

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.

-

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

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

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

Package Leaflet and Labelling

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