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
Linspec 50/100 mg/ml Solução injetável para suínos, vitelos pré-ruminantes, cães e gatos

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

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Authorisation status:
Revoked

Authorised in:
Portugal

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:
15/10/2013

Manufacturing sites for batch release:
Cenavisa S.L.
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
732/01/13DFVPT

Date of authorisation status change:
1/12/2015

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

Combined File of all Documents

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