

# EQVALAN Oral Paste for Horses

Authorised

## 18.7 mg/g

- Ivermectin

## Product identification

**Medicine name:**

EQVALAN Oral Paste for Horses 18.7 mg/g

---

**Active substance:**

Ivermectin

---

**Target species:**

Horse

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Ivermectin

18.70 milligram(s) / 1.00 gram(s)

---

**Pharmaceutical form:**

Oral paste

---

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

- 

**Horse**

- Meat and offal. 21 day

---

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

QP54AA01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Ireland

---

**Package description:**

Dose graduated disposable polypropylene syringe containing 6.42 g of a white homogenous paste. Pack of 1 syringe.

Dose graduated disposable polypropylene syringe containing 6.42 g of a white homogenous paste. Pack of 50 syringes.

Dose graduated disposable polypropylene syringe containing 8.03 g of a white homogenous paste. Pack of 1 syringe.

Dose graduated disposable polypropylene syringe containing 8.03 g of a white homogenous paste. Pack of 50 syringes.

Dose graduated disposable polypropylene syringe containing 11.77 g of a white homogenous paste. Pack of 1 syringe.

Dose graduated disposable polypropylene syringe containing 11.77 g of a white homogenous paste. Pack of 50 syringes.

---

## 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 Vetmedica GmbH

---

**Marketing authorisation date:**

1/10/2000

---

**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France SCS

---

**Responsible authority:**

Health Products Regulatory Authority

---

**Authorisation number:**

VPA10454/037/001

---

**Date of authorisation status change:**

1/10/2000

---

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