

# EQVALAN DUO ORAL PASTE

Authorised

- Ivermectin
- Praziquantel

## Product identification

**Medicine name:**

EQVALAN DUO ORAL PASTE

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**Active substance:**

Ivermectin

Praziquantel

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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:**

Ivermectin

15.50 milligram(s) / 1.00 gram(s)

Praziquantel

77.50 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Oral paste

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**Withdrawal period by route of administration:**

**Oral use:**

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**Horse**

- Meat and offal. 30 day
- Milk. no withdrawal period

Do not use in mares producing milk for human consumption.

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

QP54AA51

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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:**

Spain

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**Available in:**

Spain

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**Package description:**

- Box of 1 sachet of 1 syringe of 7,74 g
  - Box of 50 sachet of 1 syringe of 14,19 g
  - Box of 50 sachets of 1 syringe of 9,68 g
  - Box of 50 sachets of 1 syringe of 7,74 g
  - Box of 1 sachet of 1 syringe of 14,19 g
  - Box of 1 sachet of 1 syringe of 9,68 g
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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health Espana S.A.

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**Marketing authorisation date:**

3/11/2004

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

1596 ESP

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**Date of authorisation status change:**

3/11/2004

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**Reference member state:**

France

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**Procedure number:**

FR/V/0359/001

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**Concerned member states:**

Austria Belgium Cyprus Denmark Estonia Finland Germany Greece  
Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands  
Norway Portugal Slovenia 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

### Summary of Product Characteristics

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

### Package Leaflet

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

### Labelling

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

### Combined File of all Documents