

DEXAVEX 2 mg/ml solution for injection

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

DEXAVEX 2 mg/ml solution for injection

Active substance:

Dexamethasone sodium phosphate

Target species:

Cattle

Horse

Pig

Dog

Goat

Cat

Route of administration:

Intramuscular use

Intraarticular use

Intravenous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate
2.63 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

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

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption

-

Pig

- Meat and offal. 2 day

-

Goat

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

-

Cattle

- Meat and offal. 8 day

- Milk. 72 hour

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption

-

Goat

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

Intraarticular use:

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption

Intravenous use:

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption

-

Goat

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

-

Horse

- Meat and offal. 8 day
- Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption

-

Goat

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

box containing 10 vials of 50 ml

box containing 1 vial of 50 ml

Box containing 1 vial of 100 ml

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:

S P Veterinaria S.A.

Marketing authorisation date:

15/01/2013

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

2693 ESP

Date of authorisation status change:

22/01/2013

Reference member state:

Spain

Procedure number:

ES/V/0271/001

Concerned member states:

Portugal

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

Documents

Combined File of all Documents

Summary of Product Characteristics

English (PDF)

Published on: 27/06/2024

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

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