



# Veterinary Medicines Directorate

Certificate No: VMDGMP/ VMDGMP/M197/2024

## 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: **NT Laboratories Limited**

Site address: Walton Hall  
Pattenden Lane  
Marden  
Tonbridge  
Kent  
TN12 9QS

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **SAM0001** in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **18<sup>th</sup> January 2024**, 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: 15<sup>th</sup> April 2024

Name: **CONFIDENTIAL**

**Veterinary Medicines Directorate**  
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**Search for VMD on GOV.UK**

The Veterinary Medicines Directorate is an Executive Agency of the Department for Environment, Food and Rural Affairs



## Part 2

Veterinary Medicinal Products
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<b>1. MANUFACTURING OPERATIONS</b>	
1.1	Sterile Products: Not applicable
1.2	Non-sterile products: 1.2.1.5 Liquids for external use 1.2.2 Batch Certification
1.3	Biological medicinal products: Not applicable
1.4	Other products or manufacturing activity: Not applicable
1.5	Packaging: 1.5.1.5 Liquids for external use 1.5.1.8 Other solid dosage forms (powders) 1.5.2 Secondary packing
1.6	Quality Control testing: 1.6.3 Chemical/Physical

<b>2. IMPORTATION OF MEDICINAL PRODUCTS</b>	
2.1	Quality control testing of imported medicinal products Not applicable
2.2	Batch certification of imported medicinal products Not applicable

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

The certificate is restricted to the manufacture of veterinary medicinal products that fall within the scope of Schedule 6 and therefore do not require a Marketing Authorisation.

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

Signature: **CONFIDENTIAL**

Date: 15<sup>th</sup> April 2024

Name: **CONFIDENTIAL**