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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:
Poland

Available in:
Poland

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:
4/03/2021

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3076

Date of authorisation status change:
4/03/2021

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

Documents

Labelling

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.

Summary of Product Characteristics

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