



CHIEF PHARMACEUTICAL INSPECTOR

IWSF.405.52.2023.IP.1.1  
WTC/0129\_02\_01/100

**CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER****Part 1**

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

**Chief Pharmaceutical Inspector**

/the Competent Authority of Poland/

confirms the following:

the manufacturer

**Curtis Health Caps S.A.**

Wysogotowo, ul. Batorowska 52, 62-081 Przeźmierowo, POLAND

site address

**Curtis Health Caps S.A.**

Wysogotowo, ul. Batorowska 52, 62-081 Przeźmierowo, POLAND

has been inspected under the national inspection programme in connection with manufacturing authorisation No. **146/0129/15** in accordance with Art. 40 of Directive 2001/83/EC transposed in Pharmaceutical Law of 6<sup>th</sup> of September 2001 (Journal of Laws from 2022, item 2301 as amended).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **21-23/03/2023**, it is considered that it complies with the Good Manufacturing Practice requirements 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.



## Part 2

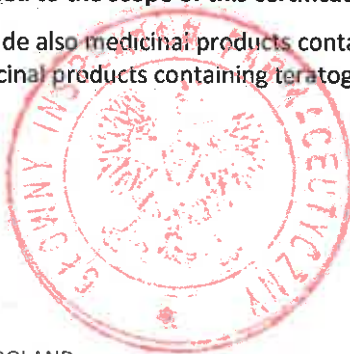
Human Medicinal Products

<b>1 MANUFACTURING OPERATIONS</b>	
<b>1.2</b>	<b>Non-sterile products</b>
	<b>1.2.1 Non-sterile products</b> 1.2.1.2 Capsules, soft shell 1.2.1.6 Liquids for internal use <b>1.2.2 Batch certification</b>
<b>1.4</b>	<b>Other products or processing activity</b>
	<b>1.4.1 Manufacture of:</b> 1.4.1.1 Herbal products
<b>1.5</b>	<b>Packaging</b>
	<b>1.5.1 Primary packing</b> 1.5.1.1 Capsules, hard shell 1.5.1.2 Capsules, soft shell 1.5.1.6 Liquids for internal use 1.5.1.13 Tablets <b>1.5.2 Secondary packing</b>
<b>1.6</b>	<b>Quality control testing</b>
	<b>1.6.2 Microbiological: non sterility</b> <b>1.6.3 Chemical/Physical</b>

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

Points 1.2.1.2, 1.2.1.6, 1.5.1.2, 1.5.1.6 include also medicinal products containing highly active substances.

Points 1.5.1.1, 1.5.1.13 includes also medicinal products containing teratogenic and highly active substances.



On the Chief Pharmaceutical Inspector authority

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 Deputy Chief Pharmaceutical Inspector

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