

Eprivalan 5 mg/ml pour-on solution for cattle

Authorised

- Eprinomectin

Product identification

Medicine name:

Eprivalan 5 mg/ml pour-on solution for 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:

Croatia

Package description:

HDPE bottle (250 ml) with induction sealed tamper evident HDPE screw on cap, containing a screw on dosing device capable of delivering measured amounts of product. The 250 ml bottle uses a 25 ml dispenser (screw-on-squeeze-and-pour measuring chamber).

HDPE bottle (1 litre) with induction sealed tamper evident HDPE screw on cap, containing a screw on dosing device capable of delivering measured amounts of product. The 1 litre bottle uses a 50 ml dispenser (screw-on-squeeze-and-pour measuring chamber).

Back-pack (2.5 litre) with induction sealed tamper evident HDPE screw on cap, containing a screw on dosing device capable of delivering measured amounts of product. The 2.5 litre back-packs are designed for use with a suitable automatic dispensing gun.

Back-pack (5 litre) with induction sealed tamper evident HDPE screw on cap, containing a screw on dosing device capable of delivering measured amounts of product. The 5 litre back-packs are designed for use with a suitable automatic dispensing gun.

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:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

15/03/2017

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/21-01/664

Date of authorisation status change:

24/09/2025

Reference member state:

Ireland

Procedure number:

IE/V/0487/001

Concerned member states:

Bulgaria Croatia Italy Luxembourg Romania Slovakia 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

English (PDF)

Published on: 20/07/2025

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Combined File of all Documents

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