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

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:
Hungary

Package description:
The product is supplied in 50 ml multi-dose Type II amber glass vials with chlorobutyl rubber stoppers and aluminium overcaps.
The product is supplied in 200 ml multi-dose Type II amber glass vials with chlorobutyl rubberstoppers and aluminium overcaps.

The product is supplied in 500 ml multi-dose Type II amber glass vials with chlorobutyl rubber stoppers and aluminium overcaps.
The product is supplied in 250 ml multi-dose Type II amber glass vials with chlorobutyl rubber stoppers and aluminium overcaps.
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 Hungary Kft.

Marketing authorisation date:
25/07/2012

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
3192/X/12 NÉBIH ÁTI

Date of authorisation status change:
25/07/2012

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