Dectomax 5 mg/ml Pour-On Solution for Cattle

Authorised

- Doramectin

Product identification

Medicine name:
Dectomax 5 mg/ml Pour-On Solution for Cattle

Active substance:
Doramectin

Target species:
Cattle

Route of administration:
Pour-on use

Product details

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

Pharmaceutical form:
Pour-on solution

Withdrawal period by route of administration:

Pour-on use:

- **Cattle**
  - Meat and offal. 35 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA03

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

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
The product will be supplied in:- 250 ml multi-dose high-density polyethylene bottles with screw-top lidsand dosing cups in a carton box
The product will be supplied in:- 1 L multi-dose high-density polyethylene bottles with screw-top lidsand dosing cups in a carton box
The product will be supplied in:- 2.5 L multi-dose high-density polyethylene bottles with screw-top lidsand draw-off adaptor in a carton box.
The product will be supplied in:-3 L multi-dose high-density polyethylene bottles with screw-top lidsand draw-off adaptor in a carton box.
.The product will be supplied in:-5 L multi-dose high-density polyethylene bottles with screw-top lidsand draw-off adaptor in a carton box.

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 Animal Health ApS

Marketing authorisation date:
18/07/2013

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
10-7389

Date of authorisation status change:
18/04/2016

Reference member state:
Ireland

Procedure number:
IE/V/0260/002

Concerned member states:
Austria Denmark France Netherlands Norway Portugal Sweden

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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Package Leaflet

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