

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

Authorised

- Magnesium chloride hexahydrate
- Calcium gluconate monohydrate

## Product identification

**Medicine name:**

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

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**Active substance:**

Magnesium chloride hexahydrate

Calcium gluconate monohydrate

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**Target species:**

Cattle

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Magnesium chloride hexahydrate

126.00 milligram(s) / 1.00 millilitre(s)

Calcium gluconate monohydrate

240.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for infusion

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**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 hours
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA12AX

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

1/02/2024

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

Bela-Pharm GmbH & Co. KG

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

105584

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**Date of authorisation status change:**

1/02/2024

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0352/002

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

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