

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:

Portugal

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:

Calier Portugal Medicamentos E Produtos Veterinarios S.A.

Marketing authorisation date:

14/12/2018

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1225/01/18DFVPT

Date of authorisation status change:

11/07/2025

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

English (PDF)

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