

ReVet RV 25 Streukügelchen für Pferd, Rind, Schaf, Ziege, Schwein, Huhn, Pute, Gans, Ente, Taube, Kaninchen, Karpfen, Forelle, Hund, Katze, Maus, Ratte, Meerschweinchen, Marderartige, Ziervögel, Amphibien und Reptilien

Authorised

- ARNICA MONTANA EX PLANTA TOTA D6
- Hypericum perforatum C6
- Ledum palustre C6
- Ruta graveolens C6
- Strychnos nux-vomica C6
- Symphytum officinale C6
- Rhus toxicodendron C6

Product identification

Medicine name:

ReVet RV 25 Streukügelchen für Pferd, Rind, Schaf, Ziege, Schwein, Huhn, Pute, Gans, Ente, Taube, Kaninchen, Karpfen, Forelle, Hund, Katze, Maus, Ratte, Meerschweinchen, Marderartige, Ziervögel, Amphibien und Reptilien

Active substance:

ARNICA MONTANA EX PLANTA TOTA D6

Hypericum perforatum C6

Ledum palustre C6

Ruta graveolens C6

Strychnos nux-vomica C6

Symphytum officinale C6

Rhus toxicodendron C6

Target species:

Pigeon

Turkey

Cattle

Amphibians

Reptile

Chicken

Duck

Goose

Ornamental bird

Dog

Goat

Sheep

Horse

Cat

Rabbit

Marten

Guinea pig

Mouse

Rat

Pig

Carp

Trout

Route of administration:

Oral use

Product details

Active substance and strength:

ARNICA MONTANA EX PLANTA TOTA D6

1.43 milligram(s) / 1.00 gram(s)

Hypericum perforatum C6

1.43 milligram(s) / 1.00 gram(s)

Ledum palustre C6

1.43 milligram(s) / 1.00 gram(s)

Ruta graveolens C6

1.43 milligram(s) / 1.00 gram(s)

Strychnos nux-vomica C6

1.43 milligram(s) / 1.00 gram(s)

Symphytum officinale C6

1.43 milligram(s) / 1.00 gram(s)

Rhus toxicodendron C6

1.43 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Pillules

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Package description:

Available only in German

Available only in German

Available only in German

Additional information

Entitlement type:

Homeopathic Registration

Marketing authorisation holder:

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Marketing authorisation date:

20/02/2023

Manufacturing sites for batch release:

Pharmazeutische Fabrik Dr. Reckeweg & Co. GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

V7006787.00.00

Date of authorisation status change:

20/02/2023

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

Documents

Package Leaflet

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