File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000030151

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:
United Kingdom (Northern 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:
13/01/2015

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 16849/4051

Date of authorisation status change:
26/11/2021

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

Combined File of all Documents

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