

# Dexamecine 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

Authorised

- Dexamethasone sodium phosphate

## Product identification

**Medicine name:**

Dexamecine 2 mg/ml solution for injection for cattle, horses, pigs, dogs and cats

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**Active substance:**

Dexamethasone sodium phosphate

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**Target species:**

Cattle

Pig

Horse

Dog

Cat

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**Route of administration:**

Intraarticular use

Intramuscular use

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## 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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#### **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

#### **Intramuscular use:**

- 

#### **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

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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:**

Italy

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**Available in:**

Italy

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**Package description:**

100 ml amber co-ex plastic (polypropylene) vials closed with bromobutyl rubber stoppers and aluminium caps. The bottles are individually packaged in a cardboard box, the package leaflet is enclosed.

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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:**

Vet-Agro Trading Sp. z o.o.

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**Marketing authorisation date:**

23/05/2012

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**Manufacturing sites for batch release:**

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

105124

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**Date of authorisation status change:**

23/05/2012

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0268/001

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**Concerned member states:**

Hungary Italy 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

Combined File of all Documents

English (PDF)

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