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:
Romania

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:
3/11/2025

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
250148

Date of authorisation status change:

3/11/2025

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

Combined File of all Documents

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