Vetroxy LA 200 mg/ml solution for injection for cattle, sheep and pigs

Authorised

- Oxytetracycline dihydrate

Product identification

Medicine name:
Vetroxy LA 200 mg/ml solution for injection for cattle, sheep and pigs
Vetroxy LA 200 mg/ml šķīdums injekcijām liellopiem, aitām un cūkām

Active substance:
Oxytetracycline dihydrate

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Oxytetracycline dihydrate

216.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 10 day
  - Meat and offal. 31 day
- **Sheep**
  - Milk. 7 day
  - Meat and offal. 9 day
- **Pig**
  - Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Amber type II glass vials of 100 ml sealed with a bromobutyl rubber stopper with aluminium overseals and packaged individually into outer cartons.

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:
Bimeda Animal Health Limited

Marketing authorisation date:
30/11/2016

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/16/0040

Date of authorisation status change:
30/11/2016

Reference member state:
Netherlands

Procedure number:
NL/V/0253/001

Concerned member states:
Estonia Greece Hungary Latvia Poland Sweden
United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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