

Lactato-RingerVet Solution for infusion for cattle, horses, sheep, goats, pigs, dogs and cats

Authorised

- Potassium chloride
- Sodium chloride
- Calcium chloride dihydrate
- Sodium lactate

Product identification

Medicine name:

Lactato-RingerVet Solution for infusion for cattle, horses, sheep, goats, pigs, dogs and cats

Ringer-Lactat-Lösung nach Hartmann B. Braun Vet Care, Infusionslösung für Rinder, Pferde, Schafe, Ziegen, Schweine, Hunde und Katzen

Active substance:

Potassium chloride

Sodium chloride

Calcium chloride dihydrate

Sodium lactate

Target species:

Cattle

Sheep

Goat

Horse

Pig

Dog

Cat

Route of administration:

Intravenous use

Product details

Active substance and strength:

Potassium chloride

0.04 gram(s) / 100.00 millilitre(s)

Sodium chloride

0.60 gram(s) / 100.00 millilitre(s)

Calcium chloride dihydrate

0.03 gram(s) / 100.00 millilitre(s)

Sodium lactate

0.31 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:

Solution for infusion

Withdrawal period by route of administration:

Intravenous use:

• **Cattle**

- Meat and offal. 0 day

- Milk. 0 day

• **Sheep**

- Meat and offal. 0 day

- Milk. 0 day

• **Goat**

- Meat and offal. 0 day

- Milk. 0 day

• **Horse**

- Meat and offal. 0 day

- Milk. 0 day

- **Pig**

- Meat and offal. 0 day

- Milk. 0 day

- **Dog**

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

Package description:

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:

B. Braun Melsungen AG

Marketing authorisation date:

19/05/2010

Manufacturing sites for batch release:

B.Braun Medical S.A.

B. Braun Melsungen AG

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-00872

Date of authorisation status change:

19/05/2010

Reference member state:

Spain

Procedure number:

ES/V/0153/001

Concerned member states:

Austria Belgium Germany Ireland Italy Netherlands

United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

English (PDF)

Published on: 22/12/2023

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Summary of Product Characteristics

English (PDF)

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Labelling

English (PDF)

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