

# Dexrapid, 2mg/ml, Solution for injection

Authorised

- Dexamethasone

## Product identification

**Medicine name:**

Dexrapid, 2mg/ml, Solution for injection

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**Active substance:**

Dexamethasone

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**Target species:**

Horse

Pig

Cattle

Dog

Cat

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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

Not authorised for use in mares producing milk for human consumption.,

- Meat and offal. 8 day

**Intravenous use:**

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**Horse**

- Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.,

- Meat and offal. 8 day

**Intramuscular use:**

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**Horse**

- Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.,

- Meat and offal. 8 day

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**Pig**

- Meat and offal. 2 day

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**Cattle**

- Milk. 72 hour

- 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:**

Vetviva Richter GmbH

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**Marketing authorisation date:**

11/11/2020

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**Manufacturing sites for batch release:**

Vetviva Richter GmbH

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

840391

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**Date of authorisation status change:**

11/11/2020

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0167/001

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**Concerned member states:**

Austria Belgium Bulgaria Denmark Finland France Germany Greece  
Hungary Ireland Lithuania Netherlands Poland Portugal Romania Slovakia  
Slovenia Spain Sweden

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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

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

### Package Leaflet

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.

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