# 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

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**Reptile**

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**Chicken**

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

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**Duck**

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

•

**Goose**

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

•

**Ornamental bird**

•

**Dog**

•

**Goat**

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

•

**Sheep**

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

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**Horse**

- Meat and offal. 0 day

- Milk. 0 hour

- **Cat**
- **Rabbit**
  - Meat and offal. 0 day
- **Ferret**
- **Small rodents**
- **Pig**
  - Meat and offal. 0 day

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**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

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.

Package Leaflet

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

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