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Epispas ReVet RV10 - Globuli für Tiere

Authorised

- Citrullus colocynthis C6
- CUPRUM METALLICUM C9
- Hyoscyamus niger C6
- LACHESIS C9

Product identification

Medicine name:
Epispas ReVet RV10 - Globuli für Tiere

Active substance:
Citrullus colocynthis C6

CUPRUM METALLICUM C9

Hyoscyamus niger C6

LACHESIS C9

Target species:
Pigeon
Cattle
Reptile
Fowl
Ornamental bird
Cattle (calf)
Dog
Goat

Sheep
Horse
Cat
Rabbit
Ferret
Small rodents
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
Citrullus colocynthis C6
2.50 milligram(s) / 1.00 gram(s)

CUPRUM METALLICUM C9
2.50 milligram(s) / 1.00 gram(s)

Hyoscyamus niger C6
2.50 milligram(s) / 1.00 gram(s)

LACHESIS C9
2.50 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Pillules

Withdrawal period by route of administration:
Oral use:

- **Pigeon**
  - Meat and offal. 0 day

- **Cattle**
  - Meat and offal. 0 day

- Milk. 0 hour

- **Fowl**
  - Eggs. 0 day
  - Meat and offal. 0 day

- **Cattle (calf)**
  - Meat and offal. 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 not 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:
7/07/1998

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30048

Date of authorisation status change:
7/07/1998

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.