

WERAVET Vomisal C30 - Injektionslösung für Tiere

Authorised

- PSYCHOTRIA IPECACUANHA C30

Product identification

Medicine name:

WERAVET Vomisal C30 - Injektionslösung für Tiere

Active substance:

PSYCHOTRIA IPECACUANHA C30

Target species:

Cattle

Dog

Goat

Sheep

Horse

Cat

Pig

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

PSYCHOTRIA IPECACUANHA C30
199.10 milligram(s) / 2.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular 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

•

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

•

Pig

- Meat and offal. 0 day

Subcutaneous 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

•

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QV03AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Package description:

Available only in German

Available only in German

Additional information**Entitlement type:**

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Dr. Assmann Veterinaerspezialitaeten GmbH

Marketing authorisation date:

1/09/1998

Manufacturing sites for batch release:

Biokanol Pharma GmbH

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-30056

Date of authorisation status change:

1/09/1998

To consult adverse reactions on veterinary medicinal products please go to

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.