File downloaded on 2026-11-28
Source URL: https://medicines.health.europa.eu/veterinary/en/600000030094

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**
  - Milk. 0 day

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

  - Meat and offal. 0 day

- **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**

- Milk. 0 day

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

- Meat and offal. 0 day

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.

Combined File of all Documents

eu-puar-frv0276001-mr-rpe_169-en.pdf