



## CHIEF PHARMACEUTICAL INSPECTOR

ISF.405.117.2025.IP.1  
WTC/0665\_01\_01/246

### CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER<sup>1</sup>

#### 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

**LuCan Laboratories (Pty) Ltd**

**1<sup>st</sup> Floor, Letšeng Building, Kingsway Road Maseru, 100, Lesotho**

site address

**LuCan Laboratories (Pty) Ltd**

**1<sup>st</sup> Floor, Letšeng Building, Kingsway Road Maseru, 100, Lesotho**

Has been inspected in connection with marketing authorisation(s) listing manufacturers located outside of the European Economic Area in accordance with Art. 111(4) of Directive 2001/83/EC transposed in the following national legislation: Pharmaceutical Law of 6th of September 2001 (Journal of Laws from 2025, item 750 as amended).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **17/07/2025 – 18/07/2025**, 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.

2025 -10- 15



Chief Pharmaceutical Inspector  
Łukasz Pietrzak

Human Medicinal Products

**1 MANUFACTURING OPERATIONS**

1.6 Quality control testing

1.6.2 Microbiological: non-sterility

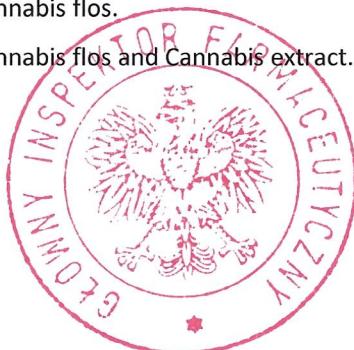
1.6.3 Chemical/Physical

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

Point 1.6.2 concerns dried Cannabis flos.

Point 1.6.3 concerns dried Cannabis flos and Cannabis extract.

2025 -10- 15



Chief Pharmaceutical Inspector

Łukasz Pietrzak

<sup>1</sup>. 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.