Tuloxxin 25 mg/ml solution for injection for pigs

Authorised

- Tulathromycin

Product identification

Medicine name:
Tuloxxin 25 mg/ml solution for injection for pigs
Tuloxxin 25 mg/ml Roztwór do wstrzykiwań

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:
Poland

Package description:
Cardboard box containing one clear type I glass vial of 50 ml, with a type I chlorobutyl/butyl film laminated rubber stopper and aluminium cap with plastic tear tab (flip-off).
Cardboard box containing one clear type I glass vial of 100 ml with a type I chlorobutyl/butyl film laminated rubber stopper and aluminium cap with plastic tear tab (flip-off).
Cardboard box containing one clear type I glass vial of 250 ml, with a type I chlorobutyl/butyl film laminated rubber stopper and aluminium cap with plastic tear tab (flip-off).

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
10/11/2020

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3040

Date of authorisation status change:
10/11/2020

Reference member state:
Ireland

Procedure number:
IE/V/0396/002

Concerned member states:
Belgium Bulgaria Croatia Czechia Estonia 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: 3/05/2024
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Package Leaflet

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Labelling

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