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Hormon ReVet RV13 - Globuli für Tiere

Authorised

- Glandula suprarenalis suis C9
- Glandula thymi suis C9
- Hypophysis suis C9
- Ovarium suis C9
- Pancreas suis C9
- Testis suis C9
- Glandula thyreoidea suis c9

Product identification

Medicine name:
Hormon ReVet RV13 - Globuli für Tiere

Active substance:
Glandula suprarenalis suis C9

Glandula thymi suis C9

Hypophysis suis C9

Ovarium suis C9

Pancreas suis C9

Testis suis C9

Glandula thyreoidea suis 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:
Glandula suprarenalis suis C9
1.43 milligram(s) / 1.00 gram(s)

Glandula thymi suis C9
1.43 milligram(s) / 1.00 gram(s)

Hypophysis suis C9
1.43 milligram(s) / 1.00 gram(s)

Ovarium suis C9
1.43 milligram(s) / 1.00 gram(s)

Pancreas suis C9
1.43 milligram(s) / 1.00 gram(s)

Testis suis C9
1.43 milligram(s) / 1.00 gram(s)

Glandula thyreoidea suis c9
1.43 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:
8-30050

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

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Labelling

This document does not exist in this language (English). You can find it in another language below.