

# Dexashot, 2mg/ml, Solution for injection

Authorised

- Dexamethasone

## Product identification

**Medicine name:**

Dexashot, 2mg/ml, Solution for injection

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**Active substance:**

Dexamethasone

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**Target species:**

Horse

Cattle

Pig

Cat

Dog

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**Route of administration:**

Intraarticular use

Intravenous use

Intramuscular use

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## Product details

**Active substance and strength:**

Dexamethasone

2.00 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**

- Milk. no withdrawal period

Do not use in horses producing milk for human consumption.,

- Meat and offal. 8 day

**Intravenous use:**

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**Horse**

- Milk. no withdrawal period

Do not use in horses producing milk for human consumption.,

- Meat and offal. 8 day

**Intramuscular 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**

- Milk. no withdrawal period

Do not use in horses producing milk for human consumption.,

- 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:**

Austria

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**Available in:**

Austria

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**Package description:**

Available only in German

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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:**

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

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**Marketing authorisation date:**

31/05/2016

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**Manufacturing sites for batch release:**

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

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

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

836943

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**Date of authorisation status change:**

31/05/2016

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0132/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia France Greece Ireland Poland Portugal  
Romania Spain

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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

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.