



Health And Youth Care Inspectorate

CERTIFICATE NUMBER: *NL/H 23/2048324*

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

Art. 63 of Regulation (EU) 536/2014

The competent authority of Netherlands confirms the following:

The manufacturer: *Synergy Health Utrecht B.V.*

Site address: *De Liesbosch 88, Nieuwegein, 3439 LC, Netherlands*

OMS Organisation Id. / OMS Location Id.: *ORG-100012299 / LOC-100067101*

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. *6785 F* in accordance with Art. 40 of Directive 2001/83/EC and Art. 61 of Regulation (EU) No 536/2014 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 *2023-04-18*, it is considered that it complies with:

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC³

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.

¹The certificate referred to in paragraph Art. 111(5) of Directive 2001/83/EC and Art. 15 of Directive 2001/20/EC, shall also be required for imports coming from third countries into a Member State.

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

³These requirements fulfil the GMP recommendations of WHO.



Part 2

Human Medicinal Products

Human Investigational Medicinal Products

1 MANUFACTURING OPERATIONS

1.6 Quality control testing

1.6.1 *Microbiological: sterility*

1.6.2 *Microbiological: non-sterility*

1.6.3 *Chemical/Physical*

1.6.4 *Biological*

2 IMPORTATION OF MEDICINAL PRODUCTS

2.1 Quality control testing of imported medicinal products

2.1.1 *Microbiological: sterility*

2.1.2 *Microbiological: non-sterility*

2.1.3 *Chemical/Physical*

2.1.4 *Biological*

2023-06-08

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

Jos van der Ven

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