

CERTIFICATE OF ANALYSIS / ORIGINElectrophoresis

Albumin bovine Fraction V receptor grade, lyophil.		Cat.N Contr	
Parameter	Method	Specification	Result
Molecular weight		ca. 67 000	
Protein (%)	Nitrogen Analyzer	98 - 100	100
(protein factor 6.22)	dry weight basis		
Purity (%)	Cellulose Acetate	98 - 100	100
(albumin, 7 % solution)	Electrophoresis		
рН	7 % solution	6.8 – 7.2	7.1
Sodium (mg/g)	Flame Photometer	0 - 10.0	4.5
Chloride (mg/g)	Coulometric Titrator	0-3.0	0.6
Calcium (mg/g)	Atomic Absorption	0-0.5	0.02
Iron (µg/g)	Atomic Absorption	0 - 5.0	0.9
Heavy metals (µg/g)	Atomic Absorption	0 - 20.0	5.1
L-Lactate (mg/g)	Enzymatic Assay	0 - 0.5	0
Uric Acid (mg/dL)	Colorimetry (7 % solution)	0-4	0
Urea nitrogen (mg/dL)	(7 % solution) Colorimetry (7 % solution)	0-7	0
Total lipids (mg/g)	Supercritical Fluid Extraction	0 – 3.5	0.4
Free fatty acids (mg/g)	Chromatography	0-1.0	0.3
Esterified fatty acids (mg/g)	Chromatography	0-1.5	< 0.1
IgG	Radialimmunodiffusion	none detected	none detected
Moisture (%)	Karl Fischer	0 – 5.0	< 5.0
Minimum shelf life			10/ 2015
Storage (°C)			+2 to +8

The product has been subjected to a heat treatment at a temperature of at least 65°C for a period of at least 3 hours and subjected to pH 5 or lower for a period of at least 3 hours.

It is certified that the product is derived from bovine blood collected from US sourced cattle slaughtered at a USDA licensed establishment located in the USA. The bovine plasma/serum was derived from US sourced animals under 30 months of age that were not stunned using a penetrating device that injects air into the cranial cavity. During the collection process appropriate precautions were taken not to contaminate the serum/plasma with specified risk materials (SRM) as defined by the USDA. The material does not contain, nor is derived from SRM as defined in Commission decision 999/2001. All cattle received ante- and post mortem health inspection under a veterinarian's supervision at the abattoir and were apparently free from infectious and contagious diseases and injurious parasites. Ruminant materials used in the manufacture of this product are not originated from BSE related herds. The record of each raw material collection is incorporated into the manufacturing records and veterinary certificates are maintained on file at the manufacturing site. At the time of manufacture of this lot of material, the US is classified as a GBRIII country and is free from Rinderpest, foot-and-mouth disease, and contagious bovine pleuropneumonia.

We do not guarantee that the product can be used for a special application. This document does not release you from performing the standard control upon receipt of incoming goods.

## SERVA Electrophoresis GmbH Quality Control

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## Dipl.-Ing. (FH) Bernhard Göckel

## Patricia Steinmann

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SERVA Electrophoresis GmbH • D-69115 Heidelberg • Carl-Benz-Str. 7 • Tel.: +49 6221 138 40-0 • Fax +49 6221 138 40-10 Geschäftsführer: Dr. Barbara Müller, Dr. Manfred Kurfürst • HRB 336136 AG Mannheim • E-Mail: info.serva.de • http://www.serva.de COMMERZBANK BLZ: 672 400 39 Kto: 190 38 30 00 • IBAN: DE90 6724 0039 0190 3830 00 • BIC: COBADEFFXXX • Ust.ID-Nr.: DE 812 517 285