



## 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: **Cod Beck Blenders Limited**

Site address: Cod Beck Estate  
Dalton  
Thirsk  
North Yorkshire  
YO7 3HR

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **ManA 10261** 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 **16 September 2020**, 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: Confidential

Date: 13 October 2020

Name: Confidential

## Part 2

|                               |
|-------------------------------|
| Veterinary Medicinal Products |
|-------------------------------|

|                                    |   |
|------------------------------------|---|
| <b>1. MANUFACTURING OPERATIONS</b> |   |
| 1.1                                | Sterile Products:<br>N/A  |
| 1.2                                | Non-sterile products:<br>1.2.1 Non-sterile products (processing operations for the following dosage forms)<br>1.2.1.8 Other solid dosage forms (Powders for administration Under Schedule 6)<br>1.2.1.16 Veterinary premixes<br>1.2.2 Batch certification |
| 1.3                                | Biological medicinal products:<br>N/A   |
| 1.4                                | Other products or manufacturing activity:<br>N/A  |
| 1.5                                | Packaging:<br>1.5.1 Primary packing<br>1.5.1.8 Other solid dosage forms (Powders for administration under Schedule 6)<br>1.5.1.16 Veterinary premixes<br>1.5.2 Secondary packing  |
| 1.6                                | Quality Control testing:<br>1.6.2 Microbiological: non-sterility<br>1.6.3 Chemical/Physical   |

|   |   |
|---|---|
| <b>2. IMPORTATION OF MEDICINAL PRODUCTS</b> |   |
| 2.1   | Quality control testing of imported medicinal products<br>N/A |
| 2.2   | Batch certification of imported medicinal products<br>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: Confidential

Date: 13 October 2020

Name: Confidential

Veterinary Medicines Directorate  
tel: Confidential  
email: Confidential