

Health And Youth Care Inspectorate

CERTIFICATE NUMBER: *NL/H 23/2049089*

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

Part 1

Issued following an inspection in accordance with
Art. 63 of Regulation (EU) 536/2014

The competent authority of Netherlands confirms the following:

The manufacturer: **Intravacc B.V.**

Site address: **Antonie Van Leeuwenhoeklaan 9, Bilthoven, 3721 MA, Netherlands**

OMS Organisation Id. / OMS Location Id.: **ORG-100006508 / LOC-100011722**

Has been inspected under the national inspection programme in connection with manufacturing
authorisation no. **16641 F** in accordance with Art. 61 of Regulation (EU) No 536/2014.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted
on **2023-11-16**, 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

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.

¹The certificate referred to in paragraph Art. 15 of Directive 2001/20/EC is also applicable to importers.

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

³These requirements fulfil the GMP recommendations of WHO.

Part 2

1 MANUFACTURING OPERATIONS	
1.1	Sterile products
	<i>1.1.3 Batch certification</i>
1.2	Non-sterile products
	<i>1.2.1 Non-sterile products (processing operations for the following dosage forms)</i> <i>1.2.1.6 Liquids for internal use</i>
	<i>1.2.2 Batch certification</i>
1.3	Biological medicinal products (list of product types)
	<i>1.3.1 Biological medicinal products (list of product types)</i> <i>1.3.1.2 Immunological products</i> <i>1.3.1.5 Biotechnology products</i>
	<i>1.3.2 Batch Certification (list of product types)</i> <i>1.3.2.2 Immunological products</i> <i>1.3.2.5 Biotechnology products</i>
1.5	Packaging
	<i>1.5.1 Primary Packaging</i> <i>1.5.1.6 Liquids for internal use</i>
	<i>1.5.2 Secondary packaging</i>
1.6	Quality control testing
	<i>1.6.3 Chemical/Physical</i> <i>1.6.4 Biological</i>

Clarifying remarks (for public users)

1.2.1.6 en 1.5.1.6; concerns nasal spray. This document also encompasses CTD IMP's

2024-02-09

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

Confidential
Health And Youth Care Inspectorate
Tel: *Confidential*
Fax: *Confidential*