

## VITAMIN-K/AGROSEED CANDILIDIS

Authorised

ενέσιμο διάλυμα 10 mg/ml για  
ιπποειδή, βοοειδή, μόσχους,  
χοίρους, σκύλο, γάτα

- Menadione sodium bisulfite

### Product identification

**Medicine name:**

VITAMIN-K/AGROSEED CANDILIDIS ενέσιμο διάλυμα 10 mg/ml για ιπποειδή, βοοειδή, μόσχους, χοίρους, σκύλο, γάτα

**Active substance:**

Menadione sodium bisulfite

**Target species:**

Cattle

Equid

Cattle (calf)

Pig

Dog

Cat

**Route of administration:**

Intramuscular and intravenous use

Subcutaneous use

## Product details

### Active substance and strength:

Menadione sodium bisulfite

10.00 milligram(s) / 1.00 millilitre(s)

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### Pharmaceutical form:

Solution for injection

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### Withdrawal period by route of administration:

#### Intramuscular and intravenous use:

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##### Cattle

- Meat and offal, milk. 0 day

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##### Equid

- Meat and offal, milk. 0 day

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##### Cattle (calf)

- Meat and offal. 0 day

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##### Pig

- Meat and offal. 0 day

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##### Dog

- Not applicable. no withdrawal period

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##### Cat

- Not applicable. no withdrawal period

### Subcutaneous use:

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**Cattle**

- Meat and offal, milk. 0 day

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**Equid**

- Meat and offal, milk. 0 day

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**Cattle (calf)**

- Meat and offal. 0 day

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**Pig**

- Meat and offal. 0 day

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**Dog**

- Not applicable. no withdrawal period

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**Cat**

- Not applicable. no withdrawal period

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QB02BA

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Package description:**

Available only in Greek

Available only in Greek

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Complete application (stand-alone)

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**Marketing authorisation holder:**

Candilidis S.A.

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**Marketing authorisation date:**

8/03/2000

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**Manufacturing sites for batch release:**

Alfasan International B.V.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

37813/05-05-2021/K-0130001

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**Date of authorisation status change:**

9/06/2022

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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Package Leaflet

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