Traumato ReVet RV 25 - Injektionslösung für Tiere

Authorised

- Arnica montana ex planta tota C6
- Hypericum perforatum C6
- Ledum palustre C6
- Strychnos nux-vomica C6
- Rhus toxicodendron C6
- Ruta graveolens C6
- Symphytum officinale C6

Product identification

Medicine name:
Traumato ReVet RV 25 - Injektionslösung für Tiere

Active substance:
Arnica montana ex planta tota C6

Hypericum perforatum C6

Ledum palustre C6

Strychnos nux-vomica C6

Rhus toxicodendron C6

Ruta graveolens C6

Symphytum officinale C6

Target species:
Cattle

Dog
Goat
Sheep
Horse
Cat
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Arnica montana ex planta tota C6
143.00 milligram(s) / 1.00 millilitre(s)

Hypericum perforatum C6
143.00 milligram(s) / 1.00 millilitre(s)

Ledum palustre C6
143.00 milligram(s) / 1.00 millilitre(s)

Strychnos nux-vomica C6
143.00 milligram(s) / 1.00 millilitre(s)

Rhus toxicodendron C6
143.00 milligram(s) / 1.00 millilitre(s)

Ruta graveolens C6
143.00 milligram(s) / 1.00 millilitre(s)

Symphytum officinale C6
143.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**

- Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Horse**
  - 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 subject to veterinary prescription

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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:
9/02/2011

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-30107

Date of authorisation status change:
9/02/2011

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.

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