Huvexxin 25 mg/ml solution for injection for pigs

Authorised

- Tulathromycin

Product identification

Medicine name:
Huvexxin, 25mg/ml, Injekční roztok
Huvexxin 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:
Czechia

Package description:
Type I colourless glass vial with a chlorobutyl rubber stopper and aluminum overseal.
Pack sizes: Cardboard box containing one vial of 250 ml.
Type I colourless glass vial with a chlorobutyl rubber stopper and aluminum overseal.
Pack sizes: Cardboard box containing one vial of 100 ml.
Type I colourless glass vial with a chlorobutyl rubber stopper and aluminum overseal.
Pack sizes: Cardboard box containing one 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:
HuVepharma

Marketing authorisation date:
30/12/2022

Manufacturing sites for batch release:

Biovet J.S.C.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/040/22-C

Date of authorisation status change:
30/12/2022

Reference member state:
Ireland

Procedure number:
IE/V/0662/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Malta 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: 3/05/2024
Download

Package Leaflet

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Labelling

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