

Noromectin 10 mg/ml Solution for Injection for sheep.

Authorised

- Ivermectin

Product identification

Medicine name:

Noromectin 10 mg/ml Solution for Injection for sheep.

Active substance:

Ivermectin

Target species:

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Ivermectin

10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Subcutaneous use:**

-

Sheep

- Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

The product will be supplied as a colourless to pale yellow solution in 1 L volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap. The product will be supplied as a colourless to pale yellow solution in 500 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap. The product will be supplied as a colourless to pale yellow solution in 250 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap. The product will be supplied as a colourless to pale yellow solution in 100 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap. The product will be supplied as a colourless to pale yellow solution in 50 ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

29/06/2001

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22664/059/001

Date of authorisation status change:

29/06/2001

Reference member state:

Ireland

Procedure number:

IE/V/0168/001

Concerned member states:

Italy Portugal Spain

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