

# EQUEST PRAMOX 19.5 MG/G + 121.7 MG/G ORAL GEL

Authorised

- Moxidectin
- Praziquantel

## Product identification

### **Medicine name:**

EQUEST PRAMOX 19.5 MG/G + 121.7 MG/G ORAL GEL

EQUEST PRAMOX 19,5 mg/g + 121,7 mg/g GEL ORAL PARA CABALLOS

---

### **Active substance:**

Moxidectin

Praziquantel

---

### **Target species:**

Horse (mare)

Horse

---

### **Route of administration:**

Oral use

---

## Product details

### **Active substance and strength:**

Moxidectin

19.50 milligram(s) / 1.00 gram(s)

Praziquantel  
121.70 milligram(s) / 1.00 gram(s)

---

**Pharmaceutical form:**

Oral gel

---

**Withdrawal period by route of administration:**

**Oral use:**

- 

**Horse (mare)**

- Milk. no withdrawal period

Not permitted for use in lactating mares producing milk for human consumption.

- 

**Horse**

- Meat and offal. 64 day

---

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

QP54AB52

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Spain

---

**Available in:**

Spain

---

**Package description:**

Box containing 1 individually boxed syringes

Box containing 20 syringes

Box containing 20 individually boxed syringes

Box containing 10 individually boxed syringes

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

---

**Marketing authorisation holder:**

Zoetis Spain S.L.

---

**Marketing authorisation date:**

21/06/2006

---

**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

---

**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

---

**Authorisation number:**

1698 ESP

---

**Date of authorisation status change:**

21/06/2006

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0161/001

---

**Concerned member states:**

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

---

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.