Therapeutic Use.

**Storage Temperature**: ≤ -10°C

**Product description:** Chrysalis<sup>™</sup> Insect Cell Qualified Fetal Bovine Serum is sterile-filtered through a 0.1 μm filter

prior to freezing.

Test	Methodology	Specification	Analysis
Biological Testing			
Endotoxin	USP<85>, EP 2.6.14	<10.0 EU/mL	<0.500 EU/mL
Hemoglobin	Fleming, AF and Woolf,	<20.0 mg/ dL	12.1 mg/dL
-	AJ (1965)		
Microbiological Testing			
Sterility	USP<71>, EP 2.6.1		
Bacteria		No Growth	No Growth
Fungi		No Growth	No Growth
Mycoplasma	Barile, MF and Kern, J (1971), EP 2.6.7	Not Detected	Not Detected
Viral Testing	9CFR 113.53c[113.46, 113.47]		
Cytopathic Agents	220.17	Not Detected	Not Detected
Hemadsorbing Agents		Not Detected	Not Detected
Bovine Viral Diarrhea Ab Titer			1100 200000
Type I		Test and Report	1:4
Type II		Test and Report	1:1
Extraneous Viruses (FAb)		7.001.011.07.04	
Bluetongue Virus		Not Detected	Not Detected
Bovine Adenovirus		Not Detected	Not Detected
Bovine Parvovirus		Not Detected	Not Detected
Bovine Viral Diarrhea(cytopathic)		Not Detected	Not Detected
Bovine Viral Diarrhea(non-cytopathic)		Test and Report	Tested
Bovine Syncytial Virus		Not Detected	Not Detected
Bovine Rabie Virus		Not Detected	Not Detected
Reovirus		Not Detected	Not Detected
Physical Testing			
Electrophoretic Pattern	Cellulose Acetate	Normal	Normal
Osmolality	USP<785>, EP 2.2.35	260 - 350 mOsm/ kg	307 m0sm/kg
pH	USP<791>	6.5 – 8.5	7.1



Test	Methodology	Specification	Analysis
Biochemistry			
Albumin		Test and Report	2.2 g/dL
Alkaline Phosphatase		Test and Report	216 U/L
ALT (SGPT)		Test and Report	6 U/L
AST (SGOT)		Test and Report	52 U/L
GGT		<10 U/L	4 U/L
Bilirubin, Total		Test and Report	0.3 mg/dL
BUN		Test and Report	15 mg/dL
Calcium		Test and Report	3.43 mM
Chloride		Test and Report	99 mM
Cholesterol		Test and Report	28 mg/dL
Cortisol		Test and Report	6.48 nM
Creatinine		Test and Report	2.8 mg/dL
Estradiol		Test and Report	180 pM
Glucose		Test and Report	6.83 mM
IgG	ELISA	<0.300 mg/mL*	0.124 mg/mL
Insulin		Test and Report	44.7 pM
Iron, Serum		Test and Report	184 ug/dL
Phosphorus		Test and Report	3.07 mM
Potassium		Test and Report	>10 mM
Progesterone		Test and Report	0.19 nM
Protein, Total	Biuret	2.5 – 6.0 g/dL	3.7 g/dL
Sodium		Test and Report	137 mM
Testosterone		Test and Report	<0.03 nM
T3		Test and Report	2.46 nM
T4		Test and Report	190 nM
Triglycerides		Test and Report	61 mg/dL
Tetracycline		<4 ng/mL	Pass
Uric Acid		Test and Report	2.7 mg/dL

Our fetal bovine serum meets all USDA requirements and has passed ante and post-mortem inspection by a licensed veterinarian. It is collected in countries recognized as being free of foot-and-mouth disease and rinderpest. The origin of all fetal bovine serum is traceable by lot number, date and country of collection. Precipitates may develop in this product upon freezing and/or thawing; this occurrence does not impact culture performance. \*Some geographic regions may test higher than 0.300 mg/mL. If this be the case, refer to GGT results less than 10 U/L to ensure FBS purity.

The testing that has been performed as part of this lot release has been reviewed by Quality Assurance personnel and has confirmed that the testing meets the specifications presented on this Certificate of Analysis.

Megha khalayi	QA Associate III
Name	Title
27 Feb 2020	
Date	×