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Eprizero 5mg/ml Pour-On Solution for Beef and Dairy Cattle

Authorised

- Eprinomectin

Product identification

Medicine name:
Eprizero 5mg/ml Pour-On Solution for Beef and Dairy Cattle

Active substance:
Eprinomectin

Target species:
Cattle (for meat production)
Cattle (dairy cow)

Route of administration:
Pour-on use

Product details

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

Pharmaceutical form:
Pour-on solution

Withdrawal period by route of administration:

Pour-on use:

- **Cattle (for meat production)**
  - Meat and offal. 10 day

- **Cattle (dairy cow)**
  - Meat and offal. 10 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA04

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

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID5) 1 litre(s): unspecified outer container with 1 Barrel (High Density PolyEthylene) with 1 litre(s)
(ID4) 5 litre(s): unspecified outer container with 1 Barrel (High Density PolyEthylene) with 5 litre(s)
(ID3) 2.5 litre(s): unspecified outer container with 1 Barrel (High Density PolyEthylene) with 2.5 litre(s)
(ID2) 1 litre(s): unspecified outer container with 1 Container (High Density PolyEthylene) with 1 litre(s), closed with Cap (High Density PolyEthylene)
(ID1) 250 millilitre(s): unspecified outer container with 1 Container (High Density PolyEthylene) with 250 millilitre(s), closed with Cap (High Density PolyEthylene)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
29/07/2014

Manufacturing sites for batch release:
Norbrook Laboratories (Ireland) Limited
Norbrook Laboratories Limited

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401700.00.00

Date of authorisation status change:
20/09/2018

Reference member state:
Germany

Procedure number:
DE/V/0327/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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