

# 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:**

Finland

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### **Available in:**

Finland

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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:**

1/10/2009

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

24307

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**Date of authorisation status change:**

1/10/2009

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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  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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

Package Leaflet

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

Combined File of all Documents

Rapidexon 2 mg.pdf