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# GastroGard 370 mg/g oral paste

Authorised

- Omeprazole

## Product identification

**Medicine name:**
GastroGard 370 mg/g oral paste

**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QA02BC01

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**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

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**Authorisation status:**
Valid

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**Authorised in:**
France

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**Available in:**
France

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**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-Carton box of 1 syringe
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 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-Bulk pack of 72 syringes

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## 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: 9/08/2026
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Package Leaflet and Labelling

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