RIMADYL Cattle 50 mg/ml Solution for Injection

Authorised

- Carprofen

Product identification

Medicine name:
RIMADYL Cattle 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**
  - Meat and offal. 21 day
  - Milk. 0 day

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE91

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

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Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Cardboard box containing one multidose amber glass (Type I) vial of 250 ml capped with a bromobutyl rubber stopper retained by an aluminium crimped seal.
Cardboard box containing one multidose amber glass (Type I) vial of 100 ml capped with a bromobutyl rubber stopper retained by an aluminium crimped seal.
Cardboard box containing one multidose amber glass (Type I) vial of 50 ml capped with a bromobutyl rubber stopper retained by an aluminium crimped seal.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Polska Sp. z o.o.

Marketing authorisation date:
12/08/2008

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1846

Date of authorisation status change:
12/08/2008

Reference member state:
Ireland

Procedure number:
IE/V/0140/001

Concerned member states:
Belgium Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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

Labelling

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

Summary of Product Characteristics

English (PDF)
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