National Organization for Medicines

CERTIFICATE NUMBER: *37852/13-5-2013*

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. 80(5) of Directive 2001/82/EC as amended

The competent authority of Greece confirms the following:
The manufacturer: ***ΚΟΠΕΡ ΦΑΡΜΑΚΟΒΙΟΜΗΧΑΝΙΑ Α.Ε. / COOPER S.A.***
Site address: ***Αριστοβούλου 64 / 64 Aristovoulou str., Αθήνα / Athens, 11853, Greece***

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. ***0000000740/13/1*** in accordance with Art. 40 of Directive 2001/83/EC and Art. 44 of Directive 2001/82/EC transposed in the following national legislation:
Δ.ΥΓ 3(α)/Γ.Π. 32221/29-4-2013, art. 57
282371/16-6-2006 Art.44

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on ***2013-03-15*** , it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC *(3)*
- The principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/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.

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(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
Veterinary Medicinal Products

1 MANUFACTURING OPERATIONS	
1.1	**Sterile Products**
	1.1.3 Batch certification
	1.1.1 Aseptically prepared (processing operations for the following dosage forms) 1.1.1.4 Small volume liquids 1.1.1.5 Solids and implants Special Requirements 1 B-lactam Antibiotics
	1.1.2 Terminally Sterilised (processing operations for the following dosage forms) 1.1.2.1 Large Volume Liquids 1.1.2.3 Small volume liquids
1.2	**Non-sterile products**
	1.2.2 Batch certification
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.6 Liquids for internal use
1.5	**Packaging**
	1.5.1 Primary Packing 1.5.1.6 Liquids for internal use
	1.5.2 Secondary packing
1.6	**Quality Control Testing**
	1.6.1 Microbiological: sterility 1.6.2 Microbiological: non-sterility 1.6.3 Chemical/Physical

2013-05-14

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

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ΘΕΩΡΗΘΗΚΕ ΓΙΑ ΤΗΝ ΑΚΡΙΒΕΙΑ
Η ΠΡΟΪΣΤΑΜΕΝΗ
ΓΕΝΙΚΗΣ ΓΡΑΜΜΑΤΕΙΑΣ
ΔΕΣΠΟΙΝΑ ΚΟΝΤΟΓΙΑΝΝΗ