

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:

Hungary

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:

30/10/2025

Manufacturing sites for batch release:

V.M.D.

Laboratoires Biove

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

30/10/2025

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

eu-puar-frv0505001-mr-rpe914-en.pdf