Receptal 0,004 mg/ml Injektionslösung für Rinder, Pferde, Schweine und Kaninchen

Authorised

This information is not available for this product.

Product identification

Medicine name:
Receptal 0,004 mg/ml Injektionslösung für Rinder, Pferde, Schweine und Kaninchen

Active substance:
This information is not available for this product.

Target species:
Cattle
Horse
Rabbit
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - 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

Intravenous use:

- **Cattle**
  - 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

Subcutaneous use:

- **Cattle**
  - 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:
QH01CA90

Legal status of supply:
Medicinal product on medical prescription for renewable delivery

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Available only in German
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Ges.m.b.H.

Marketing authorisation date:
23/03/1981

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
16887

Date of authorisation status change:
11/05/2010

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

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