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TAF SPRAY 28.5 MG/G CUTANEOUS SPRAY, SOLUTION

Authorised

- Thiamphenicol

Product identification

Medicine name:

TAF SPRAY 28.5 MG/G CUTANEOUS SPRAY, SOLUTION

Active substance:

Thiamphenicol

Target species:

Cattle

Pig

Mink

Rabbit

Equid

Sheep

Goat

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Thiamphenicol

28.50 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Cutaneous spray, solution

Withdrawal period by route of administration:

Cutaneous use:

-

Cattle

- Meat and offal. 0 day

- Milk. 0 day

Do not use on the udder of lactating animals if their milk is intended for human consumption.

-

Pig

- Meat and offal. 14 day

-

Rabbit

- Meat and offal. 0 day

-

Equid

- Meat and offal. 0 day

-

Sheep

- Meat and offal. 0 day

- Milk. 0 day

Do not use on the udder of lactating animals if their milk is intended for human consumption.

-

Goat

- Meat and offal. 0 day
- Milk. 0 day

Do not use on the udder of lactating animals if their milk is intended for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD06AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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:

16/02/2016

Manufacturing sites for batch release:

Eurovet Animal Health B.V.
IGS Aerosols GmbH

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2518

Date of authorisation status change:

16/02/2016

Reference member state:

France

Procedure number:

FR/V/0276/001

Concerned member states:

Austria Belgium Denmark Germany Ireland Italy Netherlands Poland
Portugal Spain Sweden 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.