

Pulmo/Stibium comp. "Wala" - Injektionslösung für Tiere

Authorised

- TARTARUS STIBIATUS DIL. D7 AQUOS. (HAB, VS. 5B)
- STANNUM METALLICUM DIL. D9
- PULMO BOVIS GL DIL. D5 (HAB, VS. 41A)

Product identification

Medicine name:

Pulmo/Stibium comp. "Wala" - Injektionslösung für Tiere

Active substance:

TARTARUS STIBIATUS DIL. D7 AQUOS. (HAB, VS. 5B)

STANNUM METALLICUM DIL. D9

PULMO BOVIS GL DIL. D5 (HAB, VS. 41A)

Target species:

Cattle

Dog

Goat

Sheep

Horse

Cat

Rabbit

Guinea pig

Hamster

Rat

Pig

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

TARTARUS STIBIATUS DIL. D7 AQUOS. (HAB, VS. 5B)

100.00 milligram(s) / 1.00 millilitre(s)

STANNUM METALLICUM DIL. D9

100.00 milligram(s) / 1.00 millilitre(s)

PULMO BOVIS GL DIL. D5 (HAB, VS. 41A)

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

•

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

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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

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:

31/03/1998

Manufacturing sites for batch release:

Wala-Heilmittel GmbH

Responsible authority:

Authorisation number:

8-30030

Date of authorisation status change:

31/03/1998

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.