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

---

Authorisation status:
Valid

Authorised in:
Italy

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:
12/08/2019

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

Responsible authority:
Ministry Of Health

Authorisation number:
105170

Date of authorisation status change:
12/08/2019

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

Combined File of all Documents

2402472-paren-20210113.pdf