Lincocin Soluble Powder, 400 mg/g powder for use in drinking water

Authorised

- Lincomycin hydrochloride

Product identification

Medicine name:
Lincocin Soluble Powder, 400 mg/g powder for use in drinking water
Lincocin 400 mg/g proszek do podania w wodzie do picia dla świń i kur

Active substance:
Lincomycin hydrochloride

Target species:
Chicken
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
Lincomycin hydrochloride
453.63 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:

Oral use:

- **Chicken**
  - Meat and offal. 5 day

- **Pig**
  - Meat and offal. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
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.Pack sizes:Bottle of 150 g
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.Pack sizes:Bottle of 1.5 kg

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Polska Sp. z o.o.

Marketing authorisation date:
12/03/2001

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1154

Date of authorisation status change:
12/03/2001

Reference member state:
Ireland

Procedure number:
IE/V/0410/001

Concerned member states:
Belgium France Germany Luxembourg Poland

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
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Package Leaflet

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Labelling

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