Addimag 240 mg/ml + 126 mg/ml solution for infusion for cattle

Authorised

- Calcium gluconate monohydrate
- Magnesium chloride hexahydrate

Product identification

Medicine name:
Addimag 240 mg/ml + 126 mg/ml solution for infusion for cattle
ADDIMAG FORTE SOLUTION POUR PERFUSION POUR BOVINS

Active substance:
Calcium gluconate monohydrate

Magnesium chloride hexahydrate

Target species:
Cattle

Route of administration:
Solution for infusion

Product details

Active substance and strength:
Calcium gluconate monohydrate
240.00 milligram(s) / 1.00 millilitre(s)

Magnesium chloride hexahydrate
126.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for solution for infusion

Withdrawal period by route of administration:

Solution for infusion:

- **Cattle**
  - Meat and offal. no withdrawal period
  - Milk. no withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AX

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Package description:
750 ml square shaped clear polypropylene (PP) bottle with a bromobutyl rubber stopper and an aluminium screw cap.
500 ml square shaped clear polypropylene (PP) bottle with a bromobutyl rubber stopper and an aluminium screw cap

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
13/01/2022

Manufacturing sites for batch release:
Alfasan Nederland B.V.
Bela-Pharm GmbH & Co. KG

Responsible authority:
National Veterinary Medicines Agency

Authorisation number:
FR/V/0193319 4/2021

Date of authorisation status change:
13/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0352/002/DC

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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

Labelling

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

Package Leaflet

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