

***National Food Chain Safety Office, Directorate of Veterinary Medicinal
Products***

CERTIFICATE NUMBER : **CG-HU/02V/2022.**

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER^{1,2}

Part 1

Issued following an inspection in accordance with :
Art. 94(1) of Regulation (EU) 2019/6 as amended

The competent authority of Hungary confirms the following:

The manufacturer : ***Lavet Kft.***

Site address : ***Batthyany Utca 6, Kistarcsa, 2143, Hungary***

OMS Location : ***LOC-100039871***

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. ***MIA-HU-V-LAVET*** in accordance with Art. 88 of Regulation (EU) 2019/6 .

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **2021-09-24** , it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/EC³

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

¹ The certificate referred to in paragraph Art. 80(5) of Directive 2001/82/EC, shall also be required for imports coming from third countries into a Member State.

² Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.

³ These requirements fulfil the GMP recommendations of WHO.

Part 2

1 MANUFACTURING OPERATIONS	
1.2	Non-sterile products
	<i>1.2.1 Non-sterile products (processing operations for the following dosage forms)</i> 1.2.1.6 Liquids for internal use 1.2.1.8 Other solid dosage forms Special Requirements 1 B-lactam Antibiotics 1.2.1.13 Tablets 1.2.1.16 Veterinary premixes Special Requirements 1 B-lactam Antibiotics
	<i>1.2.2 Batch certification</i>
1.5	Packaging
	<i>1.5.1 Primary Packaging</i> 1.5.1.6 Liquids for internal use 1.5.1.8 Other solid dosage forms Special Requirements 1 B-lactam Antibiotics 1.5.1.13 Tablets 1.5.1.16 Veterinary premixes Special Requirements 1 B-lactam Antibiotics
	<i>1.5.2 Secondary packaging</i>
1.6	Quality control testing
	<i>1.6.3 Chemical/Physical</i>

Any restrictions related to the scope of this certificate :

Building	Room	Line/equipment	QC testing	Products
<i>Building 'A' / 1. floor</i>				<i>liquids for internal use</i>
<i>Building 'B'</i>				<i>other solid dosage forms and vet premixes</i>
<i>Building 'B' / 2. and 3. floor</i>			<i>Chemical/Physical testing</i>	
<i>Building 'C'</i>				<i>Tablet manufacturing</i>
<i>Building 'D'</i>				<i>Storages</i>

2022-03-17

Name and signature of the authorised person of the
Competent Authority of Hungary



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