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Broncho-Akut ReVet RV 3A - Globuli für Tiere

Authorised

- Aconitum napellus C9
- FERRUM PHOSPHORICUM C9
- HEPAR SULFURIS C9

Product identification

Medicine name:
Broncho-Akut ReVet RV 3A - Globuli für Tiere

Active substance:
Aconitum napellus C9

FERRUM PHOSPHORICUM C9

HEPAR SULFURIS C9

Target species:
Pigeon
Turkey (hen)
Cattle
Reptile
Chicken
Duck
Goose
Ornamental bird
Dog
Goat

Sheep
Horse
Cat
Rabbit
Ferret
Small rodents
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
Aconitum napellus C9
3.33 milligram(s) / 1.00 gram(s)

FERRUM PHOSPHORICUM C9
3.33 milligram(s) / 1.00 gram(s)

HEPAR SULFURIS C9
3.33 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Pillules

Withdrawal period by route of administration:
Oral use:

- **Pigeon**
  - Meat and offal. 0 day
- **Turkey (hen)**
  - Eggs. 0 day
  - Meat and offal. 0 day
-

Cattle

- Meat and offal. 0 day
- Milk. 0 hour

•

Chicken

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

•

Duck

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

•

Goose

- Eggs. 0 day
- 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:
6/04/2010

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30097

Date of authorisation status change:
6/04/2010

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

Documents

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.

Package Leaflet

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

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