Analeptol 50 mg/ml + 50 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

Authorised

- Heptaminol
- Diprophylline

Product identification

Medicine name:
Analeptol 50 mg/ml + 50 mg/ml solution for injection for cattle, horses, pigs, dogs and cats
Analeptol 50 mg/ml + 50 mg/ml Injektionslösung für Rinder, Pferde, Schweine, Hunde und Katzen

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Active substance:
Heptaminol
Diprophylline

Target species:
Cattle
Horse
Pig
Cattle (calf)
Horse (foal)
Pig (piglet)
Dog
Cat

Route of administration:
Intravenous use
Intraperitoneal use
Intramuscular use

Product details

Active substance and strength:
Heptaminol
50.00 milligram(s) / 1.00 millilitre(s)

Diprophylline
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Meat and offal. 2 day
  - Milk. 48 hour
- **Horse**
  - Meat and offal. 2 day
  - Milk. 48 hour
- **Pig**
  - Meat and offal. 2 day
- **Cattle (calf)**
  - Meat and offal. 2 day
- **Horse (foal)**
  - Meat and offal. 2 day
- **Pig (piglet)**
  - Meat and offal. 2 day
- **Dog**

- **Cat**

Intraperitoneal use:

- **Cattle**
  - Meat and offal. 2 day
  - Milk. 48 hour
- **Horse**
  - Meat and offal. 2 day
  - Milk. 48 hour
- **Pig**
  - Meat and offal. 2 day
- **Cattle (calf)**
  - Meat and offal. 2 day
- **Horse (foal)**
  - Meat and offal. 2 day
- **Pig (piglet)**
  - Meat and offal. 2 day
- **Dog**
- **Cat**

Intramuscular use:

- **Cattle (calf)**
  - Meat and offal. 7 day
- **Horse (foal)**
  - Meat and offal. 7 day
- **Pig (piglet)**
  - Meat and offal. 7 day
- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR03DA51

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

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Authorisation status:
Valid

Authorised in:
Austria

Available in:
Austria

Package description:
Carton box containing 1 clear type I glass vial containing 50 ml closed with a grey bromobutyl rubber stopper and aluminium cap.
Carton box containing 1 clear type I glass vial containing 20 ml closed with a grey bromobutyl rubber stopper and aluminium cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
29/03/2023

Manufacturing sites for batch release:
Cp-Pharma Handelsgesellschaft mbH

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
841644

Date of authorisation status change:

29/03/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0379/001

Concerned member states:
Austria Belgium Estonia France Germany Hungary Ireland Italy Portugal Spain

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

Documents

Combined File of all Documents

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.

Package Leaflet

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