AQUPHARM RINGER LACTATE SOLUTION FOR INFUSION

Authorised

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

Product identification

Medicine name:
AQUPHARM RINGER LACTATE SOLUTION FOR INFUSION
Aqupharm Ringer-laktát oldatos infúzió A.U.V.

Active substance:
Sodium chloride
Sodium lactate
Calcium chloride
Potassium chloride

Target species:
Cattle
Pig
Rabbit
Cat
Horse
Sheep
Goat

Dog

Route of administration:
Intravenous use

Product details

Active substance and strength:
Sodium chloride
6.00 milligram(s) / 1.00 millilitre(s)

Sodium lactate
3.10 milligram(s) / 1.00 millilitre(s)

Calcium chloride
0.20 milligram(s) / 1.00 millilitre(s)

Potassium chloride
0.40 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
- **Pig**
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
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Horse

- Meat and offal. 0 day
- Milk. 0 day

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Sheep

- Meat and offal. 0 day
- Milk. 0 day

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Goat

- 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

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Authorisation status:
Valid

Authorised in:
Hungary

Available in:
Hungary

Package description:
1 bag of 250 ml
1 bag of 500 ml
1 bag of 1000 ml
1 bag of 3000 ml
1 bag of 5000 ml
Cardboard box containing 30 bags of 250 ml
Cardboard box containing 2 bags of 5000 ml
Cardboard box containing 4 bags of 3000 ml

Cardboard box containing 10 bags of 1000 ml
Cardboard box containing 20 bags of 500 ml

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:
Ecuphar

Marketing authorisation date:
10/11/2016

Manufacturing sites for batch release:
Laboratoire Bioluz

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
3810/X/16 NÉBIH ÁTI

Date of authorisation status change:
10/11/2016

Reference member state:
France

Procedure number:
FR/V/0303/001

Concerned member states:
Belgium Finland Hungary Iceland Ireland Netherlands
United Kingdom (Northern Ireland)

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