

File downloaded on 2026-04-29

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000030147>

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:

Ireland

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:

9/01/2015

Manufacturing sites for batch release:

Eurovet Animal Health B.V.
IGS Aerosols GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10989/063/001

Date of authorisation status change:

9/01/2015

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

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