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

Authorised

- all-rac-alfa-Tocopheryl acetate
- COLECALCIFEROL CONCENTRATE (OILY FORM)
- Retinol palmitate

Product identification

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

Active substance:
all-rac-alfa-Tocopheryl 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:

all-rac-alfa-Tocopheryl 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:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Portugal

Package description:
(ID1) 100 millilitre(s): unspecified outer container with 1 Bottle (brown glass) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)
(ID2) 600 millilitre(s): unspecified outer container with 6 Bottle (brown glass) each with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)
(ID3) 1200 millilitre(s): unspecified outer container with 12 Bottle (brown glass) each with 100 millilitre(s), closed with Stopfen (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:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
16/05/2019

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1267/01/19RFVPT

Date of authorisation status change:
23/08/2022

Reference member state:
Germany

Procedure number:
DE/V/0313/001

Concerned member states:
Austria Croatia Cyprus France Greece Iceland Ireland Italy Latvia Norway Portugal Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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