Veterinary Medicines Directorate

Promoting animal health and welfare by assuring the safety, quality and efficacy of veterinary medicines

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Certificate No: VMDGMP/M232/2025

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 amended) 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 **4-5th September 2025**, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice as required by the Veterinary Medicines Regulations 2013 (as amended).

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: 25 November 2025

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: 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 N/A
2.3	Other importation activities N/A

Any restrictions or clarifying remarks related to the scope of this certificate:
The following manufacturing and associated rooms are covered by this certificate:

- Manufacturing room (E09)
- Warehouse / bottle store room
- Packaging room (E08)
- Finished product storage room (SG010)

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

Signature: CONFIDENTIAL

Date: 25 November 2025

Name: CONFIDENTIAL