Veterinary Medicines Directorate

Certificate No: VMDGMP/M184/2023

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

UK Manufacturer

Part 1

Issued following an inspection in accordance with Schedule 2 of the Veterinary Medicines Regulations 2013 as it has effect in Northern Ireland and Great Britain.

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

The manufacturer: **River Manufacturing Limited**

Site address: 61- 63 Windmill Road
Sunbury on Thames
Middlesex
TW16 7DT

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **ManA 19314** in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **9th October 2023**, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice as required by the Veterinary Medicines Regulations 2013.

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 on GOV.UK. If it does not appear, please contact the issuing authority.

Signature: CONFIDENTIAL

Date: 06 December 2023

Name: CONFIDENTIAL

Veterinary Medicines Directorate
Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911
Website: www.vmd.defra.gov.uk

Part 2

Veterinary Medicinal Products

1.	**MANUFACTURING OPERATIONS**
1.1	Sterile Products: N/A
1.2	Non-sterile products: 1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.8 Other solid dosage forms – Ear tags 1.2.2 Batch Certification
1.3	Biological medicinal products: N/A
1.4	Other products or manufacturing activity: N/A
1.5	Packaging: 1.5.1 Primary packing 1.5.1.8 Other solid dosage forms 1.5.2 Secondary packing
1.6	Quality Control testing: N/A

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
2.3	Other importation activities N/A

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

Authorisation of 1.2.1.8 Other solid dosage forms, covers manufacture of moulded permethrin / cypermethrin impregnated ear tags.

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

CONFIDENTIAL

Signature: _____

Date: 06 December 2023

Name: CONFIDENTIAL