

## CERTIFICATE OF ANALYSIS



| Albumin bovine               |        |               | Cat.No. : | 47330 |
|------------------------------|--------|---------------|-----------|-------|
| cell culture grade, lyophil. |        | Contr.No.:    | 150281    |       |
|                              |        |               |           |       |
| Parameter                    | Method | Specification | Res       | sult  |

| Parameter            | Method                       | Specification   | Result      |
|----------------------|------------------------------|---|-------------|
| Appearance           | Visual inspection            | white to yellow with tan<br>to green cast lyophilisate    | corresponds |
| Solubility           | 4 % in H <sub>2</sub> O      | clear to slightly hazy,<br>faint yellow-green<br>solution | corresponds |
| Protein (%)          | Nitrogen analysis            | min. 96   | 99.3        |
| Purity (%)           | Agarose zone electrophoresis | min. 96   | 100         |
| pН                   | 10 % in H <sub>2</sub> O     | 6.5 - 7.5   | 6.9         |
| Endotoxin (EU/mg)    | Limulus Amebocyte Lysate     | max. 3  | < 0.1       |
| Moisture (%)         | Loss on drying               | max. 5.0  | 1.2         |
| APC, average (cfu/g) | Aerobic plate count          | max. 1000   | < 10        |
| Minimum shelf life   |                              |   | 06.10.2021  |
| Storage (°C)         |                              |   | -15 to -25  |

This product was derived from bovine blood collected at a USDA licensed establishment located in the United States. The animals received ante- and post-mortem health inspection at the abattoir by a Veterinary Medical Officer, USDA Food Safety and Inspection Services, and they were apparently free from infectious and contagious diseases and injurious parasites.

During the manufacturing process, the material was subjected to a pH of  $\leq$ 5 and a temperature in excess of 65 °C for at least 3 hours and was not commingled with any other material of animal origin. Blood collection records are incorporated into the manufacturing batch documents.

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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