

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:

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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.

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Pig

- Meat and offal. 14 day

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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.

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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:

Germany

Available in:

Germany

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:

3/12/2014

Manufacturing sites for batch release:

Eurovet Animal Health B.V.
IGS Aerosols GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402081.00.00

Date of authorisation status change:

28/01/2020

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

Summary of Product Characteristics

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

Package Leaflet

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

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