

# ALFADEXX 2 MG/ML SOLUTION FOR INJECTION FOR HORSES, CATTLE, GOATS, PIGS, DOGS AND CATS

Authorised

- DEXAMETHASONE DISODIUM PHOSPHATE

## Product identification

### **Medicine name:**

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

### **Active substance:**

DEXAMETHASONE DISODIUM PHOSPHATE

### **Target species:**

Cattle

Pig

Cat

Horse

Goat

Dog

### **Route of administration:**

Intramuscular use

Subcutaneous use

Periarticular use  
Intravenous use  
Intraarticular use

---

## Product details

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

DEXAMETHASONE DISODIUM 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. 72 hour

- 

#### **Pig**

- Meat and offal. 2 day

- 

#### **Horse**

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

Not authorized for use in horses producing milk for human consumption.

- 

#### **Goat**

- Meat and offal. 8 day
- Milk. 72 hour

### **Periarticular use:**

- 

**Horse**

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

Not authorized for use in horses producing milk for human consumption.

**Intravenous use:**

- 

**Cattle**

- Meat and offal. 8 day
- Milk. 72 hour

- 

**Pig**

- Meat and offal. 6 day

- 

**Horse**

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

Not authorized for use in horses producing milk for human consumption.

- 

**Goat**

- Meat and offal. 8 day
- Milk. 72 hour

**Intraarticular use:**

- 

**Horse**

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

Not authorized for use in horses producing milk for human consumption.

---

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

QH02AB02

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Italy

---

**Package description:**

Available only in [French](#)

Available only in [French](#)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Alfasan Nederland B.V.

---

**Marketing authorisation date:**

19/01/2023

---

**Manufacturing sites for batch release:**

Produlab Pharma B.V.

Alfasan Nederland B.V.

---

**Responsible authority:**

Ministry Of Health

---

**Authorisation number:**

105549

---

**Date of authorisation status change:**

19/01/2023

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0430/001

---

**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Netherlands Norway Poland Portugal Romania Slovakia  
Slovenia Spain Sweden

---

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

English (PDF)

Published on: 14/03/2026

[Download](#)

eu-puar-frv0430001-mr-rpe652-en.pdf