

Tullavis 25 mg/ml solution for injection for pigs

Authorised

- Tulathromycin

Product identification

Medicine name:

Tullavis 25 mg/ml solution for injection for pigs

Active substance:

Tulathromycin

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Tulathromycin

25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Pig

- Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA94

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 250 m

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 100 ml

Clear glass (Type II Ph. Eur.) vial closed with a bromobutyl rubber stopper and sealed with an aluminium cap. Cardboard box containing 1 vial of 50 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:

Industrial Veterinaria S.A.

Marketing authorisation date:

20/08/2020

Manufacturing sites for batch release:

aniMedica Herstellungs GmbH

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/3523192 1/2020

Date of authorisation status change:

20/08/2020

Reference member state:

Ireland

Procedure number:

IE/V/0892/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Italy Latvia Lithuania Netherlands Poland
Portugal Romania Slovakia Slovenia Spain 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: 22/12/2023

[Download](#)

Package Leaflet and Labelling

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

Combined File of all Documents

eu-PUAR-esv0359001-dcp-tulavis-25-mg-ml-solution-for-injection-for-pigs-en.pdf