

# Vitamin K1 10 mg/ml Solution for Injection

Authorised

- Phytomenadione

## Product identification

**Medicine name:**

Vitamin K1 10 mg/ml Solution for Injection

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**Active substance:**

Phytomenadione

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**Target species:**

Cattle

Dog

Sheep

Horse

Cat

Pig

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

### **Active substance and strength:**

Phytomenadione

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 use:**

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##### **Cattle**

- Meat and offal. 28 day

- Milk. 7 day

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##### **Sheep**

- Meat and offal. 28 day

- Milk. 7 day

- 

##### **Horse**

- Meat and offal. 28 day

- Milk. 7 day

- 

##### **Pig**

- Meat and offal. 28 day

#### **Intravenous use:**

- 

##### **Cattle**

- Meat and offal. 28 day

- Milk. 7 day

- 

**Sheep**

- Meat and offal. 28 day
- Milk. 7 day

- 

**Horse**

- Meat and offal. 28 day
- Milk. 7 day

- 

**Pig**

- Meat and offal. 28 day

**Subcutaneous use:**

- 

**Cattle**

- Meat and offal. 28 day
- Milk. 7 day

- 

**Sheep**

- Meat and offal. 28 day
- Milk. 7 day

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**Horse**

- Meat and offal. 28 day
- Milk. 7 day

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**Pig**

- Meat and offal. 28 day

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

QB02

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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:**

Ireland

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**Package description:**

50 ml amber Type II glass, multidose vial sealed with a bromobutyl rubber bung and capped with an aluminium cap containing a clear to slightly opalescent, pale yellow, aqueous, colloidal solution for injection.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Complete application (stand-alone) - Council Directive 81/851/EEC

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**Marketing authorisation holder:**

Bimeda Animal Health Limited

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**Marketing authorisation date:**

1/10/1991

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**Manufacturing sites for batch release:**

Bimeda Animal Health Limited

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA22033/048/001

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**Date of authorisation status change:**

1/10/1991

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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