

# Norocarp 50 mg/ml Solution for Injection for Cattle

Authorised

- Carprofen

## Product identification

**Medicine name:**

Norocarp 50 mg/ml Solution for Injection for Cattle

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**Active substance:**

Carprofen

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**Target species:**

Cattle

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**Route of administration:**

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Carprofen

50.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intravenous use:**

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**Cattle**

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

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 21 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AE91

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**

Valid

**Authorised in:**

Ireland

**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Norbrook Laboratories (Ireland) Limited

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**Marketing authorisation date:**

9/09/2005

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**Manufacturing sites for batch release:**

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA22664/073/001

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**Date of authorisation status change:**

9/09/2005

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0230/001

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**Concerned member states:**

Austria Belgium Denmark Finland France Germany Italy Norway Portugal  
Spain United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Combined File of all Documents