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

Authorised

- Tulathromycin

Product identification

Medicine name:
Vetdrax 100 mg/ml solution for injection for cattle, pigs and sheeps

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

- **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:

- **Pig**
  - Meat and offal. 13 day

- **Sheep**
  - Meat and offal. 16 day

- **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:
Greece

Available in:
Greece

Package description:
cardboard box containing 1 vial of 100 ml
cardboard box containing 1 vial of 250 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:
Vetpharma Animal Health S.L.

Marketing authorisation date:
2/08/2021

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
34340/18-03-2026/K-0245201

Date of authorisation status change:

17/03/2026

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

Combined File of all Documents

Package Leaflet

Labelling

Summary of Product Characteristics

English (PDF)

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