DEXALONE SOLUTION INJECTABLE POUR EQUINS BOVINS CAPRINS ET PORCINS

Authorised

- DEXAMETHASONE DISODIUM PHOSPHATE

Product identification

Medicine name:
DEXALONE SOLUTION INJECTABLE POUR EQUINS BOVINS CAPRINS ET PORCINS

Active substance:
DEXAMETHASONE DISODIUM PHOSPHATE

Target species:
Cattle
Pig
Equid
Goat

Route of administration:
Intramuscular use
Subcutaneous use
Periarticular use
Intravenous use
Intraarticular use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 3 day
  - Meat and offal. 8 day
- **Pig**
  - Meat and offal. 6 day
- **Equid**
  - Milk. 3 day
  - Meat and offal. 6 day
- **Goat**
  - Milk. 3 day
  - Meat and offal. 8 day

Subcutaneous use:

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

- **Pig**
  - Meat and offal. 6 day
- **Equid**
  - Milk. 3 day
  - Meat and offal. 6 day
- **Goat**
  - Milk. 3 day
  - Meat and offal. 8 day

Periarticular use:

- **Cattle**
  - Milk. 3 day
  - Meat and offal. 8 day
- **Pig**
  - Meat and offal. 6 day
- **Equid**
  - Milk. 3 day
  - Meat and offal. 6 day
- **Goat**
  - Milk. 3 day
  - All relevant tissues. 8 day

Intravenous use:

-

Cattle

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

•

Pig

- Meat and offal. 6 day

•

Equid

- Milk. 3 day
- Meat and offal. 6 day

•

Goat

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

Intraarticular use:

•

Cattle

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

•

Pig

- Meat and offal. 6 day

•

Equid

- Milk. 3 day
- Meat and offal. 6 day

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Goat

- Milk. 3 day

- Meat and offal. 6 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
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:
Dopharma France S.A.S.

Marketing authorisation date:
24/07/1992

Manufacturing sites for batch release:
Dopharma France

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

Authorisation number:

FR/V/4409579 2/1992

Date of authorisation status change:
24/07/2012

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

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