ΚΥΠΡΙΑΚΗ ΔΗΜΟΚΡΑΤΙΑ
ΥΠΟΥΡΓΕΙΟ ΥΓΕΙΑΣ
ΦΑΡΜΑΚΕΥΤΙΚΕΣ ΥΠΗΡΕΣΙΕΣ
1475 ΛΕΥΚΩΣΙΑ

REPUBLIC OF CYPRUS
MINISTRY OF HEALTH
PHARMACEUTICAL SERVICES
1475 LEFKOSIA

Our ref.: Ph.S. 5.4.4.51
Tel.: +357 22 608 620
Fax: +357 22 608 649
Email: phscentral@phs.moh.gov.cy

December 4th, 2024

cp FOODLAB LTD

Inspection of the Analytical Laboratory C.P. FOODLAB LIMITED
Inspection no.: 46/2024, date inspection ended: 27th of June 2024
Good Manufacturing Practice (GMP) Certificate no.: FOODLAB/2024/001

Please find, herewith attached, the GMP Certificate for the above-mentioned Laboratory.

The Certificate is issued in accordance to the provisions of article 48(15)(a) of the Medicinal Products for Human Use (Control of Quality, Supply and Prices) Law No. 70(I) of 2001 *[Article 111(5) of the Directive 2001/83/EC]*.

This Certificate is also published in the EudraGMP European Database (http://eudragmp.eudra.org/).

Dr. Helena Panayiotopoulou
Director Pharmaceutical Services
Ministry of Health
Registrar Drugs Council

Pharmaceutical Services Ministry Of Health

CERTIFICATE NUMBER: ***FOODLAB/2024/001***

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER [1,2]

Part 1

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

The competent authority of Cyprus confirms the following:
The manufacturer: ***C.P. Foodlab Limited***
Site address: ***Lakatamia, Viomichanias 19, Nicosia, 2326, Cyprus***
OMS Organisation Id. / OMS Location Id.: ***ORG-100025178 / LOC-100075079***

Has been inspected under the national inspection programme in accordance with Art. 40 of Directive 2001/83/EC, transposed in the following national legislation:PART IV, Chapters A and B of the Medicinal Products for Human Use (Control of Quality, Supply and Prices) Law No 70(I) of 2001, as amended..

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on ***2024-06-27***, it is considered that it complies with::

- The principles and guidelines of Good Manufacturing Practice laid down in Directive (EU) 2017/1572. [3]

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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[1] *The certificate referred to in paragraph Art. 111(5) of Directive 2001/83/ECis also applicable to importers.*

[2] *Guidance on the interpretation of this template can be found in the Interpretation of the Union format for GMP certificate.*

[3] *These requirements fulfil the GMP recommendations of WHO.*

PHARMACEUTICAL SERVICES * REPUBLIC OF CYPRUS * MINISTRY OF HEALTH

Part 2

Human Medicinal Products

1 MANUFACTURING OPERATIONS	
1.6	**Quality control testing**
	1.6.1 Microbiological: sterility
	1.6.2 Microbiological: non-sterility
	1.6.3 Chemical/Physical

2024-12-02

Name and signature of the authorised person of the Competent Authority of Cyprus

Helena Panayiotopoulou
Pharmaceutical Services Ministry Of Health
Tel:*+357 2260 8716*
Fax: