Summary of Product Characteristics

1 NAME OF THE VETERINARY MEDICINAL PRODUCT

Linco-Spectin 100, 222/444.7 mg/g Powder for use in drinking water for pigs and chickens

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each gram contains:

Active substances:

Lincomycin (as lincomycin hydrochloride)	222 mg
Spectinomycin (as spectinomycin sulphate)	444.7 mg

Excipients:
For the full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Product imported from Spain:
Powder for use in drinking water.
White pale powder

4 CLINICAL PARTICULARS

As per VPA 10387/040/001

5 PHARMACOLOGICAL or IMMUNOLOGICAL PROPERTIES

As per VPA 10387/040/001

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Sodium benzoate
Lactose monohydrate

6.2 Major incompatibilities

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

6.3 Shelf-life

Shelf life of the veterinary medicinal product as packaged for sale: 5 years
Shelf life after first opening the immediate packaging: 6 months
Shelf life after dilution according to directions: 24 hours

6.4 Special precautions for storage

This veterinary medicinal product does not require any special storage conditions.

6.5 Nature and composition of immediate packaging

White high density polyethylene (HDPE) bottle containing 1.5 kg powder for use in drinking water with a white tamper evident low density polyethylene (LDPE) lid.

White high density polyethylene (HDPE) bottle containing 150 g powder for use in drinking water with a white tamper evident low density polyethylene (LDPE) lid with an aluminium cap.
Not all pack sizes may be marketed.

6.6 Special precautions for the disposal of unused veterinary medicinal products or waste materials derived from the use of such products

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

Kernfarm BV
De Corridor 14D
Breukelen
36212 ZB
Netherlands

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PVPA22031/001/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 10 August 2018

10 DATE OF REVISION OF THE TEXT

April 2021