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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:
Hungary

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:
12/11/2020

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
4215/X/20 NÉBIH ÁTI

Date of authorisation status change:
12/11/2020

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

Combined File of all Documents

2402721-paren-20201217.pdf