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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)

Pharmaceutical form:
Suspension for injection

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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Package Leaflet and Labelling

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