Source URL: https://medicines.health.europa.eu/veterinary/en/600000991053

Lincoral-S 222 mg/g + 444.7 mg/g powder for use in drinking water

Authorised

- Spectinomycin sulfate tetrahydrate
- Lincomycin hydrochloride monohydrate

Product identification

Medicine name:

Lincoral-S 222 mg/g + 444.7 mg/g powder for use in drinking water Lincoral-S (251,7 mg + 672,4 mg)/g Proszek do podania w wodzie do picia

Active substance:

Spectinomycin sulfate tetrahydrate

Lincomycin hydrochloride monohydrate

Target species:

Chicken

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Spectinomycin sulfate tetrahydrate 672.40 milligram(s) / 1.00 gram(s) Lincomycin hydrochloride monohydrate 251.70 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

Oral use:

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Chicken

- Meat and offal. 5 day

•

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FF52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

1.5 kg thermos-sealed bags made of polyethylene/aluminum/polyester 150 g thermos-sealed sachets made of polyethylene/aluminum/polyester

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

HuVepharma

Marketing authorisation date:

22/08/2023

Manufacturing sites for batch release:

Huvepharma S.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3268

Date of authorisation status change:

22/08/2023

Reference member state:

Ireland

Procedure number:

IE/V/0668/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain 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.

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