Veterinary Medicines Directorate

Certificate No: VMDGMP/M190/2023

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

UK Manufacturer

Part 1

Issued following an inspection in accordance with 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: **Hyperdrug Pharmaceuticals Limited**

Site address:
Station Industrial Estate
Middleton in Teesdale
Barnard Castle
Co. Durham
DL12 0NG
United Kingdom

Has been inspected under the national inspection programme in connection with Manufacturing Authorisation no. **SAM29895** in accordance with Good Manufacturing Practice (GMP).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **03rd October 2023**, 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 GOV.UK. If it does not appear, please contact the issuing authority.

Signature: CONFIDENTIAL

Date: 13 December 2023

Name: CONFIDENTIAL

Part 2

Veterinary Medicinal Products

1.	MANUFACTURING OPERATIONS
1.1	Sterile Products: N/A
1.2	Non-sterile products: 1.2.1.1 Capsules, hard shell 1.2.1.5 Liquids for external use 1.2.1.6 Liquids for internal use 1.2.1.8 Other solid dosage forms 1.2.2 Batch certification
1.3	Biological medicinal products: N/A
1.4	Other products or manufacturing activity: N/A
1.5	Packaging: 1.5.1 Primary packaging 1.5.1.1 Capsules, hard shell 1.5.1.5 Liquids for external use 1.5.1.6 Liquids for internal use 1.5.1.8 Other solid dosage forms 1.5.2 Secondary packaging
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:

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

Signature: CONFIDENTIAL

Date: 13 December 2023

Name: CONFIDENTIAL