

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:

Greece

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:

4/05/2026

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

National Organization For Medicines

Authorisation number:

55304/05-05-2026/K-0263901

Date of authorisation status change:

24/02/2026

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

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