

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 for infusion for cattle

Addimag vet 160 mg/ml + 84 mg/ml infusionsvätska, lösning för nötkreatur

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:

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Cattle

- Meat and offal. no withdrawal period zero days
 - Milk. no withdrawal period zero days
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

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:

8/04/2022

Manufacturing sites for batch release:

Alfasan Nederland B.V.

Bela-Pharm GmbH & Co. KG

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

61334

Date of authorisation status change:

8/04/2022

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

Summary of Product Characteristics

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.

Labelling

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