Acticarp 50 mg/ml, Solution for Injection

Not authorised

- Carprofen

Product identification

Medicine name:
Acticarp 50 mg/ml, Solution for Injection

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

Subcutaneous use:

- **Cattle**
  - Milk. no withdrawal period zero hours
  - 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:
Revoked

Authorised in:
Belgium

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:
Ecuphar

Marketing authorisation date:
14/03/2012

Manufacturing sites for batch release:
Fundacio Privada Dau

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V415694

Date of authorisation status change:
14/04/2022

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

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.