**Source URL:** https://medicines.health.europa.eu/veterinary/en/60000050726

# Equimax oral gel for horses

Authorised

- Ivermectin
- Praziquantel

### Product identification

#### **Medicine name:**

Equimax oral gel for horses Equimax

#### **Active substance:**

**Ivermectin** 

Praziquantel

### **Target species:**

Horse

#### Route of administration:

Oral use

# **Product details**

## **Active substance and strength:**

**Ivermectin** 

18.70 milligram(s) / 1.00 gram(s)

Praziquantel

140.30 milligram(s) / 1.00 gram(s)

#### **Pharmaceutical form:**

Oral gel

### Withdrawal period by route of administration:

#### Oral use:

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#### Horse

- Meat and offal. 35 day

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

**QP54AA51** 

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Germany

### Package description:

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 7.49 grams of product and is fitted with variable dose capacity. Product presentations: Box of 48 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 7.49 grams of product and is fitted with variable dose capacity. Product presentations: Box of 40 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 7.49 grams of product and is fitted with variable dose capacity. Product presentations: Box of 12 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 7.49 grams of product and is fitted with variable dose capacity. Product presentations: Box of 2 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 7.49 grams of product and is fitted with variable dose capacity. Product presentations: Box of 1 syringe or Blister of

one syringe.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 6.42 grams of product and is fitted with variable dose capacity. Product presentations: Box of 48 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 6.42 grams of product and is fitted with variable dose capacity. Product presentations: Box of 40 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 6.42 grams of product and is fitted with variable dose capacity. Product presentations: Box of 12 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 6.42 grams of product and is fitted with variable dose capacity. Product presentations: Box of 2 syringes.

An adjustable multidose syringe consisting of high density polyethylene (white) and low density polyethylene (white). The syringe contains 6.42 grams of product and is fitted with variable dose capacity. Product presentations: Box of 1 syringe or Blister of one syringe.

### Additional information

### **Entitlement type:**

Marketing Authorisation

### Legal basis of product authorisation:

Complete application (stand-alone) - Council Directive 81/851/EEC

### Marketing authorisation holder:

Virbac

### Marketing authorisation date:

8/02/2002

### Manufacturing sites for batch release:

Sofarimex-Industria Quimica E Farmaceutica S.A.

Virbac

### **Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

#### **Authorisation number:**

400551.00.00

#### Date of authorisation status change:

5/07/2007

#### **Reference member state:**

Ireland

#### **Procedure number:**

IE/V/0501/001

#### **Concerned member states:**

Austria Belgium Czechia Denmark Finland France Germany Greece Hungary Italy Luxembourg Netherlands Norway Poland Portugal Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

### **Documents**

Combined File of all Documents

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