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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

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**Active substance:**

Thiamphenicol

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**Target species:**

Cattle

Pig

Mink

Rabbit

Equid

Sheep

Goat

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**Route of administration:**

Cutaneous use

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## 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:**

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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

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**Equid**

- Meat and offal. 0 day

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**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.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QD06AX

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Available in:**

France

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Eurovet Animal Health B.V.

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**Marketing authorisation date:**

12/11/2014

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.  
IGS Aerosols GmbH

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/4917402 1/2014

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**Date of authorisation status change:**

24/10/2019

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**Reference member state:**

France

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**Procedure number:**

FR/V/0276/001

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**Concerned member states:**

Austria Belgium Denmark Germany Ireland Italy Netherlands Poland  
Portugal Spain Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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