

Urtica/Stannum comp. "Wala" - Injektionslösung für Tiere

Authorised

- URTICA URENS EX HERBA FERM 33C DIL. D2 (HAB, VS. 33C)
- STANNUM METALLICUM DIL. D9
- CALCIUM CARBONICUM HAHNEMANNI DIL D6

Product identification

Medicine name:

Urtica/Stannum comp. "Wala" - Injektionslösung für Tiere

Active substance:

URTICA URENS EX HERBA FERM 33C DIL. D2 (HAB, VS. 33C)

STANNUM METALLICUM DIL. D9

CALCIUM CARBONICUM HAHNEMANNI DIL D6

Target species:

Cattle

Dog

Goat

Sheep

Horse

Cat

Rabbit

Rodents

Pig

Route of administration:

Intramuscular use
Subcutaneous use

Product details**Active substance and strength:**

URTICA URENS EX HERBA FERM 33C DIL. D2 (HAB, VS. 33C)

100.00 milligram(s) / 1.00 millilitre(s)

STANNUM METALLICUM DIL. D9

100.00 milligram(s) / 1.00 millilitre(s)

CALCIUM CARBONICUM HAHNEMANNI DIL D6

100.00 milligram(s) / 1.00 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

•

Rabbit

- Meat and offal. 0 day

•

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

•

Rabbit

- Meat and offal. 0 day

-

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:

SaluVet GmbH

Marketing authorisation date:

12/08/1997

Manufacturing sites for batch release:

Wala-Heilmittel GmbH

Responsible authority:

Authorisation number:

8-30018

Date of authorisation status change:

12/08/1997

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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

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.