

Acticarp 50 mg/ml, Solution for Injection

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:

Valid

Authorised in:

Netherlands

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:

Accord Healthcare B.V.

Marketing authorisation date:

18/04/2012

Manufacturing sites for batch release:

Fundacio Privada Dau

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 109311

Date of authorisation status change:

19/01/2022

Reference member state:

Netherlands

Procedure number:

NL/V/0156/001

Concerned member states:

Austria Belgium Italy Latvia Lithuania Luxembourg Poland Portugal
Slovakia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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