



ISF.405.107.2026.IP.1
WTC/0108_02_01/213

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

AFLOFARM FARMACJA POLSKA Sp. z o.o.

ul. Partyzancka 133/151, 95-200 Pabianice, POLAND

site address

AFLOFARM FARMACJA POLSKA Sp. z o.o.

ul. Szkolna 31, 95-054 Ksawerów, POLAND

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. **051/0108/15** 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 **23/06/2026 - 26/06/2026**, 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.

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

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Chief Pharmaceutical Inspector

Łukasz Pietrzak

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Human Medicinal Products

1 MANUFACTURING OPERATIONS	
1.2	Non-sterile products
	1.2.1 Non-sterile products 1.2.1.1 Capsules, hard shell 1.2.1.8 Other solid dosage forms: powders 1.2.1.12 Suppositories 1.2.1.13 Tablets 1.2.1.17 Other non-sterile medicinal product: raw materials used for preparing medicinal products in the pharmacy according to doctor's prescription or Pharmacopoeia formula 1.2.2 Batch certification
1.3	Biological medicinal products
	1.3.1 Biological medicinal products 1.3.1.8 Other biological medicinal products: product containing microorganisms
1.4	Other products or processing activity
	1.4.1 Manufacture of: 1.4.1.1 Herbal products
1.5	Packaging
	1.5.1 Primary packing 1.5.1.1 Capsules, hard shell 1.5.1.5 Liquids for external use 1.5.1.8 Other solid dosage forms: powders 1.5.1.11 Semi-solids 1.5.1.12 Suppositories 1.5.1.13 Tablets 1.5.1.17 Other non-sterile medicinal products: raw materials used for preparing medicinal products in the pharmacy according to doctor's prescription or Pharmacopoeia formula 1.5.2 Secondary packing
1.6	Quality control testing
	1.6.2 Microbiological: non sterility 1.6.3 Chemical/Physical

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2 IMPORTATION OF MEDICINAL PRODUCTS	
2.1	Quality control testing of imported medicinal products
	2.1.2 Microbiological: non sterility 2.1.3 Chemical/Physical
2.2	Batch certification of imported medicinal products
	2.2.2 Non-sterile products
2.3	Other importation activities
	2.3.1 Site of physical importation

Any restrictions or clarifying remarks related to the scope of this certificate:

Points: 1.2.1.1, 1.2.1.8, 1.2.1.12, 1.2.1.13, 1.5.1.1, 1.5.1.5, 1.5.1.8, 1.5.1.11, 1.5.1.12, 1.5.1.13 include also manufacturing of medicinal products containing highly potent substances .

Points: 1.2.1.1, 1.2.1.13, 1.5.1.1, 1.5.1.13 include also manufacturing of medicinal products containing substances causing allergic or intolerance reactions.

Points: 1.2.1.1, 1.5.1.1 include also manufacturing of medicinal products containing microorganisms.

Point 1.3.1.8 concern medicinal product in the form of capsules, hard shell containing microorganisms.

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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.