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Huvexxin 25 mg/ml solution for injection for pigs

Authorised

- Tulathromycin

Product identification

Medicine name:
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:
Germany

Available in:
Germany

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:
20/12/2022

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7007045.00.00

Date of authorisation status change:
20/12/2022

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)

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Documents

Combined File of all Documents

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