

Cardio ReVet RV 4 - Injektionslösung für Tiere

Authorised

- APIS MELLIFICA C12
- Apocynum cannabinum C12
- Digitalis purpurea C12
- Prunus laurocerasus C12
- Strychnos nux-vomica C12
- Urginea maritima C12

Product identification

Medicine name:

Cardio ReVet RV 4 - Injektionslösung für Tiere

Active substance:

APIS MELLIFICA C12

Apocynum cannabinum C12

Digitalis purpurea C12

Prunus laurocerasus C12

Strychnos nux-vomica C12

Urginea maritima C12

Target species:

Reptile

Ornamental bird

Dog

Cat

Rabbit
Ferret
Small rodents

Route of administration:

Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

APIS MELLIFICA C12

1.67 gram(s) / 10.00 millilitre(s)

Apocynum cannabinum C12

1.67 gram(s) / 10.00 millilitre(s)

Digitalis purpurea C12

1.67 gram(s) / 10.00 millilitre(s)

Prunus laurocerasus C12

1.67 gram(s) / 10.00 millilitre(s)

Strychnos nux-vomica C12

1.67 gram(s) / 10.00 millilitre(s)

Urginea maritima C12

1.67 gram(s) / 10.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Rabbit

- Meat and offal. 0 day

Intravenous use:

-

Rabbit

- Meat and offal. 0 day

Subcutaneous use:

-

Rabbit

- 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:

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Marketing authorisation date:

21/07/2010

Manufacturing sites for batch release:

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-30101

Date of authorisation status change:

21/07/2010

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.

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