

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

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

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:

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Cattle

- Milk. 120 hour
- Meat and offal. 259 day

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Horse

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

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

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Pig

- Meat and offal. 194 day

Intramuscular use:

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Cattle

- Milk. 120 hour

- Meat and offal. 259 day

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Horse

- Meat and offal. 250 day

- Milk. no withdrawal period

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

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Pig

- Meat and offal. 194 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA11JA

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

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:

4/07/2019

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/19/0038

Date of authorisation status change:

4/07/2019

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

Summary of Product Characteristics

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.

Labelling

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

Combined File of all Documents

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