

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

Authorisation status:

Valid

Authorised in:

Netherlands

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:

5/02/2018

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 120334

Date of authorisation status change:

24/01/2022

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

Combined File of all Documents

English (PDF)

Published on: 1/02/2024

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