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

Authorised

- Eprinomectin

Product identification

Medicine name:
Unomec 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:

- **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:
Poland

Package description:
High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:1L 'Squeeze pour' packs.Pack sizes 1L.
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 the following:3L 'Flexi' packs.Pack sizes 3L.
High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:5L 'Flexi' packs.Pack sizes 5 L.

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:
14/12/2017

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2731

Date of authorisation status change:
14/12/2017

Reference member state:
Ireland

Procedure number:
IE/V/0354/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Finland France Germany Greece Hungary Italy Netherlands Norway Poland Portugal Romania Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 12/04/2026
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Package Leaflet

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Labelling

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Combined File of all Documents