

Linspec 50/100 mg/ml Solution for injection for dogs, cats, pigs and pre-ruminant calves

Not
authorised

- Lincomycin hydrochloride
- Spectinomycin sulfate tetrahydrate

Product identification

Medicine name:

Linspec 50/100 mg/ml Solution for injection for dogs, cats, pigs and pre-ruminant calves

Active substance:

Lincomycin hydrochloride
Spectinomycin sulfate tetrahydrate

Target species:

Cattle
Dog
Cat
Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Lincomycin hydrochloride

54.49 milligram(s) / 1.00 millilitre(s)

Spectinomycin sulfate tetrahydrate

151.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 21 day

-

Pig

- Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FF52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

France

Package description:

250 ml multi-dose translucent polypropylene vial with bromobutyl stopper and aluminium cap with a flip off seal. Vials is placed in an outer container (paper box).

100 ml multi-dose translucent polypropylene vial with bromobutyl stopper and aluminium cap with a flip off seal. Vial is placed in an outer container (paper box).

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:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

4/10/2010

Manufacturing sites for batch release:

Cenavisa S.L.

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/4226186 9/2010

Date of authorisation status change:

20/09/2024

Reference member state:

Ireland

Procedure number:

IE/V/0238/001

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

Documents

Summary of Product Characteristics

English (PDF)

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