

# 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  
DEXA-JECT 2 MG/ML SOLUTION INJECTABLE POUR BOVINS CHEVAUX PORCINS CHIENS ET CHATS

### 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:**

France

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**Available in:**

France

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**Package description:**

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.

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.

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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:**

20/07/2012

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**Manufacturing sites for batch release:**

Dopharma B.V.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/5391841 3/2012

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**Date of authorisation status change:**

25/08/2017

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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)

Published on: 3/05/2024

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### Package Leaflet and Labelling

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