Pereprin 5 mg/ml pour on solution for cattle, sheep and goats

Authorised

- Eprinomectin

Product identification

Medicine name:
Pereprin 5 mg/ml pour on solution for cattle, sheep and goats

Active substance:
Eprinomectin

Target species:
Cattle
Goat
Sheep

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
- **Goat**
  - Meat and offal. 1 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 2 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:
Portugal

Package description:
White high-density polyethylene 5 L bottle closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 5 L bottle per cardboard box together with a spigot cap.
White high-density polyethylene backpack bottles of 1 L closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 1 L bottle per cardboard box together with a spigot cap

and backpack strap.
White high-density polyethylene 250 ml bottle closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 250 ml bottle per cardboard box together with a spigot cap.
White high-density polyethylene backpack bottles of 2.5 L closed with black screw closure made of polypropylene with a polyethylene-lined sealing disk (wadding) for induction sealing inside. One 2.5 L bottle per cardboard box together with a spigot cap and backpack strap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
22/12/2025

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1764/01/25DFVPT

Date of authorisation status change:
22/12/2025

Reference member state:
Ireland

Procedure number:

IE/V/0607/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Germany Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

Generic of:

600000047578

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Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics

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