Linco-Spectin 100 222 mg/g - 444.7 mg/g Powder for use in drinking water

Authorised

- Spectinomycin sulfate
- Lincomycin hydrochloride

Product identification

Medicine name:
Linco-Spectin 100 222 mg/g - 444.7 mg/g Powder for use in drinking water
LINCO-SPECTIN 100, 222/444,7 MG/G POUDRE POUR ADMINISTRATION DANS L'EAU DE BOISSON POUR PORCINS ET POULETS

Active substance:
Spectinomycin sulfate
Lincomycin hydrochloride

Target species:
Pig
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:

Spectinomycin sulfate

575.90 milligram(s) / 1.00 gram(s)

Lincomycin hydrochloride

241.90 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

Oral use:

- **Pig**
  - Meat and offal. no withdrawal period

0 days; Animals must not be slaughtered for human consumption during treatment.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FF52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

France

Package description:

Linco-Spectin 100 222 mg/g - 444.7 mg/g pwdr. for drinking water 1.5 kg

Linco-Spectin 100 222 mg/g - 444.7 mg/g pwdr. for drinking water 150 g

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Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:
21/07/1992

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8295078 3/1992

Date of authorisation status change:
21/07/2012

Reference member state:
Belgium

Procedure number:
BE/V/0029/001

Concerned member states:
Austria Cyprus Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Poland Portugal Slovakia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.