Nympho ReVet RV20 - Globuli für Tiere

Authorised

- AURUM METALLICUM C9
- BUFO RANA C9
- Origanum majorana C6
- Platinum metallicum C9

Product identification

Medicine name:
Nympho ReVet RV20 - Globuli für Tiere

Active substance:
AURUM METALLICUM C9

BUFO RANA C9

Origanum majorana C6

Platinum metallicum 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:
AURUM METALLICUM C9
2.50 milligram(s) / 1.00 gram(s)

BUFO RANA C9
2.50 milligram(s) / 1.00 gram(s)

Origanum majorana C6
2.50 milligram(s) / 1.00 gram(s)

Platinum metallicum 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:
25/05/1998

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30033

Date of authorisation status change:
25/05/1998

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

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