Roxacin 100 mg/ml šķīdums injekcijām liellopiem un cūkām

Authorised

- Enrofloxacin

Product identification

Medicine name:
Roxacin 100 mg/ml šķīdums injekcijām liellopiem un cūkām

Active substance:
Enrofloxacin

Target species:
Cattle
Pig

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Enrofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Milk. 3 day
  - Meat and offal. 5 day

Subcutaneous use:

- **Cattle**
  - Milk. 4 day
  - Meat and offal. 12 day

Intramuscular use:

- **Pig**
  - Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

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Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Laboratorios Calier S.A.

Marketing authorisation date:
16/02/2001

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/01/1292

Date of authorisation status change:
18/02/2001

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

Documents

Summary of Product Characteristics

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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.
