Dectomax Inspuitbare Oplossing 10 mg/ml Oplossing voor injectie

Not authorised

- Doramectin

Product identification

Medicine name:
Dectomax Inspuitbare Oplossing 10 mg/ml Oplossing voor injectie

Active substance:
Doramectin

Target species:
Sheep
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

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

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Sheep**

  - Meat and offal. 70 day

  - Milk. no withdrawal period

Not permitted for use in lactating animals producing milk for human consumption. Do not use in pregnant ewes, which are intended to produce milk for human consumption, within 70 days of expected parturition.

Subcutaneous use:

- **Cattle**

  - Milk. no withdrawal period

Not permitted for use in lactating animals producing milk for human consumption. Do not use in pregnant cows or heifers, which are intended to produce milk for human consumption, within 2 months of expected parturition.

  - Meat and offal. 70 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Belgium

Package description:

Dectomax Solution for Injection 250 ml Vial Solution for injection

Dectomax Solution for Injection 500 ml Vial Solution for injection

Dectomax Solution for Injection 200 ml Vial Solution for injection
Dectomax Solution for 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:
Elanco GmbH

Marketing authorisation date:
30/10/1995

Manufacturing sites for batch release:
Elanco France S.A.S.
Norbrook Laboratories Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V172365

Date of authorisation status change:
7/08/2024

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.