



REPUBLIC OF CYPRUS/
MINISTRY OF HEALTH
PHARMACEUTICAL SERVICES
1475 LEFKOSIA

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July 16th, 2018

cp FOODLAB LTD

Inspection of the Analytical Laboratory cp FOODLAB
Inspection no.: 15/2018, date inspection ended: 15th February 2018
Good Manufacturing Practice (GMP) Certificate no.: FDLB/2018/001

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. Loizos Panayi
Director Pharmaceutical Services
Ministry of Health
Registrar Drugs Council





Pharmaceutical Services Ministry Of Health

CERTIFICATE NUMBER: *FDLB/2018/001*

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: *cp FOODLAB Ltd*

Site address: *25 POLYPHONTI STREET, STROVOLOS, NICOSIA, 2047, Cyprus*

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 *2018-02-15* , it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC³

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. 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.

¹ The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, shall also be required for imports coming from third countries into a Member State.

² Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.

³ These requirements fulfil the GMP recommendations of WHO.



Part 2

Human Medicinal Products

1 MANUFACTURING OPERATIONS

1.6	Quality control testing
	<i>1.6.1 Microbiological: sterility</i>
	<i>1.6.2 Microbiological: non-sterility</i>
	<i>1.6.3 Chemical/Physical</i>

2018-07-10

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



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