DEXAMETHASONE VMD 2 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS, GOATS, HORSES, DOGS AND CATS

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
DEXAMETHASONE VMD 2 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS, GOATS, HORSES, DOGS AND CATS

Active substance:
Dexamethasone sodium phosphate

Target species:
Cattle
Pig
Cat
Equid
Goat
Dog

Route of administration:
Intramuscular use
Subcutaneous use

Periarticular use
Intravenous use
Intraarticular use

Product details

Active substance and strength:
Dexamethasone sodium 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
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

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

Periarticular use:

- **Equid**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

Intravenous use:

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

- **Pig**
  - Meat and offal. 6 day

- **Equid**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

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

Intraarticular use:

- **Equid**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Box of 1 vial of 25 mL
Box of 1 vial of 50 mL
Box of 1 vial of 100 mL

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
26/09/2025

Manufacturing sites for batch release:
V.M.D.
Laboratoires Biove

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

Authorisation number:
FR/V/6759815 5/2025

Date of authorisation status change:

26/09/2025

Reference member state:
France

Procedure number:
FR/V/0505/001

Concerned member states:
Belgium Bulgaria Estonia Hungary Latvia Lithuania Luxembourg Netherlands Poland Romania

Generic of:
600000032138

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet and Labelling

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

Summary of Product Characteristics

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

Combined File of all Documents

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