DICLOVET 50 mg/ml SOLUCION INYECTABLE

Authorised

- Diclofenac sodium

Product identification

Medicine name:
DICLOVET 50 mg/ml SOLUCION INYECTABLE

Active substance:
Diclofenac sodium

Target species:
Cattle
Pig
Horse (non food-producing)

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Diclofenac sodium
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 15 day

- **Pig**
  - Meat and offal. 12 day

- **Horse (non food-producing)**
  - Meat and offal. no withdrawal period

Carne: Su uso no está autorizado en équidos cuya carne se destine al consumo humano

- **Cattle**
  - Milk. 6 day

- **Horse (non food-producing)**
  - Milk. no withdrawal period

Leche: Su uso no está autorizado en équidos cuya leche se destine al consumo humano

Intravenous use:

- **Horse (non food-producing)**
  - Meat and offal. no withdrawal period

Carne: Su uso no está autorizado en équidos cuya carne se destine al consumo humano

- **Horse (non food-producing)**
  - Milk. no withdrawal period

Leche: Su uso no está autorizado en équidos cuya leche se destine al consumo humano

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AB05

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

Authorisation status:
Valid

Authorised in:
Spain

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro Iberica S.L.

Marketing authorisation date:
20/03/2013

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
2760 ESP

Date of authorisation status change:
20/03/2013

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

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