Dexashot, 2mg/ml, Solution for injection

Authorised

- Dexamethasone

Product identification

Medicine name:
Dexashot, 2mg/ml, Solution for injection
DEXASHOT 2 MG/ML SOLUTION INJECTABLE POUR BOVINS CHEVAUX PORCINS CHIENS ET CHATS

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

- **Cat**
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

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:
4/09/2019

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1158702 8/2019

Date of authorisation status change:
10/06/2021

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 and Labelling

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