



ISF.405.100.2026.IP.2
WTC/0328_02_01/207

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER¹

Part 1

Issued following an inspection in accordance with Art. 94(1) of Regulation (EU) 2019/6

Chief Pharmaceutical Inspector

/the Competent Authority of Poland/

confirms the following:

the manufacturer

J.S. HAMILTON POLAND Sp. z o. o.

ul. Chwaszczyńska 180, 81-571 Gdynia, POLAND

site address

J.S. HAMILTON POLAND Sp. z o. o.

ul. Chwaszczyńska 180, 81-571 Gdynia, POLAND

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. **097/0328/15** in accordance with Art. 88 of Regulation (EU) 2019/6 Art. 38 and Art. 51a point 2 Pharmaceutical Law of 6th of September 2001 (Journal of Laws from 2026, item 612 as amended).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **30/06/2026 – 02/07/2026**, it is considered that it complies with the Good Manufacturing Practice laid down in Directive 91/412/EEC.

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.

2026 -09- 1 1



Chief Pharmaceutical Inspector

Eukasz Pietrzak

Part 2

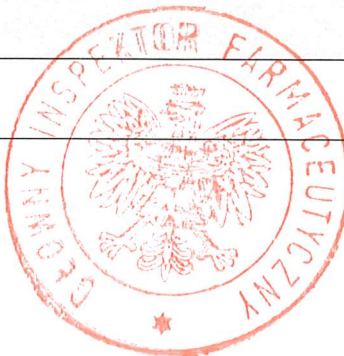
Veterinary Medicinal Products

1 MANUFACTURING OPERATIONS

1.6 Quality control testing

1.6.3 Chemical/Physical

2026 -09- 1 1



Chief Pharmaceutical Inspector

[Signature]
Łukasz Pietrzak

¹ The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 94(1) of Regulation (EU) 2019/6 is also applicable to importers.



ISF.405.100.2026.IP.1
WTC/0328_02_01/206

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER¹

Part 1

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

Chief Pharmaceutical Inspector

/the Competent Authority of Poland/

confirms the following:

the manufacturer

J.S. HAMILTON POLAND Sp. z o. o.
ul. Chwaszczyńska 180, 81-571 Gdynia, POLAND

site address

J.S. HAMILTON POLAND Sp. z o. o.
ul. Chwaszczyńska 180, 81-571 Gdynia, POLAND

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. **097/0328/15** in accordance with Art. 40 of Directive 2001/83/EC transposed in Pharmaceutical Law of 6th of September 2001 (Journal of Laws from 2026, item 612 as amended).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **30/06/2026 – 02/07/2026**, it is considered that it complies with the Good Manufacturing Practice laid down in Directive (EU) 2017/1572.

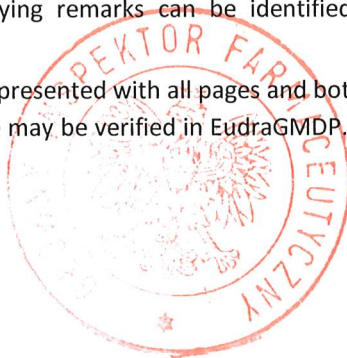
This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended 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.

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2026 -09- 1 1



Chief Pharmaceutical Inspector

Łukasz Pietrzak

Part 2

Human Medicinal Products

1 MANUFACTURING OPERATIONS

1.6 Quality control testing

1.6.3 Chemical/Physical

2 IMPORTATION OF MEDICINAL PRODUCTS

2.1 Quality control testing of imported medicinal products

2.1.3 Chemical/Physical

2026 -09- 1 1



Chief Pharmaceutical Inspector

Łukasz Pietrzak

¹ The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 94(1) of Regulation (EU) 2019/6 is also applicable to importers.