

# OESTRACTON 52.4

## micrograms/ml solution for injection for cattle, horses, pigs

Authorised

- Gonadorelin (6-D-phenylalanine) acetate

### Product identification

**Medicine name:**

OESTRACTON 52.4 micrograms/ml solution for injection for cattle, horses, pigs  
Oestracton 52,4 microgram/ml oplossing voor injectie voor runderen, paarden, varkens

**Active substance:**

Gonadorelin (6-D-phenylalanine) acetate

**Target species:**

Pig  
Cattle  
Horse

**Route of administration:**

Subcutaneous use  
Intramuscular use

## Product details

### Active substance and strength:

Gonadorelin (6-D-phenylalanine) acetate  
52.40 microgram(s) / 1.00 millilitre(s)

---

### Pharmaceutical form:

Solution for injection

---

### Withdrawal period by route of administration:

#### Subcutaneous use:

- 

##### Pig

- Meat and offal. 0 day

#### Intramuscular use:

- 

##### Cattle

- Meat and offal. 0 day
- Milk. 0 hour

- 

##### Horse

- Meat and offal. 0 day
- Milk. 0 hour

- 

##### Pig

- Meat and offal. 0 day

---

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

QH01CA01

---

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Netherlands

---

**Package description:**

(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)

(ID2) 60 millilitre(s): unspecified outer container with 6 Vial (Glass) each with 10 millilitre(s)

(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (Glass) with 10 millilitre(s)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Vetoquinol S.A.

---

**Marketing authorisation date:**

6/01/2014

---

**Manufacturing sites for batch release:**

Wirtschaftsgenossenschaft deutscher Tieraerzte e G

---

**Responsible authority:**

Medicines Evaluation Board

---

**Authorisation number:**

REG NL 112668

---

**Date of authorisation status change:**

8/02/2022

---

**Reference member state:**

Germany

---

**Procedure number:**

DE/V/0154/001

---

**Concerned member states:**

Belgium Czechia France Hungary Ireland Netherlands

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

Combined File of all Documents

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