

RemaTuss - Hustenglobuli für Tiere

Authorised

- AMMONIUM CARBONICUM C200
- BRYONIA C200
- DACTYLOPIUS COCCUS C200
- DROSER A C200
- RUMEX CRISPUS C200
- EUSPONGIA OFFICINALIS C200
- STANNUM METALLICUM C200

Product identification

Medicine name:

RemaTuss - Hustenglobuli für Tiere

Active substance:

AMMONIUM CARBONICUM C200

BRYONIA C200

DACTYLOPIUS COCCUS C200

DROSER A C200

RUMEX CRISPUS C200

EUSPONGIA OFFICINALIS C200

STANNUM METALLICUM C200

Target species:

Cattle

Dog
Goat
Sheep
Horse
Cat
Rabbit
Ferret
Small rodents
Pig

Route of administration:

Oral use

Product details

Active substance and strength:

AMMONIUM CARBONICUM C200

1.43 milligram(s) / 1.00 gram(s)

BRYONIA C200

1.43 milligram(s) / 1.00 gram(s)

DACTYLOPIUS COCCUS C200

1.43 milligram(s) / 1.00 gram(s)

DROSER A C200

1.43 milligram(s) / 1.00 gram(s)

RUMEX CRISPUS C200

1.43 milligram(s) / 1.00 gram(s)

EUSPONGIA OFFICINALIS C200

1.43 milligram(s) / 1.00 gram(s)

STANNUM METALLICUM C200

1.43 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Pillules

Withdrawal period by route of administration:

Oral use:

-

Cattle

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

-

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:

Remedia Homoeopathie Mag. Pharm. Robert Muentz Ges.m.b.H.

Marketing authorisation date:

13/06/2021

Manufacturing sites for batch release:

Remedia Homoeopathie Mag. Pharm. Robert Muentz Ges.m.b.H.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

840705

Date of authorisation status change:

13/06/2021

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.

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.

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