

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

Authorised

- Dexamethasone sodium phosphate

## Product identification

### Medicine name:

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

### Active substance:

Dexamethasone sodium phosphate

### Target species:

Cattle

Pig

Cat

Horse

Goat

Dog

### Route of administration:

Intramuscular use

Subcutaneous use

Periarticular use  
Intravenous use  
Intraarticular use

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## Product details

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

Dexamethasone sodium phosphate  
2.63 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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

#### **Intramuscular use:**

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#### **Cattle**

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

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#### **Pig**

- Meat and offal. 2 day

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#### **Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day

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#### **Goat**

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

### **Periarticular use:**

- 

### **Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day

### **Intravenous use:**

- 

### **Cattle**

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

- 

### **Pig**

- Meat and offal. 6 day

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### **Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day

- 

### **Goat**

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

### **Intraarticular use:**

- 

### **Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH02AB02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Romania

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**Package description:**

Available only in French

Available only in French

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

13/09/2021

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

Alfasan Nederland B.V.

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

260132

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**Date of authorisation status change:**

10/09/2026

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**Reference member state:**

France

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**Procedure number:**

FR/V/0430/001

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**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

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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)

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