Veterinary Medicines Directorate

Certificate No: VMDGMP/M109/2020

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

UK Manufacturer

Part 1

Issued following an inspection in accordance with Art. 80(5) of Directive 2001/82/EC as amended.

The competent authority of: UNITED KINGDOM (***Veterinary***) confirms the following:

The manufacturer: **Pharmaq Limited**

Site address: Unit 15
Sandleheath Industrial Estate
Fordingbridge
Hampshire
SP6 1PA

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **ManA11003** in accordance with Art 44 of Directive 2001/82/EC transposed in the following national legislation:

The Current Veterinary Medicines Regulations

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **12th to 13th October 2020**, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/EEC.

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.

Signature: Confidential

Date: 23 December 2020 Name: Confidential

Part 2

Veterinary Medicinal Products

1.	**MANUFACTURING OPERATIONS**
1.1	Sterile Products: N/A
1.2	Non-sterile products: 1.2.1.5 Liquids for external use 1.2.2 Batch certification
1.3	Biological medicinal products: 1.3.2.2 Batch certification (immunological products)
1.4	Other products or manufacturing activity: N/A
1.5	Packaging only: 1.5.1 Primary packaging 1.5.1.5 Liquids for external use (ESPA) 1.5.1.8 Other solid dosage forms (non-sterile powders) 1.5.2 Secondary packaging
1.6	Quality Control testing: 1.6.3 Chemical/ Physical

2.	**IMPORTATION OF MEDICINAL PRODUCTS**
2.1	Quality control testing of imported medicinal products N/A
2.2	Batch certification of imported medicinal products N/A

Any restrictions or clarifying remarks related to the scope of this certificate:

Units 6, 7, 8, 15 are involved with GMP operations.

Name and signature of the authorised person of the Competent Authority of the UK:

Signature: Confidential

Date: 23 December 2020

Name: Confidential

Veterinary Medicines Directorate
tel: Confidential
email: Confidential