

# 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

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## Product details

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

DEXAMETHASONE DISODIUM 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**

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

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#### **Pig**

- Meat and offal. 2 day

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#### **Horse**

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

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

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#### **Goat**

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

### **Periarticular use:**

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**Horse**

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

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

**Intravenous use:**

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**Cattle**

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

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**Pig**

- Meat and offal. 6 day

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**Horse**

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

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

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**Goat**

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

**Intraarticular use:**

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**Horse**

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

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

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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:**

Belgium

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**Available in:**

Belgium

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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:**

11/01/2022

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

Alfasan Nederland B.V.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V594373

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**Date of authorisation status change:**

11/01/2022

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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

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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

Combined File of all Documents

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