



CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

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: **Cloverleaf Industries Limited**

Site address: Stondon Farm
Ongar Road
Stondon Massey
Essex
CM15 0LD

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **SAM0020** in accordance with Good Manufacturing Practice (GMP)

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **23 June 2021**, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice 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: 26 August 2021

Name: Confidential

Part 2

Veterinary Medicinal Products

1. MANUFACTURING OPERATIONS	
1.1	Sterile Products: N/A
1.2	Non-sterile products: 1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.8 Other solid dosage forms (powders) 1.2.2 Batch Certification
1.3	Biological medicinal products: N/A
1.4	Other products or manufacturing activity: N/A
1.5	1.5.1 Primary packing: 1.5.1.8 Other solid dosage forms (powders) 1.5.2 Secondary packing
1.6	Quality Control testing: 1.6.3 Chemical/Physical

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:

Schedule 6 (Exemption for Small Pet Animals) Non-sterile only products for use in fish ponds and aquariums where the administration route is via water.

The active substances permitted for use in manufacture are as per the list of approved substances published on the VMD's pages accessed via the GOV.UK website.

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

Signature: Confidential

Date: 26 August 2021

Name: Confidential

Veterinary Medicines Directorate
tel: Confidential
email: Confidential