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Equinor 370 mg/g Oral Paste for Horses

Authorised

- Omeprazole

Product identification

Medicine name:
Equinor 370 mg/g Oral Paste for Horses

Active substance:
Omeprazole

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Omeprazole
370.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral paste

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA02BC01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Norway

Available in:

Norway

Package description:

Immediate packageImmediate packaging: 7 ml oral syringe containing 7.57 g of paste composed of polyethylene barrel, plunger and end cap, withpolypropylene dosing rings.Outer package and sales presentations- Carton box of 1 oral syringe Immediate packageImmediate packaging: 7 ml oral syringe containing 7.57 g of paste composed of polyethylene barrel, plunger and end cap, withpolypropylene dosing rings.Outer package and sales presentations- Carton box of 7 oral syringes Immediate packageImmediate packaging: 7 ml oral syringe containing 7.57 g of paste composed of polyethylene barrel, plunger and end cap, withpolypropylene dosing rings.Outer package and sales presentations- Bucket of 72 oral syringes. Immediate package Immediate packaging: 7 ml oral syringe containing 7.57 g of paste composed of polyethylene barrel, plunger and end cap, with polypropylene dosing rings. Outer package and sales presentations - Carton box of 14 oral syringes Immediate package Immediate packaging: 7 ml oral syringe containing 7.57 g of paste composed of polyethylene barrel, plunger and end cap, with polypropylene dosing rings. Outer package and sales presentations - Carton box of 21 oral syringes

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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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
16/03/2018

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

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
17-12023

Date of authorisation status change:
16/03/2018

Reference member state:
Ireland

Procedure number:
IE/V/0308/001

Concerned member states:
Austria Belgium Denmark France Germany Netherlands Norway Poland 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: 12/04/2026
Download

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

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