VITAMIN AD3EC oralna emulzija, za goveda, konje, svinje, ovce, koze, kuniće, nerčeve, činčile, kokoši, purane, golubove

Not authorised

- Sodium ascorbate
- DL-ALPHA-TOCOPHEROL
- Colecalciferol
- Retinol palmitate

Product identification

Medicine name:
VITAMIN AD3EC oralna emulzija, za goveda, konje, svinje, ovce, koze, kuniće, nerčeve, činčile, kokoši, purane, golubove

Active substance:
Sodium ascorbate

DL-ALPHA-TOCOPHEROL

Colecalciferol

Retinol palmitate

Target species:
Cattle
Horse
Pig

Sheep
Goat
Rabbit
Mink
Chinchilla
Chicken
Turkey
Pigeon

Route of administration:
Oral use

Product details

Active substance and strength:
Sodium ascorbate
100.00 milligram(s) / 1.00 millilitre(s)

DL-ALPHA-TOCOPHEROL
30.00 milligram(s) / 1.00 millilitre(s)

Colecalciferol
5000.00 international unit(s) / 1.00 millilitre(s)

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

Pharmaceutical form:
Oral emulsion

Withdrawal period by route of administration:
Oral use:

- **Cattle**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Mink**
  - Meat and offal. 0 day
- **Chinchilla**
  - Meat and offal. 0 day
- **Chicken**
  - Eggs. 0 day
- **Turkey**
  - Meat and offal. 0 day
- **Pigeon**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

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

Authorisation status:
Revoked

Authorised in:
Croatia

Package description:
Available only in Croatian
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:
1/10/2020

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/20-01/603

Date of authorisation status change:
13/12/2024

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

Documents

Summary of Product Characteristics

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