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Stomato ReVet RV24 - Injektionslösung für Tiere

Authorised

- APIS MELLIFICA C4
- Natrium tetraboracicum C4
- Kreosotum C6

Product identification

Medicine name:
Stomato ReVet RV24 - Injektionslösung für Tiere

Active substance:
APIS MELLIFICA C4

Natrium tetraboracicum C4

Kreosotum C6

Target species:
Cattle
Goat
Sheep
Horse
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

APIS MELLIFICA C4
0.33 gram(s) / 1.00 millilitre(s)

Natrium tetraboracicum C4
0.33 gram(s) / 1.00 millilitre(s)

Kreosotum C6
0.33 gram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular 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

- **Pig**
  - Meat and offal. 0 day

Intravenous 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
- **Pig**
  - Meat and offal. 0 day

Subcutaneous 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

•

Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV03AX

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Austria

Package description:
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:
6/07/1998

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30039

Date of authorisation status change:
6/07/1998

To consult adverse reactions on veterinary medicinal products please go to 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

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