DEXANIL 2 mg/ml SOLUCIÓN INYECTABLE PARA BOVINO, CABALLOS Y PORCINO

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
DEXANIL 2 mg/ml SOLUCIÓN INYECTABLE PARA BOVINO, CABALLOS Y PORCINO

Active substance:
Dexamethasone sodium phosphate

Target species:
Cattle
Horse
Pig

Route of administration:
Intramuscular use
Intraarticular use
Intravenous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate
2.63 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 7 day
  - Milk. 60 hour
- **Horse**
  - Meat and offal. 11 day
  - Milk. no withdrawal period

leche: Su uso no está autorizado en yeguas cuya leche se utiliza para consumo humano

- **Pig**
  - Meat and offal. 2 day
- **Cattle**
  - Meat and offal. 7 day
  - Milk. 60 hour
- **Horse**
  - Meat and offal. 11 day
  - Milk. no withdrawal period

leche: Su uso no está autorizado en yeguas cuya leche se utiliza para consumo humano

Intraarticular use:

- **Horse**
  - Meat and offal. 11 day
  - Milk. no withdrawal period

leche: Su uso no está autorizado en yeguas cuya leche se utiliza para consumo humano

- **Horse**
  - Meat and offal. 11 day
  - Milk. no withdrawal period

leche: Su uso no está autorizado en yeguas cuya leche se utiliza para consumo humano

Intravenous use:

- **Horse**
  - Meat and offal. 11 day
  - Milk. no withdrawal period

leche: Su uso no está autorizado en yeguas cuya leche se utiliza para consumo humano

- **Horse**
  - Meat and offal. 11 day
  - Milk. no withdrawal period

leche: Su uso no está autorizado en yeguas cuya leche se utiliza para consumo humano

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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:
29/10/2015

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3319 ESP

Date of authorisation status change:
1/01/2023

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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