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

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

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:
Poland

Available in:
Poland

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
30/03/2018

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2761

Date of authorisation status change:
30/03/2018

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

Package Leaflet

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

Summary of Product Characteristics

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

Diatrim - PuAR.pdf