CALIERCORTIN 4 mg/ml solution for injection for cattle, pigs, horses, dogs and cats

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
CALIERCORTIN 4 mg/ml solution for injection for cattle, pigs, horses, dogs and cats

Active substance:
Dexamethasone sodium phosphate

Target species:
Cattle
Dog
Horse
Cat
Pig

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
5.30 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Milk. 4 day
  - Meat and offal. 16 day
- **Horse**
  - Meat and offal. 16 day
  - Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.

- **Pig**
  - Meat and offal. 4 day

Subcutaneous use:

- **Cattle**
  - Milk. 4 day
  - Meat and offal. 16 day
- **Horse**

- Meat and offal. 16 day
- Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.

- **Pig**
  - Meat and offal. 4 day

Intramuscular use:

- **Cattle**
  - Milk. 4 day
  - Meat and offal. 16 day
- **Horse**
  - Meat and offal. 16 day
  - Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.

- **Pig**
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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Authorisation status:
Valid

Authorised in:
Greece

Package description:

(ID2) 50 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber)
(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 10 millilitre(s), closed with Stopfen (bromobutyl rubber)

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:
Laboratorios Calier S.A.

Marketing authorisation date:
28/02/2019

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
21526/01-03-2019/K-0230701

Date of authorisation status change:
4/07/2021

Reference member state:
Germany

Procedure number:
DE/V/0179/001

Concerned member states:
Austria Bulgaria Croatia Cyprus Greece Italy Poland Portugal Romania

Spain

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

Documents

Summary of Product Characteristics

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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