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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:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

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:
20/05/2010

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2156 ESP

Date of authorisation status change:
20/05/2010

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

Combined File of all Documents

Summary of Product Characteristics

English (PDF)
Published on: 5/04/2023
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Labelling

English (PDF)
Published on: 5/04/2023
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Package Leaflet

English (PDF)
Published on: 5/04/2023
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