VITAMIN AD3E, emulzija za injekciju, goveda, ovce, koze, konji, svinje, kunići, psi, mačke

Not authorised

- DL-ALPHA-TOCOPHEROL
- Colecalciferol
- Retinol palmitate

Product identification

Medicine name:
VITAMIN AD3E, emulzija za injekciju, goveda, ovce, koze, konji, svinje, kunići, psi, mačke

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Active substance:
DL-ALPHA-TOCOPHEROL

Colecalciferol

Retinol palmitate

Target species:
Cattle
Sheep
Goat
Horse
Pig
Rabbit
Dog
Cat
Cattle

Sheep
Goat
Horse
Pig
Rabbit
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
DL-ALPHA-TOCOPHEROL
20.00 milligram(s) / 1.00 millilitre(s)

Colecalciferol
25000.00 international unit(s) / 1.00 millilitre(s)

Retinol palmitate
50000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 243 day
- **Sheep**
  - Meat and offal. 187 day
- **Goat**
  - Meat and offal. 187 day
- **Horse**
  - Meat and offal. 243 day

- **Pig**
  - Meat and offal. 215 day
- **Rabbit**
  - Meat and offal. 122 day
- **Dog**
- **Cat**

Subcutaneous use:

- **Cattle**
  - Meat and offal. 243 day
- **Sheep**
  - Meat and offal. 187 day
- **Goat**
  - Meat and offal. 187 day
- **Horse**
  - Meat and offal. 243 day
- **Pig**
  - Meat and offal. 215 day
- **Rabbit**
  - Meat and offal. 122 day
- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
Croatia

Package description:
Available only in Croatian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Krka-Farma d.o.o.

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Krka d.d. Novo Mesto

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/17-01/509

Date of authorisation status change:
19/12/2023

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.
