

Karidox Doxycycline 500 mg/g Powder for use in Drinking Water for Pigs, Chickens and Turkeys

Authorised

- Doxycycline hyclate

Product identification

Medicine name:

Karidox Doxycycline 500 mg/g Powder for use in Drinking Water for Pigs, Chickens and Turkeys

Active substance:

Doxycycline hyclate

Target species:

Chicken (for reproduction)

Chicken (broiler)

Pig (for fattening)

Turkey (for reproduction)

Turkey (for meat production)

Route of administration:

In drinking water use

Product details

Active substance and strength:

Doxycycline hyclate

580.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

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Chicken (for reproduction)

- Meat and offal. 5 day
- Eggs. no withdrawal period

Eggs: Not authorised for use in birds producing eggs for human consumption

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Chicken (broiler)

- Meat and offal. 5 day
- Eggs. no withdrawal period

Eggs: Not authorised for use in birds producing eggs for human consumption

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Pig (for fattening)

- Meat and offal. 4 day

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Turkey (for reproduction)

- Meat and offal. 12 day
- Eggs. no withdrawal period

Eggs: Not authorised for use in birds producing eggs for human consumption

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Turkey (for meat production)

- Meat and offal. 12 day
- Eggs. no withdrawal period

Eggs: Not authorised for use in birds producing eggs for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

bag of 1 kg
bag of 200 g

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:

Laboratorios Karizoo S.A.

Marketing authorisation date:

21/12/2012

Manufacturing sites for batch release:

Laboratorios Karizoo S.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 31223/4001

Date of authorisation status change:

10/01/2018

Reference member state:

Spain

Procedure number:

ES/V/0178/001

Concerned member states:

Germany Hungary Lithuania Netherlands Poland Portugal Romania
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents