# 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 Injektionslösung für Rinder, Schweine, Pferde, Hunde und Katzen
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)

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**Pharmaceutical form:**
Solution for injection

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QH02AB02

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

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**Authorisation status:**
Valid

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**Authorised in:**
Germany

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**Package description:**

(ID2): 1 unspecified outer container with 1 Vial (Glass) with 50 millilitre(s) (50 millilitre(s))
(ID1): 1 unspecified outer container with 1 Vial (Glass) with 10 millilitre(s) (10 millilitre(s))

## 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:**
8/02/2019

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

**Responsible authority:**
Federal Office Of Consumer Protection And Food Safety

**Authorisation number:**
402472.00.00

**Date of authorisation status change:**
8/02/2019

**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

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

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