Dexashot, 2mg/ml, Solution for injection

Authorised

- Dexamethasone

Product identification

Medicine name:
Dexashot, 2mg/ml, Solution for injection

Active substance:
Dexamethasone

Target species:
Horse
Cattle
Pig
Cat
Dog

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

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

  - Meat and offal. 8 day

Intravenous use:

- **Horse**
  - Milk. no withdrawal period

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

  - Meat and offal. 8 day

Intramuscular use:

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

- **Pig**
  - Meat and offal. 2 day

- **Horse**
  - Milk. no withdrawal period

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

- 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:
Spain

Available in:
Spain

Package description:
Plastic 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:
Vet-Agro Multi-Trade Company Sp. z o.o.

Marketing authorisation date:
29/07/2016

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

Responsible authority:

Spanish Agency For Medicines And Medical Devices

Authorisation number:
3455 ESP

Date of authorisation status change:
30/07/2016

Reference member state:
Czechia

Procedure number:
CZ/V/0132/001

Concerned member states:
Austria Belgium Bulgaria Croatia France Greece Ireland Poland Portugal Romania Spain

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

Documents

Summary of Product Characteristics

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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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