

# Dexa-ject 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

Authorised

- Dexamethasone sodium phosphate

## Product identification

### Medicine name:

Dexa-ject 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

### Active substance:

Dexamethasone sodium phosphate

### Target species:

Horse

Cattle

Dog

Cat

Pig

### Route of administration:

Intraarticular use

Intramuscular use

Intravenous use

## 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:**

Estonia

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**Available in:**

Estonia

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**Package description:**

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

Cardboard box with 1 colourless, type I glass vial of 50 ml which is closed with a bromobutyl 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:**

Dopharma Research B.V.

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**Marketing authorisation date:**

25/09/2012

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**Manufacturing sites for batch release:**

Dopharma B.V.

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**Responsible authority:**

State Agency Of Medicines

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**Authorisation number:**

1730

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**Date of authorisation status change:**

25/09/2012

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0293/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland France  
Germany Greece Hungary Iceland Italy Latvia Lithuania Netherlands  
Norway Poland Romania Slovakia Spain Sweden  
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

English (PDF)

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