Veterinary Medicines Directorate

Certificate No: VMDGMP/M158/2023

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

UK Manufacturer

Part 1

Issued following an inspection in accordance 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: **Waterlife Research Industries Limited**

Site address: 476 Bath Road
Longford
West Drayton
Middlesex
UB7 0ED

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **SAM0003** in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **11 October 2022**, 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: 6th January 2023

Name: Confidential

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) 1.2.1.5 Liquids for external use 1.2.1.13 Tablets 1.2.2 Batch Certification
1.3	Biological medicinal products: N/A
1.4	Other products or manufacturing activity: N/A
1.5	1.5.1 Primary packing: 1.5.1.5 Liquids for external use 1.5.1.13 Tablets 1.5.2 Secondary packing
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
2.3	Other importation activities N/A

Any restrictions or clarifying remarks related to the scope of this certificate:
Schedule 6 (Exemption for Small Pet Animals) Non-sterile only products for use in fish ponds and aquariums where the administration route is via water.

The active substances permitted for use in manufacture are as per the list of approved substances published on the VMD's pages accessed via the GOV.UK website.

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

Signature: Confidential

Date: 6th January 2023

Name: Confidential

Veterinary Medicines Directorate
email: Confidential