Addimag 160 mg/ml + 84 mg/ml solution for infusion for cattle

Authorised

- Magnesium chloride hexahydrate
- Calcium gluconate monohydrate

Product identification

Medicine name:
Addimag 160 mg/ml + 84 mg/ml Solution pour perfusion
Addimag 160 mg/ml + 84 mg/ml Oplossing voor infusie
Addimag 160 mg/ml + 84 mg/ml Infusionslösung
Addimag 160 mg/ml + 84 mg/ml solution for infusion for cattle

Active substance:
Magnesium chloride hexahydrate
Calcium gluconate monohydrate

Target species:
Cattle

Route of administration:
Intravenous use

Product details

Active substance and strength:
Magnesium chloride hexahydrate

84.00 milligram(s) / 1.00 millilitre(s)

Calcium gluconate monohydrate
160.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for infusion

Withdrawal period by route of administration:
Intravenous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12AX

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
500 ml square shaped clear polypropylene (PP) bottle with a bromobutyl rubber stopper and an aluminium screw cap
750 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:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
23/12/2022

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V660951

Date of authorisation status change:
23/12/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0352/001

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)

Generic of:
600000066129

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

Documents

Combined File of all Documents

Package Leaflet

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

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.