Vitamin AD3E Krka emulzija za injiciranje za govedo, konje, prašiče, ovce, koze, kunce, pse in mačke

Not

authorised

- Retinol palmitate
- Colecalciferol
- ALPHATOCOPHEROL ACETATE

Product identification

Medicine name:

Vitamin AD3E Krka emulzija za injiciranje za govedo, konje, prašiče, ovce, koze, kunce, pse in mačke

Active substance:

Retinol palmitate Colecalciferol ALPHATOCOPHEROL ACETATE

Target species:

Cattle Horse Pig Sheep Goat

Rabbit

Dog

Cat

Product details

Active substance and strength:

Retinol palmitate 50000.00 international unit(s) / 1.00 millilitre(s) Colecalciferol 25000.00 international unit(s) / 1.00 millilitre(s) ALPHATOCOPHEROL ACETATE 20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration: Intramuscular use:

ntramuscular us

- Cattle
 - Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni
- Horse
 - Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni
- . Pig
 - Meat and offal. 0 day Meso in organi: 0 dni
- . Sheep
 - Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni
- . Goat
 - Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni
- . Rabbit

- Meat and offal. 0 day Meso in organi: 0 dni

- . Dog
- . Cat

Subcutaneous use:

. Cattle

- Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni

. Horse

- Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni

- Pig
 - Meat and offal. 0 day Meso in organi: 0 dni
- . Sheep
 - Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni
- . Goat
 - Meat and offal, milk. 0 day Meso in organi: 0 dni Mleko: 0 dni
- Rabbit
 - Meat and offal. 0 day Meso in organi: 0 dni
- . Dog
- . Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes: OA11BA

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Slovenia

Package description:

Available only in <u>Slovenian</u>

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder: KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date: 10/10/2002

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

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number: NP/V/0377/002

Date of authorisation status change:

2/10/2023

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

Documents

Package Leaflet and Labelling

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

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

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