



National Organization for Medicines

CERTIFICATE NUMBER: 23374/21-3-13

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER ⁽¹⁾

Part 1

Issued following an inspection in accordance with :
Art. 111(5) of Directive 2001/83/EC as amended
Art. 15 of Directive 2001/20/EC

The competent authority of Greece confirms the following:

The manufacturer: **ΦΑΡΜΑΤΕΝ ΑΒΕΕ / PHARMATHEN SA**

Site address: **Δερβενακίων 6 / Dervenakion 6, Παλλήνη Αττικής / Pallini Attiki, 15351, Greece**

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. **0000006501/13/1** in accordance with Art. 40 of Directive 2001/83/EC and Art. 13 of Directive 2001/20/EC transposed in the following national legislation:

ΔΥΓ 3(α) 83657/24-1-2006, Art. 54

ΔΥΓ 3/89292/03, Art. 12

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **2013-01-30**, it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC ⁽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. 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 EudraGMP. 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.

(3) These requirements fulfil the GMP recommendations of WHO.



Part 2

Human Medicinal Products	
Human Investigational Medicinal Products	
1 MANUFACTURING OPERATIONS	
1.1	Sterile Products
	<i>1.1.3 Batch certification</i>
	<i>1.1.1 Aseptically prepared (processing operations for the following dosage forms)</i> 1.1.1.1 Large Volume Liquids 1.1.1.4 Small volume liquids
	<i>1.1.2 Terminally Sterilised (processing operations for the following dosage forms)</i> 1.1.2.1 Large Volume Liquids 1.1.2.3 Small volume liquids
1.2	Non-sterile products
	<i>1.2.2 Batch certification</i>
	<i>1.2.1 Non-sterile products (processing operations for the following dosage forms)</i> 1.2.1.1 Capsules, hard shell 1.2.1.5 Liquids for external use 1.2.1.13 Tablets
1.5	Packaging
	<i>1.5.1 Primary Packing</i> 1.5.1.1 Capsules, hard shell 1.5.1.2 Capsules, soft shell 1.5.1.5 Liquids for external use 1.5.1.13 Tablets
	<i>1.5.2 Secondary packing</i>
1.6	Quality Control Testing
	1.6.1 Microbiological: sterility 1.6.2 Microbiological: non-sterility 1.6.3 Chemical/Physical



2. IMPORTATION OF MEDICINAL PRODUCTS	
2.1	Quality control testing of imported medicinal products
	2.1.1 Microbiological: sterility 2.1.2 Microbiological: non-sterility 2.1.3 Chemical/Physical
2.2	Batch certification of imported medicinal products
	2.2.1 <i>Sterile Products</i> 2.2.1.1 Aseptically prepared 2.2.1.2 Terminally sterilised
	2.2.2 <i>Non-sterile products</i>
2.3	Other importation activities
	2.3.1 <i>Site of physical importation</i>

Any restrictions related to the scope of this certificate :

Storage of semifinished and finished products, packaging materials, starting materials, cosmetic products and medical devices, in the warehouse in Afoi Ksintara road, Pikermi Attiki, Greece Storage of semifinished and finished products, packaging materials, starting materials, cosmetic products and medical devices, in the warehouse in Marathonos Avenue, Pallini, Attiki, Greece.

Clarifying remarks (for public users)

Storage of semifinished and finished products, packaging materials, starting materials, cosmetic products and medical devices, in the warehouse in Afoi Ksintara road, Pikermi Attiki, Greece Storage of semifinished and finished products, packaging materials, starting materials, cosmetic products and medical devices, in the warehouse in Marathonos Avenue, Pallini, Attiki, Greece.



2013-04-29

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

National Organization for Medicines

Tel:

Fax:

