Omeproshield 370 mg/g oral paste for horses

Authorised

- Omeprazole

Product identification

Medicine name:
Omeproshield 370 mg/g oral paste for horses
Omeproshield 370 mg/g pasta voor oraal gebruik voor paarden

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:

QA02BC01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:
12/06/2015

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France SCS

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 114950

Date of authorisation status change:
24/02/2022

Reference member state:
Ireland

Procedure number:
IE/V/0488/001

Concerned member states:
Austria Belgium Germany Italy Luxembourg Netherlands Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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Summary of Product Characteristics

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