Noromectin 18.7 mg/g Oral Paste for Horses

Authorised

- Ivermectin

Product identification

Medicine name:
Noromectin 18.7 mg/g Oral Paste for Horses
Noromectin 18.7 mg/g Pasta voor oraal gebruik
Noromectin 18.7 mg/g Pâte orale
Noromectin 18.7 mg/g Paste zum Einnehmen

Active substance:
Ivermectin

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Ivermectin
1.87 gram(s) / 100.00 gram(s)

Pharmaceutical form:
Oral paste

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 34 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Low-density polyethylene pre-filled syringes containing 7.49 g of product in cartons of 1 syringe.
Low-density polyethylene pre-filled syringes containing 7.49 g of product in cartons of 2 syringes.
Low-density polyethylene pre-filled syringes containing 7.49 g of product in cartons of 10 syringes.
Low-density polyethylene pre-filled syringes containing 7.49 g of product in cartons of 50 syringes.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
13/01/2003

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V243135

Date of authorisation status change:
13/01/2003

Reference member state:
Ireland

Procedure number:
IE/V/0124/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Iceland Italy Luxembourg Netherlands Portugal 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: 13/04/2025
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Package Leaflet

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Combined File of all Documents

Labelling

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