

Huvexxin 100 mg/ml solution for injection for cattle, pigs and sheep

Authorised

- Tulathromycin

Product identification

Medicine name:

Huvexxin 100 mg/ml solution for injection for cattle, pigs and sheep

Active substance:

Tulathromycin

Target species:

Sheep

Pig

Cattle

Route of administration:

Intramuscular use

Subcutaneous 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:

Intramuscular use:

-

Sheep

- Meat and offal. 16 day

-

Pig

- Meat and offal. 13 day

Subcutaneous use:

-

Cattle

- Meat and offal. 22 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 an aluminium overseal. Pack sizes: Cardboard box containing one vial of 100 ml.

Type I colourless glass vial with a chlorobutyl rubber stopper and an aluminium overseal.Pack sizes: Cardboard box containing one vial of 20 ml.

Type I colourless glass vial with a chlorobutyl rubber stopper and an aluminium overseal.Pack sizes: Cardboard box containing one vial of 250 ml.

Type I colourless glass vial with a chlorobutyl rubber stopper and an aluminium 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:

V7007046.00.00

Date of authorisation status change:

20/12/2022

Reference member state:

Ireland

Procedure number:

IE/V/0662/002

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

Combined File of all Documents

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