Norocarp 50 mg/ml Solution for Injection for Cattle

Authorised

- Carprofen

Product identification

Medicine name:
Norocarp 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

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:

Belgium

Package description:

Norocarp Injection for Cattle in 12 x 50ml multidose amber glass (Type I) vials, sealed with 20mm bromobutyl bungs and 20mm aluminium seals.
Norocarp Injection for Cattle in 10 x 50ml multidose amber glass (Type I) vials, sealed with 20mm bromobutyl bungs and 20mm aluminium seals.
Norocarp Injection for Cattle in 6 x 50ml multidose amber glass (Type I) vials, sealed with 20mm bromobutyl bungs and 20mm aluminium seals.
Norocarp Injection for Cattle in 5 x 50ml multidose amber glass (Type I) vials, sealed with 20mm bromobutyl bungs and 20mm aluminium seals.
Norocarp Injection for Cattle in 1 x 50ml multidose amber glass (Type I) vial, sealed with 20mm bromobutyl bung and 20mm aluminium seal.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
2/09/2009

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V346674

Date of authorisation status change:
2/09/2009

Reference member state:
Ireland

Procedure number:
IE/V/0230/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Italy Norway Portugal Spain 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: 29/12/2024
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Package Leaflet

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Combined File of all Documents

Labelling

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