



*Inspectie voor de Gezondheidszorg - Geneesmiddelen en Medische
Technologie*

CERTIFICATE NUMBER: *NL/H 16/1008994A*

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

Part 1

Issued following an inspection in accordance with :
Art. 111(5) of Directive 2001/83/EC as amended

The competent authority of Netherlands confirms the following:

The manufacturer: *Dr. Fisher Farma B.V.*

Site address: *Schutweg 23, LELYSTAD, 8243PC, Netherlands*

Has been inspected under the national inspection programme in connection with manufacturing
authorisation no. *4384 F* in accordance with Art. 40 of Directive 2001/83/EC transposed in the following
national legislation:

Art. 100 of the Medicines Act

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on
2016-04-07, it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC 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. 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 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, shall also be required for imports
coming from third countries into a Member State.

² Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.

³ These requirements fulfil the GMP recommendations of WHO.



Part 2

Human Medicinal Products	
1 MANUFACTURING OPERATIONS	
1.1	Sterile products
	<i>1.1.3 Batch certification</i>
1.2	Non-sterile products
	<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> 1.3.1.1 Blood products 1.3.1.2 Immunological products 1.3.1.5 Biotechnology products
1.5	Packaging
	<i>1.5.2 Secondary packing</i>

2016-09-29

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

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