

# MULTIBIO D

Authorised

- Dexamethasone acetate
- COLISTIN SULFATE
- Ampicillin

## Product identification

**Medicine name:**

MULTIBIO D

---

**Active substance:**

Dexamethasone acetate

COLISTIN SULFATE

Ampicillin

---

**Target species:**

Horse

Cattle

Pig

---

**Route of administration:**

Intramuscular use

Subcutaneous use

Intraperitoneal use

---

## Product details

### **Active substance and strength:**

Dexamethasone acetate

0.25 milligram(s) / 1.00 millilitre(s)

COLISTIN SULFATE

250000.00 international unit(s) / 1.00 millilitre(s)

Ampicillin

100.00 milligram(s) / 1.00 millilitre(s)

---

### **Pharmaceutical form:**

Suspension for injection

---

### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

- 

#### **Horse**

- Meat and offal. 14 day

Nu se va administra la iepile al căror lapte este destinat consumului uman.

- 

#### **Cattle**

- Meat and offal. 14 day

- Milk. 2 day

- 

#### **Pig**

- Meat and offal. 14 day

#### **Subcutaneous use:**

- 

#### **Cattle**

- Meat and offal. 14 day

- Milk. 2 day

- 

**Horse**

- Meat and offal. 14 day

Nu se va administra la iepele al căror lapte este destinat consumului uman.

- 

**Pig**

- Meat and offal. 14 day

**Intraperitoneal use:**

- 

**Cattle**

- Meat and offal. 14 day

- Milk. 2 day

- 

**Horse**

- Meat and offal. 14 day

Nu se va administra la iepele al căror lapte este destinat consumului uman.

- 

**Pig**

- Meat and offal. 14 day

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01RV01

---

**Legal status of supply:**

This information is not available for this product.

---

**Authorisation status:**

Valid

---

**Authorised in:**

Romania

---

**Available in:**

Romania

---

**Package description:**

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Complete application (stand-alone) - Directive No 2001/82/EC

---

**Marketing authorisation holder:**

Virbac

---

**Marketing authorisation date:**

29/03/2007

---

**Manufacturing sites for batch release:**

Virbac

---

**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

---

**Authorisation number:**

150063

---

**Date of authorisation status change:**

28/08/2025

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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