

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
VITAMIN AD3E PRO INJECTIONE

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:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

(ID3): 1 unspecified outer container with 12 Bottle (Glass) with 100 millilitre(s) (1200 millilitre(s))

(ID2): 1 unspecified outer container with 6 Bottle (Glass) with 100 millilitre(s) (600 millilitre(s))

(ID1): 1 unspecified outer container with 1 Bottle (Glass) with 100 millilitre(s) (100 millilitre(s))

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:

27/11/2019

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Ministry Of Health

Authorisation number:

105352

Date of authorisation status change:

27/11/2019

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

Combined File of all Documents

English (PDF)

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