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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:
Poland

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:
1/02/2021

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3067

Date of authorisation status change:
1/02/2021

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

Combined File of all Documents

Summary of Product Characteristics

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Labelling

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.

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