Veterinary Medicines Directorate

Certificate No: VMDGMP/M154/2022-v2

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: **Naqua Limited**

Site address: Laboratory E12, Ground Floor,
East Block, Building 500,
Discovery Park,
Ramsgate Road,
Sandwich,
Kent
CT13 9ND

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **ManA1338** in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **14th and 15th September 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 in EudraGMP or on GOV.UK. If it does not appear, please contact the issuing authority.

Signature: CONFIDENTIAL

Date: 8th December 2023 Name: CONFIDENTIAL

Part 2

Veterinary Medicinal Products

1.	**MANUFACTURING OPERATIONS**
1.1	Sterile Products: N/A
1.2	Non-sterile products: 1.2.1.5 Liquids for external use 1.2.1.8 Other solid dosage forms 1.2.2 Batch certification
1.3	Biological medicinal products: N/A
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 1.5.1.8 Other solid dosage forms 1.5.2 Secondary packaging
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:** 2.2.2 Non-sterile products
2.3	**Other importation activities** 2.3.1 Site of physical importation

Any restrictions or clarifying remarks related to the scope of this certificate:
Reissued certificate 08 December 2023 to list specific manufacturing rooms.

The following manufacturing and associated rooms are covered by this certificate:

- Manufacturing room (E12)
- Warehouse / bottle store room
- Pesticide product manufacturing room (S6)
- Finished product storage room (SG010)

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

Signature:

Date: 8th December 2023

Name: CONFIDENTIAL