Vitamin AD3E pro injectione, solution for injection for horses, cattle, pigs, and dogs

Not authorised

- DL-ALPHA TOCOPHEROL ACETATE
- COLECALCIFEROL CONCENTRATE (OILY FORM)
- Retinol palmitate

Product identification

Medicine name:
Vitamin AD3E pro injectione, solution for injection for horses, cattle, pigs, and dogs
Vitamin AD3E, Injektionslösung für Rind, Pferd, Schwein und Hund

Active substance:
DL-ALPHA TOCOPHEROL ACETATE
COLECALCIFEROL CONCENTRATE (OILY FORM)
Retinol palmitate

Target species:
Cattle
Dog
Horse
Pig

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

DL-ALPHA TOCOPHEROL ACETATE
50.00 milligram(s) / 1.00 millilitre(s)

COLECALCIFEROL CONCENTRATE (OILY FORM)
100.00 milligram(s) / 1.00 millilitre(s)

Retinol palmitate
176.46 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Milk. 120 hour
  - Meat and offal. 259 day
- **Horse**
  - Meat and offal. 250 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 194 day

Intramuscular use:

- **Cattle**
  - Milk. 120 hour

- Meat and offal. 259 day

- **Horse**
  - Meat and offal. 250 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 194 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

Legal status of supply:
Medicinal product on medical prescription for renewable delivery

Authorisation status:
Revoked

Authorised in:
Austria

Package description:
Available only in German
Available only in German
Available only in German

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:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
12/07/2019

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
839012

Date of authorisation status change:
12/07/2019

Reference member state:
Germany

Procedure number:
DE/V/0313/001

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

Documents

Combined File of all Documents

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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