

# 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

---

**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:**

Finland

---

**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:**

Generic application (Article 18 of Regulation (EU) 2019/6)

---

**Marketing authorisation holder:**

Alfasan Nederland B.V.

---

**Marketing authorisation date:**

4/09/2024

---

**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

Bela-Pharm GmbH & Co. KG

---

**Responsible authority:**

Finnish Medicines Agency

---

**Authorisation number:**

38472

---

**Date of authorisation status change:**

4/09/2024

---

**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](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

NLV0352001-002\_Addimag\_PuAR final.pdf