Cartilago comp. "Wala" - Injektionslösung für Tiere

Authorised

- ECHINACEA PALLIDA E PLANTA TOTA FERM 33C DIL. D2 (HAB, VS. 33C)
- ARTICULATIO INTERPHALANGEA BOVIS GL DIL. D16 (HAB, VS. 41B)
- QUARZ DIL. D29 AQUOS.

Product identification

Medicine name:
Cartilago comp. "Wala" - Injektionslösung für Tiere

Active substance:
ECHINACEA PALLIDA E PLANTA TOTA FERM 33C DIL. D2 (HAB, VS. 33C)
ARTICULATIO INTERPHALANGEA BOVIS GL DIL. D16 (HAB, VS. 41B)
QUARZ DIL. D29 AQUOS.

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:
ECHINACEA PALLIDA E PLANTA TOTA FERM 33C DIL. D2 (HAB, VS. 33C)
100.00 milligram(s) / 1.00 millilitre(s)

ARTICULATIO INTERPHALANGEA BOVIS GL DIL. D16 (HAB, VS. 41B)
100.00 milligram(s) / 1.00 millilitre(s)

QUARZ DIL. D29 AQUOS.
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 hour
- Milk. 0 day

•

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:
9/01/1997

Manufacturing sites for batch release:

Wala-Heilmittel GmbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30014

Date of authorisation status change:
9/01/1997

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.