

Acticarp 50 mg/ml, Solution for Injection

Not authorised

- Carprofen

Product identification

Medicine name:

Acticarp 50 mg/ml, Solution for Injection
Acticarp Cattle 50 mg/ml Solution injectable

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:

Surrendered

Authorised in:

Luxembourg

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:

Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:

V/991/09/12/1262

Date of authorisation status change:

28/12/2023

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.