LODEVIL

Authorised

- Sodium acetate
- Sodium hydrogen carbonate
- Sodium chloride
- Potassium chloride
- Magnesium chloride
- Glucose

Product identification

Medicine name:
LODEVIL

Active substance:
Sodium acetate

Sodium hydrogen carbonate

Sodium chloride

Potassium chloride

Magnesium chloride

Glucose

Target species:
Cattle

Route of administration:
Intravenous use

Product details

Active substance and strength:

Sodium acetate
4.08 gram(s) / 1.00 litre(s)

Sodium hydrogen carbonate
4.20 gram(s) / 1.00 litre(s)

Sodium chloride
3.51 gram(s) / 1.00 litre(s)

Potassium chloride
0.45 gram(s) / 1.00 litre(s)

Magnesium chloride
0.20 gram(s) / 1.00 litre(s)

Glucose
10.80 gram(s) / 1.00 litre(s)

Pharmaceutical form:

Solution for infusion

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB05BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Available in:

Luxembourg

Package description:

Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
22/01/1982

Manufacturing sites for batch release:
Vetoquinol S.A.

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 808/09/07/1472

Date of authorisation status change:
24/11/2011

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

Documents

Summary of Product Characteristics

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