

Finnish Medicines Agency

CERTIFICATE NUMBER: **FIMEA/2026/000881**

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER^{1, 2}

Part 1

Issued following an inspection in accordance with
Art. 94(1) of Regulation (EU) 2019/6 as amended
Art. 111(5) of Directive 2001/83/EC as amended
Art. 63 of Regulation (EU) 536/2014 as amended

The competent authority of Finland confirms the following:

The manufacturer: ***Eurofins BioPharma Product Testing Finland Oy***

Site address: ***Neulaniementie 2, Kuopio, 70210, Finland***

OMS Organisation Id. / OMS Location Id.: ***ORG-100053886 / LOC-100095610***

Has been inspected under the national inspection programme in connection with manufacturing
authorisation no. ***FIMEA/2024/006407*** in accordance with Art. 88 of Regulation (EU) 2019/6, Art. 61 of
Regulation (EU) No 536/2014 and Art. 40 of Directive 2001/83/EC.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted
on ***2026-03-26***, it is considered that it complies with:

- The principles and guidelines of Good Manufacturing Practice laid down in Directive (EU) 2017/1572 and/or Commission Delegated Regulation (EU) 2017/1569, as reflected by the product categories stated in Part 2.³
- The principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/EEC.³

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. Updates to restrictions or clarifying remarks can be identified through the EudraGMDP website (<http://eudragmdp.ema.europa.eu/>). This certificate is valid only when presented with all pages and both Parts 1 and 2.

The authenticity of this certificate may be verified in EudraGMDP. If it does not appear, please contact the issuing authority.

¹ The certificate referred to in paragraph Art. 80(5) of Directive 2001/82/EC, Art. 111(5) of Directive 2001/83/EC and Art. 15 of Directive 2001/20/EC is also applicable to importers.

² Guidance on the interpretation of this template can be found in the Interpretation of the Union format for GMP certificate.

³ These requirements fulfil the GMP recommendations of WHO.

Part 2

Human Medicinal Products
Human Investigational Medicinal Products
Veterinary Medicinal Products

1 MANUFACTURING OPERATIONS

1.6	Quality control testing
	1.6.2 Microbiological: non-sterility

2 IMPORTATION OF MEDICINAL PRODUCTS

2.1	Quality control testing of imported medicinal products
	2.1.2 Microbiological: non-sterility

2026-06-02

Name and signature of the authorised person of the
Competent Authority of Finland

Confidential
Finnish Medicines Agency
Tel: **Confidential**
Fax: **Confidential**