

B. Braun Vet Care Hartmann's Lactated Ringers Solution for infusion for cattle, horse, sheep, goat, pig, dog and cat

Authorised

- Potassium chloride
- Sodium chloride
- Calcium chloride dihydrate
- Sodium (S)-lactate solution

Product identification

Medicine name:

B. Braun Vet Care Hartmann's Lactated Ringers Solution for infusion for cattle, horse, sheep, goat, pig, dog and cat

Active substance:

Potassium chloride

Sodium chloride

Calcium chloride dihydrate

Sodium (S)-lactate solution

Target species:

Cattle

Sheep

Goat

Horse

Pig

Dog

Cat

Route of administration:

Intravenous use

Product details

Active substance and strength:

Potassium chloride

0.40 milligram(s) / 1.00 millilitre(s)

Sodium chloride

6.00 milligram(s) / 1.00 millilitre(s)

Calcium chloride dihydrate

0.27 milligram(s) / 1.00 millilitre(s)

Sodium (S)-lactate solution

6.24 milligram(s) / 1.00 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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

box containing 10 bottles of 1000 ml

box containing 10 bottles of 500 ml

box containing 2 bags of 5000 ml

box containing 20 bottles of 250 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

B. Braun Melsungen AG

Marketing authorisation date:

27/04/2010

Manufacturing sites for batch release:

B BRAUN MEDICAL S.A.

B. Braun Melsungen AG

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

30/07/2015

Reference member state:

Spain

Procedure number:

ES/V/0153/001

Concerned member states:

Austria Belgium Denmark France Germany Ireland Italy Netherlands Poland
Portugal Romania Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 22/10/2025

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Package Leaflet

English (PDF)

Published on: 22/10/2025

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Labelling

English (PDF)

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Combined File of all Documents

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