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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)

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**Pharmaceutical form:**
Cutaneous spray, solution

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**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:**
Belgium

**Available in:**
Belgium

**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:**
Dechra Regulatory B.V.

**Marketing authorisation date:**
6/01/2015

**Manufacturing sites for batch release:**
Eurovet Animal Health B.V.
IGS Aerosols GmbH

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
BE-V467937

**Date of authorisation status change:**
6/01/2015

**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.

Labelling

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

Combined File of all Documents

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