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# VETECARDIOL

Not authorised

- HEPTAMINOL ACEFYLLINE

## Product identification

**Medicine name:**

VETECARDIOL

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**Active substance:**

HEPTAMINOL ACEFYLLINE

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**Target species:**

Cattle

Pig

Cat

Horse

Horse (mare)

Sheep

Goat

Dog

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**Route of administration:**

Intramuscular use

Subcutaneous use

Intravenous use

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## Product details

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

HEPTAMINOL ACEFYLLINE

100.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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

#### **Intramuscular use:**

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#### **Cattle**

- Milk. 2 day
- Meat and offal. 2 day

- 

#### **Pig**

- Meat and offal. 2 day

- 

#### **Horse**

- Meat and offal. 2 day

- 

#### **Horse (mare)**

- Milk. 2 day

- 

#### **Sheep**

- Meat and offal. 2 day
- Milk. 2 day

- 

#### **Goat**

- Milk. 2 day

- Meat and offal. 2 day

### **Subcutaneous use:**

•

#### **Cattle**

- Meat and offal. 2 day

- Milk. 2 day

•

#### **Pig**

- Meat and offal. 2 day

•

#### **Horse**

- Meat and offal. 2 day

•

#### **Horse (mare)**

- Milk. 2 day

•

#### **Sheep**

- Meat and offal. 2 day

- Milk. 2 day

•

#### **Goat**

- Meat and offal. 2 day

- Milk. 2 day

### **Intravenous use:**

•

#### **Cattle**

- Milk. 2 day

- Meat and offal. 2 day

•

**Pig**

- Meat and offal. 2 day

•

**Horse**

- Meat and offal. 2 day

•

**Horse (mare)**

- Milk. 2 day

•

**Sheep**

- Meat and offal. 2 day

- Milk. 2 day

•

**Goat**

- Meat and offal. 2 day

- Milk. 2 day

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

QC01DX08

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

France

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Intervet

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**Marketing authorisation date:**

4/01/1980

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**Manufacturing sites for batch release:**

Trirx Segre

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/8407177 0/1980

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**Date of authorisation status change:**

20/03/2025

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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

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