

# CORTEXONAVET 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

Authorised

- Dexamethasone sodium phosphate

## Product identification

**Medicine name:**

CORTEXONAVET 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

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**Active substance:**

Dexamethasone sodium phosphate

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**Target species:**

Horse

Cattle

Dog

Cat

Pig

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**Route of administration:**

Intraarticular use

Intramuscular use

Intravenous 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:**

#### **Intraarticular use:**

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##### **Horse**

- Meat and offal. 8 day

#### **Intramuscular use:**

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##### **Cattle**

- Meat and offal. 8 day

- Milk. 72 hour

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##### **Horse**

- Meat and offal. 8 day

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##### **Pig**

- Meat and offal. 2 day

#### **Intravenous use:**

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##### **Horse**

- 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:**

Hungary

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**Available in:**

Hungary

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**Package description:**

Cardboard box with 1 colourless, type I glass vial of 50 ml, which is closed with a bromobutyl type I rubber stopper and sealed with an aluminium cap.

Cardboard box with 1 colourless, type I glass vial of 100 ml, which is closed with a bromobutyl type I rubber stopper and sealed with an aluminium cap.

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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:**

Laboratorios Syva S.A.

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**Marketing authorisation date:**

21/03/2018

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**Manufacturing sites for batch release:**

Laboratorios Syva S.A.

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

3949/X/18 NÉBIH ÁTI

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**Date of authorisation status change:**

21/03/2018

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0351/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Hungary Italy Poland Portugal Romania  
Spain

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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

Summary of Product Characteristics

Combined File of all Documents