

Ladoxyn 500 mg/g granules for oral solution for pigs, chickens and turkeys

Authorised

- Doxycycline

Product identification

Medicine name:

Ladoxyn 500 mg/g granules for oral solution for pigs, chickens and turkeys

Ladoxyn 500 mg/g grânulos para solução oral para suínos galinhas e perus

Active substance:

Doxycycline

Target species:

Pig

Chicken

Turkey

Route of administration:

Oral use

Product details

Active substance and strength:

Doxycycline

500.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Granules for oral solution

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

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Pig

- Meat and offal. 4 day

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Chicken

- Meat and offal. 5 day

Not permitted for use in laying birds producing eggs for human consumption.

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Turkey

- Meat and offal. 12 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

1 kg round polypropylene container with polypropylene lid and inner bag of LDPE.

5 kg round polypropylene container with polypropylene lid and inner bag of LDPE.

5 kg stand up bag with zip lock

1 kg stand up bag with zip lock

5 kg square polypropylene container with polypropylene lid and inner bag of LDPE.

1 kg square polypropylene container with polypropylene lid and inner bag of LDPE.

100 g polypropylene container with polypropylene lid and inner bag of LDPE

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Lavet Kft.

Marketing authorisation date:

8/06/2009

Manufacturing sites for batch release:

Lavet Kft.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

181/01/09RFVPT

Date of authorisation status change:

1/04/2013

Reference member state:

Hungary

Procedure number:

HU/V/0104/001

Concerned member states:

Austria Czechia Denmark Germany Greece Italy Portugal

United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Combined File of all Documents

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