Veterinary Medicines Directorate

Certificate No: VMDGMP/M145/2022

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Non-UK Manufacturer

Part 1

Issued following an inspection in accordance with Art. 80(5) of Directive 2001/82/EC as amended.

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

The manufacturer: **Zoetis LLC**

Site address: **2000 Rockford Road**
Charles City
Iowa
USA
IA 50616-9101

Has been inspected in connection with marketing authorisation(s) listing manufacturers located outside of the United Kingdom in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **25th to 29th April 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

Name: CONFIDENTIAL

Date: 20th July 2022

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: 1.1.1 Aseptically prepared 1.1.1.1 Large volume liquids 1.1.1.4 Small volume liquids
1.2	Non-sterile products: Not applicable
1.3	Biological medicinal products: 1.3.1 Biological medicinal products 1.3.1.2 Immunological products 1.3.1.8 Other biological medicinal products (Biological active starting materials)
1.4	Other products or manufacturing activity: Not applicable
1.5	Packaging 1.5.1 Primary packaging 1.5.2 Secondary packaging
1.6	Quality Control testing: 1.6.2 Microbiological: non sterility 1.6.3 Chemical/Physical 1.6.4 Biological

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

This certificate does not cover areas exclusively used for the manufacture of non-UK products.

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

CONFIDENTIAL

Signature: _____

Date: 20th July 2022

Name: CONFIDENTIAL