# DEXAFORT SUSPENSION INJECTABLE

Authorised

- DEXAMETHASONE DISODIUM PHOSPHATE
- DEXAMETHASONE 21-PHENYLPROPIONATE

## Product identification

**Medicine name:**
DEXAFORT SUSPENSION INJECTABLE

**Active substance:**
DEXAMETHASONE DISODIUM PHOSPHATE
DEXAMETHASONE 21-PHENYLPROPIONATE

**Target species:**
Cattle
Pig
Cat
Equid
Goat
Dog

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
DEXAMETHASONE DISODIUM PHOSPHATE
1.32 milligram(s) / 1.00 millilitre(s)

DEXAMETHASONE 21-PHENYLPROPIONATE
2.67 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**
Suspension for injection

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**Withdrawal period by route of administration:**
**Intramuscular use:**

- **Cattle**
  - Meat and offal. 63 day
  - Milk. 8 day
- **Pig**
  - Meat and offal. 14 day
- **Equid**
  - Meat and offal. 63 day
  - Milk. 8 day
- **Goat**
  - Meat and offal. 63 day
  - Milk. 8 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

**Authorisation status:**
Valid

**Authorised in:**
France

**Package description:**
Available only in French

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Legal basis not covered by Directive 2001/82/EC

**Marketing authorisation holder:**
Intervet

**Marketing authorisation date:**
30/06/1992

**Manufacturing sites for batch release:**
Vet Pharma Friesoythe GmbH

**Responsible authority:**
French Agency For Food, Environmental And Occupational Health & Safety

**Authorisation number:**
FR/V/1475011 1/1992

**Date of authorisation status change:**
30/06/2012

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 and Labelling

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