

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

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

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 1 oral syringe

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 7 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- 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

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:

11/09/2013

Manufacturing sites for batch release:

Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:

Danish Medicines Agency

Authorisation number:

51170

Date of authorisation status change:

11/09/2013

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)

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