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Taurador 5 mg/ml Pour-on Solution for Cattle

Authorised

- Doramectin

Product identification

Medicine name:
Taurador 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

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Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

The veterinary product will be supplied in:250mL standard high density polyethylene bottles with 28mm polypropylene/high density polyethylene caps.
The veterinary product will be supplied in:1L standard high density polyethylene bottles with 28mm polypropylene/high density polyethylene caps.
The veterinary product will be supplied in:1L white flat bottomed heavy duty high density polyethylene back-packs with 38mm white polypropylene easy peel caps.
The veterinary product will be supplied in:2.5L white flat bottomed heavy duty high density polyethylene back-packs with 38mm white polypropylene easy peel caps.
The veterinary product will be supplied in:5L white flat bottomed heavy duty high density polyethylene back-packs with 38mm white polypropylene easy peel caps.
The veterinary product will be supplied in:10L white high density polyethylene Jerry cans with high density polyethylene caps.
The veterinary product will be supplied in:20L white high density polyethylene Jerry cans with high density polyethylene caps.

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
29/11/2013

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/111/001

Date of authorisation status change:
29/11/2013

Reference member state:
Ireland

Procedure number:
IE/V/0617/001

Concerned member states:
Austria Belgium France Luxembourg Netherlands Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents