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DEXADRESON 2 MG/ML SOLUTION INJECTABLE POUR CHEVAUX BOVINS CAPRINS PORCINS CHIENS ET CHATS

Authorised

- DEXAMETHASONE DISODIUM PHOSPHATE

Product identification

Medicine name:
DEXADRESON 2 MG/ML SOLUTION INJECTABLE POUR CHEVAUX BOVINS CAPRINS PORCINS CHIENS ET CHATS

Active substance:
DEXAMETHASONE DISODIUM PHOSPHATE

Target species:
Cattle
Pig
Equid
Horse
Goat
Dog
Cat

Route of administration:
Intramuscular use

Subcutaneous use
Periarticular use
Intravenous use
Intraarticular use

Product details

Active substance and strength:
DEXAMETHASONE DISODIUM PHOSPHATE
2.63 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. 3 day
- **Pig**
  - Meat and offal. 2 day
- **Equid**
  - Meat and offal. 8 day
- **Horse**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

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Goat

- Meat and offal. 8 day
- Milk. 3 day

Periarticular use:

- **Horse**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine..

- **Equid**
  - Meat and offal. 8 day

Intravenous use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 6 day
- **Equid**
  - Meat and offal. 8 day
- **Horse**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

-

Goat

- Meat and offal. 8 day
- Milk. 3 day

Intraarticular use:

- **Horse**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

- **Equid**
  - Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French
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:
Intervet International GmbH

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

Authorisation number:
FR/V/4665198 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.