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GLUCADEX 2 MG/ML SOLUTION FOR INJECTION FOR HORSES, CATTLE, GOATS, PIGS, DOGS AND CATS

Not authorised

- DEXAMETHASONE DISODIUM PHOSPHATE

Product identification

Medicine name:
GLUCADEX 2 MG/ML SOLUTION FOR INJECTION FOR HORSES, CATTLE, GOATS, PIGS, DOGS AND CATS

Active substance:
DEXAMETHASONE DISODIUM PHOSPHATE

Target species:
Cattle
Pig
Cat
Horse
Goat
Dog

Route of administration:
Intramuscular use
Subcutaneous use

Periarticular use
Intravenous use
Intraarticular use

Product details

Active substance and strength:
DEXAMETHASONE DISODIUM 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. 8 day
  - Milk. 72 hour
- **Pig**
  - Meat and offal. 2 day
- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

- **Goat**
  - Meat and offal. 8 day
  - Milk. 72 hour

Periarticular use:

- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

Intravenous use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 72 hour
- **Pig**
  - Meat and offal. 6 day
- **Horse**
  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

  - Meat and offal. 8 day
- **Goat**
  - Meat and offal. 8 day
  - Milk. 72 hour

Intraarticular use:

- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorized for use in horses producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

Authorisation status:
Revoked

Authorised in:
Spain

Package description:
Available only in French
Available only in French

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:
Kepro B.V.

Marketing authorisation date:
18/12/2019

Manufacturing sites for batch release:
Kepro B.V.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
3848 ESP

Date of authorisation status change:

19/06/2024

Reference member state:
France

Procedure number:
FR/V/0374/001

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

Documents

Summary of Product Characteristics

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Labelling

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

Package Leaflet

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

Combined File of all Documents

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