

# Equizol 400 mg gastro-resistant granules for horses

Authorised

- Omeprazole

## Product identification

**Medicine name:**

Equizol 400 mg gastro-resistant granules for horses

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**Active substance:**

Omeprazole

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**Target species:**

Horse

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Omeprazole

80.00 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Gastro-resistant granules

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**Withdrawal period by route of administration:****Oral use:**

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**Horse**

- Meat and offal. 2 day
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

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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:**

Netherlands

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**Available in:**

Netherlands

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**Package description:**

(ID7) 1000 gram(s): Box (cardboard) with 200 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

(ID6) 560 gram(s): Box (cardboard) with 112 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

(ID5) 500 gram(s): Box (cardboard) with 100 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

(ID4) 420 gram(s): Box (cardboard) with 84 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

(ID3) 280 gram(s): Box (cardboard) with 56 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

(ID2) 140 gram(s): Box (cardboard) with 28 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

(ID1) 70 gram(s): Box (cardboard) with 14 Bag (polyethylene; aluminium; paper) each with 5 gram(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

CP-Pharma Handelsgesellschaft mbH

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**Marketing authorisation date:**

17/05/2018

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**Manufacturing sites for batch release:**

CP-Pharma Handelsgesellschaft mbH

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 119732

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**Date of authorisation status change:**

7/02/2022

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0318/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Finland France Hungary Ireland Italy  
Netherlands Norway Poland Portugal Slovakia Spain Sweden  
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)

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