

# Ovogest 1000 I.E./ml Pulver und Lösungsmittel zur Herstellung einer Injektionslösung für Tiere

Authorised

- Serum gonadotrophin

## Product identification

### **Medicine name:**

Ovogest 1000 I.E./ml Pulver und Lösungsmittel zur Herstellung einer Injektionslösung für Tiere

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### **Active substance:**

Serum gonadotrophin

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### **Target species:**

Cattle

Horse

Pig

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### **Route of administration:**

Intravenous use

Other use

Intramuscular use

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## Product details

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

Serum gonadotrophin

5000.00 international unit(s) / 5.00 millilitre(s)

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### **Pharmaceutical form:**

Powder and solvent for solution for injection

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### **Withdrawal period by route of administration:**

#### **Intravenous use:**

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#### **Cattle**

- Milk. 0 day

- Meat and offal. 0 day

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#### **Horse**

- Milk. 0 day

- Meat and offal. 0 day

- 

#### **Pig**

- Meat and offal. 0 day

#### **Other use:**

- 

#### **Cattle**

- Milk. 0 day

- Meat and offal. 0 day

- 

#### **Horse**

- Milk. 0 day

- Meat and offal. 0 day

- 

**Pig**

- Meat and offal. 0 day

**Intramuscular use:**

- 

**Cattle**

- Milk. 0 day

- Meat and offal. 0 day

- 

**Horse**

- Milk. 0 day

- Meat and offal. 0 day

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**Pig**

- Meat and offal. 0 day

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

QG03GA01

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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:**

Germany

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**Package description:**

Available only in German

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**Additional information**

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Intervet Deutschland GmbH

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**Marketing authorisation date:**

17/11/2005

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**Manufacturing sites for batch release:**

Intervet International B.V.

Intervet International GmbH

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

6021798.00.00

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**Date of authorisation status change:**

17/11/2005

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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

Combined File of all Documents

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