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GastroGard 370 mg/g oral paste

Authorised

• Omeprazole

Product identification

Medicine name:

GastroGard 370 mg/g oral paste
GASTROGARD 370 MG/G PATE ORALE POUR CHEVAUX

Active substance:

Omeprazole

Target species:

Horse

Route of administration:

Oral use

Product details

Active substance and strength:

Omeprazole 2.28 gram(s) / 1.00 Syringe

Pharmaceutical form:

Oral paste

Withdrawal period by route of administration:

Oral use:

•

Horse

- Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

OA02BC01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Immediate packaging: 10 ml syringe containing 6.16 g of paste composed of white polypropylene syringes barrel with a white LDPE cap, a rubber rod tip and a polypropylene plunger rod, with dose divisions calibrated by body weight. Outer packaging and sales presentations-Bulk pack of 72 syringes Immediate packaging: 10 ml syringe containing 6.16 g of paste composed of white polypropylene syringes barrel with a white LDPE cap, a rubber rod tip and a polypropylene plunger rod, with dose divisions calibrated by body weight. Outer packaging and sales presentations-Carton box of 14 syringes Immediate packaging: 10 ml syringe containing 6.16 g of paste composed of white polypropylene syringes barrel with a white LDPE cap, a rubber rod tip and a polypropylene plunger rod, with dose divisions calibrated by body weight. Outer packaging and sales presentations-Carton box of 7 syringes Immediate packaging: 10 ml syringe containing 6.16 g of paste composed of white polypropylene syringes barrel with a white LDPE cap, a rubber rod tip and a polypropylene plunger rod, with dose divisions calibrated by body weight. Outer packaging and sales presentations-Carton box of 1 syringe

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

2/02/2004

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1026735 8/2004

Date of authorisation status change:

2/02/2004

Reference member state:

Ireland

Procedure number:

IE/V/0489/001

Concerned member states:

Austria Belgium Denmark Finland France Germany Greece Italy Luxembourg Netherlands Norway 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: 30/03/2025

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Package Leaflet and Labelling

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