# 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

**Active substance:**
Dexamethasone sodium phosphate

**Target species:**
Cattle
Pig
Horse
Dog
Cat

**Route of administration:**
Intraarticular use
Intramuscular 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:**

- **Cattle**
  - Milk. 72 hour
  - Meat and offal. 8 day
- **Pig**
  - Meat and offal. 2 day
- **Horse**
  - Meat and offal. 8 day

**Intramuscular use:**

- **Cattle**
  - Milk. 72 hour
  - Meat and offal. 8 day
- **Pig**
  - Meat and offal. 2 day
- **Horse**

- Meat and offal. 8 day

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

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

**Authorisation status:**
Valid

**Authorised in:**
Hungary

**Available in:**
Hungary

**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.

## 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:**
Vet-Agro Trading Sp. z o.o.

**Marketing authorisation date:**
14/03/2018

**Manufacturing sites for batch release:**
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

**Responsible authority:**
Directorate Of Veterinary Medicinal Products

**Authorisation number:**
3942/1/18 NÉBIH ÁTI

**Date of authorisation status change:**
14/03/2018

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0268/001

**Concerned member states:**
Hungary Italy United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

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Combined File of all Documents