

# WERAVET Vomisal C30 - Injektionslösung für Tiere

Authorised

- PSYCHOTRIA IPECACUANHA C30

## Product identification

**Medicine name:**

WERAVET Vomisal C30 - Injektionslösung für Tiere

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**Active substance:**

PSYCHOTRIA IPECACUANHA C30

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**Target species:**

Cattle

Dog

Goat

Sheep

Horse

Cat

Pig

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

### Active substance and strength:

PSYCHOTRIA IPECACUANHA C30

199.10 milligram(s) / 2.00 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

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

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##### Goat

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

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##### Sheep

- 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

#### Intravenous use:

- 

### **Cattle**

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

- 

### **Goat**

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

- 

### **Sheep**

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

- 

### **Horse**

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

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### **Pig**

- Meat and offal. 0 day

## **Subcutaneous use:**

- 

### **Cattle**

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

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### **Goat**

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

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**Sheep**

- 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:**

QV03AX

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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:**

Austria

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**Package description:**

Available only in German

Available only in German

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Homeopathic medicinal products (Article 19 of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dr. Assmann Veterinaerspezialitaeten GmbH

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**Marketing authorisation date:**

1/09/1998

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**Manufacturing sites for batch release:**

Biokanol Pharma GmbH

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

8-30056

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**Date of authorisation status change:**

1/09/1998

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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.

Labelling

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.