

Dexacortin 2 mg/mL ml solution for injection for cattle, pigs, dogs and cats

Not
authorised

- Dexamethasone sodium phosphate
- Dexamethasone sodium phosphate

Product identification

Medicine name:

Dexacortin 2 mg/mL ml solution for injection for cattle, pigs, dogs and cats

Active substance:

Dexamethasone sodium phosphate

Dexamethasone sodium phosphate

Target species:

Cattle

Pig

Cattle

Pig

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate

2.00 milligram(s) / 1.00 millilitre(s)

Dexamethasone sodium phosphate

2.00 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

Intravenous use:

-

Cattle

- Meat and offal. 8 day

- Milk. 72 hour

-

Pig

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Slovenia

Package description:

Clear glass type I vials, closed with a bromobutylrubber stopper and sealed with an aluminium cap. One vial of 100 ml in a carton box.

Clear glass type I vials, closed with a bromobutylrubber stopper and sealed with an aluminium cap. One vial of 50 ml in a carton box.

Clear glass type I vials, closed with a bromobutylrubber stopper and sealed with an aluminium cap. One vial of 20 ml in a carton box.

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:

Intervet International B.V.

Marketing authorisation date:

16/02/2010

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

DC/V/0086/001

Date of authorisation status change:

21/08/2025

Reference member state:

Portugal

Procedure number:

PT/V/0136/001

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.

Labelling

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

Combined File of all Documents