



IWSF.405.52.2023.IP.2.1
WTC/0129_02_01/101

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with Art. 63(4) of Regulation No 536/2014

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. 61(1) of Regulation (EU) No 536/2014, Art. 38 and Art. 51a point 5 Pharmaceutical Law of 6th 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 Commission Delegated Regulation (EU) 2017/1569.

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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Part 2

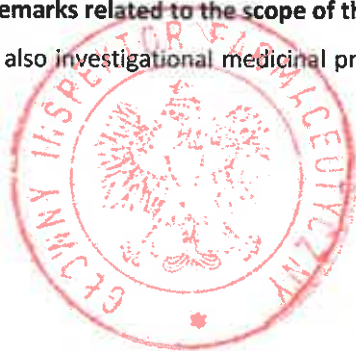
Human Investigational Medicinal Products

1 MANUFACTURING OPERATIONS OF INVESTIGATIONAL MEDICINAL PRODUCTS

1.2	Non-sterile products
	1.2.1 Non-sterile products 1.2.1.6 Liquids for internal use
1.5	Packaging
	1.5.1 Primary packing 1.5.1.6 Liquids for internal use
1.6	Quality control testing
	1.6.2 Microbiological: non sterility 1.6.3 Chemical/Physical

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

Points 1.2.1.6, 1.5.1.6 include also investigational medicinal products containing teratogenic and highly active substances.



On the Chief Pharmaceutical Inspector authority

Paweł Kulka
Deputy Chief Pharmaceutical Inspector