French Agency for veterinary medicinal products - French Agency for food, environmental and occupational health safety

CERTIFICATE NUMBER: 18/220389

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with:

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

The competent authority of France confirms the following:

The manufacturer: CEBIPHAR

Site address: 1 RUE DE LA BODINIERE, FONDETTES, 37230, France

Has been inspected under the national inspection programme in connection with manufacturing

authorisation no. V 0533/05 in accordance with Art. 44 of Directive 2001/82/EC.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **2018-08-31**, it is considered that it complies with:

• The principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/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.

Online EudraGMDP, Ref key: 51240 Issuance Date: 2018-11-20 Signatory: Confidential Page 1 of 2

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

Veterinary Medicinal Products

1 MANUFACTURING OPERATIONS							
1.6	Quality control testing						
	1.6.2 Microbiological: non-sterility				7		
	1.6.3 Chemical/Physical					•	

Clarifying remarks (for public users)

This authorisation includes quality control for biological veterinary medicinal products and for sterile and non-sterile chemical veterinary medicinal products for all pharmaceutical forms. This certificate is valid until August 28th 2023.

2018-11-20

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

Confidential

French Agency for veterinary medicinal products -French Agency for food, environmental and occupational health safety

Tel: *Confidential* Fax: *Confidential*