Noromectin Injection 10 mg/ml Oplossing voor injectie

Authorised

- Ivermectin

Product identification

Medicine name:
Noromectin Injection 10 mg/ml Oplossing voor injectie
Noromectin Injection 10 mg/ml Solution injectable
Noromectin Injection 10 mg/ml Injektionslösung

Active substance:
Ivermectin

Target species:
Pig
Cattle
Cattle (lactating cow)

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pig**
  - Meat and offal. 18 day

- **Cattle**
  - Milk. no withdrawal period

NL - Niet gebruiken bij melkkoeien tijdens de lactatie, noch tijdens de droogstand wanneer de melk bestemd is voor humaan gebruik. Niet gebruiken bij vaarzen binnen 60 dagen voor het kalven. FR - Ne pas utiliser chez les vaches laitières pendant la lactation, ni pendant la période de tarissement quand le lait est destiné à un usage pour l’homme. Ne pas utiliser chez les génisses durant les 60 jours qui précèdent le vêlage.

  - Meat and offal. 49 day

- **Cattle (lactating cow)**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:

Noromectin Injection 1 l Vial Solution for injection
Noromectin Injection 500 ml Vial Solution for injection
Noromectin Injection 250 ml Vial Solution for injection
Noromectin Injection 100 ml Vial Solution for injection
Noromectin Injection 50 ml Vial Solution for injection

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
9/12/2002

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V241525

Date of authorisation status change:
10/03/2020

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

Documents

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

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