

Acticarp 50 mg/ml Solution for Injection for Cattle

Not authorised

- Carprofen

Product identification

Medicine name:

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

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:

Expired

Authorised in:

United Kingdom (Northern Ireland)

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:

9/05/2012

Manufacturing sites for batch release:

Fundacio Privada Dau

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

VM 32742/4005

Date of authorisation status change:

20/09/2022

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