

# 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)

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**Pharmaceutical form:**

Solution for injection

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

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**Pig**

- Meat and offal. 0 day

**Intramuscular use:**

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**Cattle**

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

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**Horse**

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

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**Pig**

- Meat and offal. 0 day

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

QH01CA01

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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:**

Netherlands

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Vetoquinol S.A.

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**Marketing authorisation date:**

6/01/2014

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**Manufacturing sites for batch release:**

Wirtschaftsgenossenschaft deutscher Tieraerzte e G

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 112668

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**Date of authorisation status change:**

8/02/2022

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0154/001

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**Concerned member states:**

Belgium Czechia France Hungary Ireland Netherlands

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

Combined File of all Documents

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