

# 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 orale per cavalli

**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:**

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**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

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Package description:**

Immediate 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 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

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**Additional information****Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

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**Marketing authorisation date:**

3/07/2015

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France SCS

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

104738

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**Date of authorisation status change:**

3/07/2015

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0488/001

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**Concerned member states:**

Austria Belgium Germany Italy Luxembourg Netherlands Spain  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 2/07/2025

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Summary of Product Characteristics

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