

CERTIFICATE OF ANALYSIS

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| Hyaluronidase from ovine testes lyophil. EC 3.2.1.35 | Cat.No. : 25118 Contr.No.: 120008 |
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| Parameter | Method | Specification | Result |
|---------------------------|---|---------------------|-------------|
| Molecular weight | | ca. 55 000 | |
| Appearance | | white lyophilisate | corresponds |
| Solubility | 1 mg/ml 0.02 M Phosphate buffer pH 6.9 cont. 0.45 % NaCl and 0.01 % BSA | clear and colorless | corresponds |
| Activity (U/mg) | Hyaluronidase | min. 1 000 | 1 190 |
| Minimum shelf life | | | 03/ 2014 |
| Storage (°C) | | | -15 to -25 |

Unit definition

1 Unit is that amount of enzymatic activity which produces the same reduction in turbidity in a mixture of hyaluronic acid and albumin as one I.U. (International Unit) of a standard hyaluronidase preparation. 1 I-unit = 1 USP-unit = 1 NF-unit.

The product was prepared from ovine testes of New Zealand origin, collected from healthy animals intended for human consumption, and slaughtered in abattoirs under veterinary control.

The raw material was frozen at below minus 15°C immediately after slaughter and kept at this temperature until the time of manufacture.

Furthermore, the preparation was exposed to a pH of less than 4.0 for greater than 24 hours during the purification, and all reasonable precautions were taken to prevent contamination of the product during and after processing.

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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