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

Active substance:
Ketoprofen

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Ketoprofen

150.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 2 day
- **Pig**
  - Meat and offal. 3 day
- **Cattle**
  - Milk. 0 hour

Intravenous use:

- **Cattle**
  - Meat and offal. 2 day
- **Horse**
  - Meat and offal. 1 day
- **Cattle**
  - Milk. 0 hour
- **Horse**
  - Milk. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AE03

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

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

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

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:
Ecuphar Veterinaria S.L.U.

Marketing authorisation date:
21/06/2010

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
170050

Date of authorisation status change:
12/01/2025

Reference member state:
Spain

Procedure number:
ES/V/0115/001

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)

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

English (PDF)
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