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Vetdrax 100 mg/ml solution for injection for cattle, pigs and sheep

Authorised

- Tulathromycin

Product identification

Medicine name:

Vetdrax 100 mg/ml solution for injection for cattle, pigs and sheep

Active substance:

Tulathromycin

Target species:

Cattle

Pig

Sheep

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Tulathromycin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

•

Cattle

- Meat and offal. 22 day

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Cattle

- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition

Intramuscular use:

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Pig

- Meat and offal. 13 day

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Sheep

- Meat and offal. 16 day

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Sheep

- Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA94

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

Package description:

Available only in Spanish

Available only in Spanish

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:

Vetpharma Animal Health S.L.

Marketing authorisation date:

7/05/2021

Manufacturing sites for batch release:

Mevet S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

4005 ESP

Date of authorisation status change:

8/05/2021

Reference member state:

Spain

Procedure number:

ES/V/0391/001

Concerned member states:

Austria Belgium Denmark France Germany Greece Hungary Italy
Netherlands Poland Portugal Romania United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 1/02/2024

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Package Leaflet

English (PDF)

Published on: 26/12/2023

[Download](#)

Labelling

English (PDF)

Published on: 26/12/2023

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Combined File of all Documents

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