

# Miktiolith ReVet RV 18 - Globuli für Tiere

Authorised

- Berberis vulgaris C4
- LYTTA VESICATORIA C6
- LYCOPODIUM CLAVATUM C9
- Sabal serrulata C4

## Product identification

**Medicine name:**

Miktiolith ReVet RV 18 - Globuli für Tiere

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**Active substance:**

Berberis vulgaris C4

LYTTA VESICATORIA C6

LYCOPODIUM CLAVATUM C9

Sabal serrulata C4

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**Target species:**

Cattle

Cattle (calf)

Dog

Goat

Sheep

Horse

Cat

Rabbit

Ferret  
Small rodents  
Pig

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Berberis vulgaris C4  
2.50 milligram(s) / 1.00 gram(s)

LYTTA VESICATORIA C6  
2.50 milligram(s) / 1.00 gram(s)

LYCOPODIUM CLAVATUM C9  
2.50 milligram(s) / 1.00 gram(s)

Sabal serrulata C4  
2.50 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Pillules

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**Withdrawal period by route of administration:**

**Oral use:**

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**Cattle**

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

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**Cattle (calf)**

- Meat and offal. 0 day

•

**Goat**

- Meat and offal. 0 day

- Milk. 0 hour

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**Sheep**

- Meat and offal. 0 day

- Milk. 0 hour

•

**Horse**

- Meat and offal. 0 day

- Milk. 0 hour

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**Rabbit**

- Meat and offal. 0 day

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**Pig**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QV03AX

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria

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**Package description:**

Available only in German

Available only in German

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Homeopathic medicinal products (Article 19 of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

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**Marketing authorisation date:**

23/04/1998

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**Manufacturing sites for batch release:**

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

8-30032

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**Date of authorisation status change:**

23/04/1998

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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

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