

Epromec 5 mg/ml Pour-on Solution for beef and dairy cattle

Authorised

- Eprinomectin

Product identification

Medicine name:

Epromec 5 mg/ml Pour-on Solution for beef and dairy cattle

Active substance:

Eprinomectin

Target species:

Cattle

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:**

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Cattle

- Meat and offal. 15 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:5L 'Flexi' packs.Pack sizes 5 L.

High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:3L 'Flexi' packs.Pack sizes 3L.

High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:2.5 L 'Flexi' packs.Pack sizes 2.5L.

High density polyethylene container with a polypropylene tamper evident screw cap which consists of:1L 'Squeeze pour' packs.Pack sizes 1L.

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:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

22/05/2015

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10987/107/001

Date of authorisation status change:

22/05/2015

Reference member state:

Ireland

Procedure number:

IE/V/0355/001

Concerned member states:

France Italy Netherlands Poland 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