

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:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Przedsiębiorstwo 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.

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