

# Rapidexon 2 mg/ml solution for injection

Authorised

- Dexamethasone sodium phosphate

## Product identification

**Medicine name:**

Rapidexon 2 mg/ml solution for injection

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**Active substance:**

Dexamethasone sodium phosphate

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**Target species:**

Horse

Cattle

Pig

Dog

Cat

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**Route of administration:**

Intraarticular use

Intramuscular use

Intravenous use

Intrabursal use

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## Product details

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

Dexamethasone sodium phosphate  
2.00 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**

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

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##### **Pig**

- Meat and offal. 2 day

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##### **Horse**

- Meat and offal. 8 day

#### **Intravenous use:**

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##### **Horse**

- Meat and offal. 8 day

#### **Intrabursal 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:**

Netherlands

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### **Package description:**

50 ml vial glass type I uncoloured with bromobutyl rubber stopper type I secured with aluminium cap

25 ml vial (filled in 30 ml vial) glass type I uncoloured with bromobutyl rubber stopper type I secured with aluminium cap

100 ml vial glass type I uncoloured with bromobutyl rubber stopper type I secured with aluminium

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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:**

Eurovet Animal Health B.V.

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### **Marketing authorisation date:**

31/03/2008

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### **Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 101405

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**Date of authorisation status change:**

6/09/2024

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0284/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Finland France Greece Hungary Ireland  
Lithuania Poland Portugal Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

Published on: 1/05/2025

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