



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: **Custom Powders Limited**

Site address: Gateway
Crewe
Cheshire
CW1 6YT

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **ManA12014** in accordance with Art 44 of Directive 2001/82/EC transposed in the following national legislation:

The Current Veterinary Medicines Regulations

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **2 March 2018**, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/EEC.

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. If it does not appear, please contact the issuing authority.

Signature: _____

Date: 1 May 2018

Name: John O'Neill

Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS

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Part 2

Veterinary Medicinal Products

1. MANUFACTURING OPERATIONS	
1.1	Sterile Products: N/A
1.2	Non-sterile products: N/A
1.3	Biological medicinal products: N/A
1.4	Other products or manufacturing activity: 1.4.1.3 Bulk granules for veterinary premixes or addition to drinking water
1.5	Packaging only: N/A
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

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

None

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



Signature: _____

Date: 1 May 2018

Name: John O'Neill

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