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Carprodolor 50 mg/ml solution for injection for cattle

Authorised

- Carprofen

Product identification

Medicine name:
Carprodolor 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**
  - Meat and offal. 21 day
  - Milk. 0 hour

Subcutaneous use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE91

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

Package description:
50 ml amber glass (Type I) vial capped with chlorbutyl rubber stopper retained byan aluminium crimped seal in a cardboard 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:
Le Vet. Beheer B.V.

Marketing authorisation date:
24/09/2013

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
50330

Date of authorisation status change:
24/09/2013

Reference member state:
Ireland

Procedure number:
IE/V/0620/001

Concerned member states:
Austria Belgium Czechia Denmark Estonia Finland France Hungary Iceland Italy Latvia Lithuania Luxembourg Malta Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024
Updated on: 14/03/2026
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