

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

Active substance:

Carprofen

Target species:

Cattle

Route of administration:

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Carprofen

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intravenous use:****• Cattle**

- Milk. 0 hour
- Meat and offal. 21 day

Subcutaneous use:**• Cattle**

- Milk. 0 hour
- Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Expired

Authorised in:

Portugal

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

European Medicines Agency

Marketing authorisation date:

14/12/2021

Manufacturing sites for batch release:

Laboratori Fundacio Dau

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

443/01/12DFVPT

Date of authorisation status change:

14/12/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0156/001

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000036598>