

# Dexrapid, 2mg/ml, Solution for injection

Authorised

- Dexamethasone

## Product identification

**Medicine name:**

Dexrapid, 2mg/ml, Solution for injection

---

**Active substance:**

Dexamethasone

---

**Target species:**

Horse

Pig

Cattle

Dog

Cat

---

**Route of administration:**

Intraarticular use

Intravenous use

Intramuscular use

---

## Product details

**Active substance and strength:**

Dexamethasone

2.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for injection

---

**Withdrawal period by route of administration:**

**Intraarticular use:**

- 

**Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day

**Intravenous use:**

- 

**Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day

**Intramuscular use:**

- 

**Horse**

- Milk. no withdrawal period

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

- Meat and offal. 8 day

- 

**Pig**

- Meat and offal. 2 day

- 

**Cattle**

- Milk. 72 hour

- 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

---

**Package description:**

Glass Vial 1 x 100.0 millilitre(s)

---

## 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:**

Vetviva Richter GmbH

---

**Marketing authorisation date:**

6/10/2020

---

**Manufacturing sites for batch release:**

Vetviva Richter GmbH

---

**Responsible authority:**

Directorate Of Veterinary Medicinal Products

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

6/10/2020

---

**Reference member state:**

Czechia

---

**Procedure number:**

CZ/V/0167/001

---

**Concerned member states:**

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

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)