Diatrim 200 mg/ml + 40 mg/ml solution for injection for cattle, pigs, dogs and cats

Authorised

- Trimethoprim
- Sulfadiazine

Product identification

Medicine name:
Diatrim 200 mg/ml + 40 mg/ml solution for injection for cattle, pigs, dogs and cats
Diatrim 200 mg/ml Solution injectable

Active substance:
Trimethoprim
Sulfadiazine

Target species:
Cattle
Pig
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Trimethoprim
40.00 milligram(s) / 1.00 millilitre(s)

Sulfadiazine
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 48 hour
  - Meat and offal. 12 day
- **Pig**
  - Meat and offal. 20 day

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW10

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

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Authorisation status:
Valid

Authorised in:
Luxembourg

Package description:
1 vial in a cardboard box of uncoloured glass type II filled with 250 ml with a fluoropolymer coated chlorobutyl stopper type I secured with an aluminium cap.
1 vial in a cardboard box of uncoloured glass type II filled with 50 ml with a fluoropolymer coated chlorobutyl stopper type I secured with an aluminium cap.
1 vial in a cardboard box of uncoloured glass type II filled with 100 ml with a fluoropolymer coated chlorobutyl stopper type I secured with an aluminium cap.

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:
Dechra Regulatory B.V.

Marketing authorisation date:
14/03/2018

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V/914/18/03/1670

Date of authorisation status change:
14/03/2018

Reference member state:
Netherlands

Procedure number:
NL/V/0282/001

Concerned member states:
Austria Belgium Croatia France Germany Hungary Ireland Italy Lithuania Luxembourg Poland Portugal Slovenia Spain
United Kingdom (Northern Ireland)

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.

Combined File of all Documents

Diatrim - PuAR.pdf