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

Authorised

- Diprophylline
- Heptaminol

Product identification

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

Active substance:
Diprophylline
Heptaminol

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

Heptaminol
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

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

Intramuscular use:

•

Cattle (calf)

- Meat and offal. 7 day

- **Horse (foal)**
  - Meat and offal. 7 day

- **Pig (piglet)**
  - Meat and offal. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR03DA51

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

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:
10/03/2023

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 129522

Date of authorisation status change:
1/08/2023

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

English (PDF)

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