

DECTOMAX 10 mg/ml Solution for Injection for Cattle, Sheep and Pigs

Authorised

- Doramectin

Product identification

Medicine name:

DECTOMAX 10 mg/ml Solution for Injection for Cattle, Sheep and Pigs
Dectomax 10 mg/ml Injektionslösung für Rinder, Schafe und Schweine

Active substance:

Doramectin

Target species:

Sheep

Pig

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

-

Pig

- Meat and offal. 77 day

Subcutaneous use:

-

Cattle

- 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:

Valid

Authorised in:

Austria

Package description:

Available only in German

Available only in German

Available only in German

Available only in German

The product is supplied in 50 ml multi-dose Type III amber glass vials with chlorobutyl rubber stoppers and aluminium overcaps.

The product is supplied in 200 ml multi-dose Type III amber glass vials with chlorobutyl rubberstoppers and aluminium overcaps.

The product is supplied in 500 ml multi-dose Type III amber glass vials with chlorobutyl rubber stoppers and aluminium overcaps.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis Österreich GmbH

Marketing authorisation date:

22/08/2012

Manufacturing sites for batch release:

Zoetis Belgium

Zoetis Manufacturing & Research Spain S.L.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-01110

Date of authorisation status change:

22/08/2012

Reference member state:

Ireland

Procedure number:

IE/V/0260/001

Concerned member states:

Austria Bulgaria Croatia Cyprus Czechia Estonia France Greece Hungary Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

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Labelling

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Package Leaflet

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