



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: **Kusuri Products Ltd**

Site address: Danefield Business Park
Heathfield Industrial Estate
Wentworth Road
Newton Abbot
Devon
TQ12 6TL
United Kingdom

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. SAM0002 in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **25th July 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 EudraGMP or on GOV.UK. If it does not appear, please contact the issuing authority.

Signature: CONFIDENTIAL

Date: 04 October 2022

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
1.3	Biological medicinal products: N/A
1.4	Other products or manufacturing activity: N/A
1.5	Packaging only: 1.5.1 Primary Packing 1.5.1.5 Liquids for external use 1.5.1.8 Other solid dosage forms 1.5.2 Secondary packing
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

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

1. Restricted to the manufacture of products that fall within the scope of Schedule 6 (Exemptions for Small Pet Animals) of "The Veterinary Medicines Regulations 2013 (2013 No 2033)" and therefore do not require a Marketing Authorisation.

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

CONFIDENTIAL

Signature: _____

CONFIDENTIAL

Date: 04 October 2022

Name:

CONFIDENTIAL