

# DINALGEN 150 mg/ml solution for injection for cattle, pigs and horses

Authorised

- Ketoprofen

## Product identification

**Medicine name:**

DINALGEN 150 mg/ml solution for injection for cattle, pigs and horses

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**Active substance:**

Ketoprofen

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**Target species:**

Cattle

Pig

Horse

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**Route of administration:**

Intramuscular use

Intravenous use

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## Product details

**Active substance and strength:**

Ketoprofen

150.00 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:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. 2 day

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**Pig**

- Meat and offal. 3 day

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**Cattle**

- Milk. 0 hour

**Intravenous use:**

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**Cattle**

- Meat and offal. 2 day

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**Horse**

- Meat and offal. 1 day

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**Cattle**

- Milk. 0 hour

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**Horse**

- Milk. no withdrawal period

Milk: Not permitted for use in animals producing milk for human consumption.

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

QM01AE03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**Package description:**

box containing 5 vials of 250 ml  
box containing 10 vials of 100 ml  
box containing 5 vials of 100 ml  
box containing 1 vial of 100 ml  
box containing 1 vial of 250 ml

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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:**

Ecuphar Veterinaria S.L.U.

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**Marketing authorisation date:**

16/11/2010

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**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2022

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**Date of authorisation status change:**

16/11/2010

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0115/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Estonia Finland France Germany  
Hungary Ireland Italy Latvia Lithuania Netherlands Poland Portugal  
Romania Slovakia Slovenia Sweden 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

English (PDF)

Published on: 6/04/2023

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### Package Leaflet

English (PDF)

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Labelling

English (PDF)

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