Vitamin K1 10 mg/ml Solution for Injection

Authorised

- Phytomenadione

Product identification

Medicine name:
Vitamin K1 10 mg/ml Solution for Injection

Active substance:
Phytomenadione

Target species:
Cattle
Dog
Sheep
Horse
Cat
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Phytomenadione
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Dog**
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Horse**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Cat**
- **Pig**
  - Meat and offal. 28 day

Intravenous use:

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

- **Dog**
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Horse**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Cat**
- **Pig**
  - Meat and offal. 28 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Dog**
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Horse**
  - Meat and offal. 28 day
  - Milk. 7 day

- **Cat**
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QB02

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

Authorisation status:
Valid

Authorised in:
Ireland

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
1/10/1991

Manufacturing sites for batch release:

Bimeda Animal Health Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22033/048/001

Date of authorisation status change:
1/10/1991

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

Documents

Summary of Product Characteristics

Source URL: *https://medicines.health.europa.eu/veterinary/600000064224*