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
Taneven 300 mg/ml Injektionssuspension für Pferde, Rinder, Schafe, Ziege, Hunde, Katzen

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:
Germany

Available in:
Germany

Package description:

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

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:
27/10/2020

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402721.00.00

Date of authorisation status change:
27/10/2020

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

English (PDF)
Published on: 29/07/2025
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