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# 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:**

- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

**Intramuscular use:**

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

- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 2 day

**Intravenous use:**

- **Horse**
  - Meat and offal. 8 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QH02AB02

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Denmark

**Available in:**
Denmark

**Package description:**
Cardboard box with 1 colourless, type I glass vial of 100 ml which is closedwith a bromobutyl rubber stopper and sealed with an aluminium cap.
Cardboard box with 1 colourless, type I glass vial of 50 ml which is closedwith a bromobutyl rubber stopper and sealed with an aluminium cap.

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Dopharma Research B.V.

**Marketing authorisation date:**
2/08/2012

**Manufacturing sites for batch release:**
Dopharma B.V.

**Responsible authority:**
Danish Medicines Agency

**Authorisation number:**
49408

**Date of authorisation status change:**
2/08/2012

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0293/001

**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)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

English (PDF)

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