SPIROVET 600 000 IU/ML SOLUTION FOR INJECTION FOR CATTLE

Not authorised

- Spiramycin

Product identification

Medicine name:
SPIROVET 600 000 IU/ML SOLUTION FOR INJECTION FOR CATTLE

Active substance:
Spiramycin

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Spiramycin
600000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**

- Milk. 14 day

Mastitis (30 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 100 kg bodyweight) twice at 24h of interval).

- Meat and offal. 75 day

Mastitis (30 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 100 kg bodyweight) twice at 24h of interval).

- Milk. no withdrawal period No withdrawal period

- Meat and offal. 75 day

Respiratory infections (100 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 30 kg bodyweight) twice at 48h of interval).

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Czechia

Package description:

Box containing 1 glass vial of 100 ml
Box containing 1 glass vial of 250 ml
Box containing 1 plastic vial of 50 ml
Box containing 1 plastic vial of 100 ml
Box containing 1 plastic vial of 250 ml

Box containing 1 glass vial of 50 ml

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:
Ceva Sante Animale

Marketing authorisation date:
17/09/2012

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/104/12-C

Date of authorisation status change:
17/09/2012

Reference member state:
France

Procedure number:
FR/V/0244/001

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

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

Labelling

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

Combined File of all Documents

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