Medicines and Healthcare products Regulatory Agency CERTIFICATE NUMBER : UK API 14420 Insp GMP 14420/5065-0009 [V] CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER(1),(2)

Part 1

Issued following an inspection in accordance with : Regulation 5 of the current Veterinary Medicines Regulations

The competent authority of United Kingdom confirms the following :

The Manufacturer : LACSA (PTY) LIMITED

Site address : LACSA (PTY) LIMITED, 72 BALLANTRAE ROAD, MEREBANK, DURBAN, 4052, SOUTH AFRICA

Has been inspected in connection with marketing authorisation(s) listing manufacturers located outside of the European Economic Area in accordance with API-A80(1)-D2001/82/EC Regulation 5 of The current Veterinary Medicines Regulations

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 20/11/2024, it is considered that it complies with

• The principles and guidelines of Good Manufacturing Practice laid down in Regulation 5 of the current Veterinary Medicines Regulations

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

- (1) Guidance on the interpretation of this template can be found in the Help menu of MHRA-GMDP database.
- (2) These requirements fulfil the GMP recommendations of WHO.

Part 2

Veterinary Medicinal Products

Manufacture of active substance. Names of substances subject to inspection :

- [2000008781] LACTULOSE CONCENTRATE
- 3. MANUFACTURING OPERATIONS ACTIVE SUBSTANCES LACTULOSE CONCENTRATE

3.1

Manufacture of Active Substance by Chemical Synthesis 3.1.4 Other isomerization reaction **General Finishing Steps**

3.5.2 Primary Packaging

3.5.3 Secondary Packaging

Quality Control Testing

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3.6.1 Physical / Chemical testing

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3.6.2 Microbiological testing (excluding sterility testing)

Any restrictions related to the scope of this certificate:

3.5

3.6

| Building | Room | Line/equipment | QC Testing | Products | |
|------------|--------------|--------------------------------|--------------------------|---------------------------|--|
| | | | | Bulk Lactulose solution | |
| | | | | | |
| 07/03/2025 | Name and s | signature of the authorised pe | rson of the Competent Au | thority of United Kingdom | |
| | Confidential | | | | |
| | Medicines a | ind Healthcare products Regi | ulatory Agency | | |
| | Tel : Confid | ential | | | |