

Taneven 300 mg/ml suspension for injection for horses, cattle, sheep, goats, dogs and cats

Authorised

- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:

Taneven 300 mg/ml suspension for injection for horses, cattle, sheep, goats, dogs and cats

Active substance:

Benzylpenicillin (procaine) monohydrate

Target species:

Cattle

Dog

Goat

Sheep

Horse

Cat

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Benzylpenicillin (procaine) monohydrate
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

- Meat and offal. 10 day
- Milk. 120 hour

-

Goat

- Milk. 120 hour
- Meat and offal. 10 day

-

Sheep

- Meat and offal. 10 day
- Milk. 120 hour

-

Horse

- Meat and offal. 10 day
- Milk. no withdrawal period

Not authorised for use mares producing milk for human consumption.

Intramuscular use:

-

Cattle

- Meat and offal. 10 day
- Milk. 120 hour

•

Goat

- Milk. 120 hour
- Meat and offal. 10 day

•

Sheep

- Meat and offal. 10 day
- Milk. 120 hour

•

Horse

- Meat and offal. 10 day
- Milk. no withdrawal period

Not authorised for use mares producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

(ID1) 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 100 millilitre(s), closed with Stopper (Bromobutyl Rubber) and Cap (Aluminium)

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:

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Marketing authorisation date:

10/12/2020

Manufacturing sites for batch release:

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-3027

Date of authorisation status change:

10/12/2020

Reference member state:

Germany

Procedure number:

DE/V/0337/001

Concerned member states:

Austria Bulgaria Hungary Italy Luxembourg Poland Romania

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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Package Leaflet

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

Combined File of all Documents

Labelling

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