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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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