

# Acticarp 50 mg/ml, Solution for Injection

Not  
authorised

- Carprofen

## Product identification

**Medicine name:**

Acticarp 50 mg/ml, Solution for Injection

Acticarp 50 mg/ml solução injetável para bovinos

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**Active substance:**

Carprofen

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**Target species:**

Cattle

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**Route of administration:**

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Carprofen

50.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intravenous use:**

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**Cattle**

- Milk. no withdrawal period 0 hours
- Meat and offal. 21 day

**Subcutaneous use:**

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**Cattle**

- Milk. no withdrawal period zero hours
- Meat and offal. 21 day

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

QM01AE91

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Revoked

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**Authorised in:**

Portugal

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**Package description:**

50 ml amber glass (Type I) vial closed with Flurotec (coated chlorobutyl) rubber stopper and aluminium flip off. One vial in a carton box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ecuphar

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**Marketing authorisation date:**

23/02/2012

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**Manufacturing sites for batch release:**

Fundacio Privada Dau

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

443/01/12DFVPT

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**Date of authorisation status change:**

7/11/2024

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0156/001

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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

English (PDF)

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