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Cardio ReVet RV 4 - Globuli für Tiere

Authorised

- APIS MELLIFICA C4
- Apocynum cannabinum C1
- Digitalis purpurea C6
- Prunus laurocerasus C3
- Strychnos nux-vomica C6
- Urginea maritima C1

Product identification

Medicine name:
Cardio ReVet RV 4 - Globuli für Tiere

Active substance:
APIS MELLIFICA C4
Apocynum cannabinum C1
Digitalis purpurea C6
Prunus laurocerasus C3
Strychnos nux-vomica C6
Urginea maritima C1

Target species:
Cattle
Reptile
Ornamental bird
Dog

Goat
Sheep
Horse
Cat
Rabbit
Ferret
Small rodents
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
APIS MELLIFICA C4
1.67 milligram(s) / 1.00 gram(s)

Apocynum cannabinum C1
1.67 milligram(s) / 1.00 gram(s)

Digitalis purpurea C6
1.67 milligram(s) / 1.00 gram(s)

Prunus laurocerasus C3
1.67 milligram(s) / 1.00 gram(s)

Strychnos nux-vomica C6
1.67 milligram(s) / 1.00 gram(s)

Urginea maritima C1
1.67 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Pillules

Withdrawal period by route of administration:
Oral 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:
Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Marketing authorisation date:
7/04/2010

Manufacturing sites for batch release:
Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30100

Date of authorisation status change:
7/04/2010

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

Documents

Package Leaflet

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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.

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