

KARIDOX 500 mg/g

Authorised

- Doxycycline hyclate

Product identification

Medicine name:

KARIDOX 500 mg/g

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:

Spain

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:

19/11/2012

Manufacturing sites for batch release:

Laboratorios Karizoo S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

2675 ESP

Date of authorisation status change:

21/11/2012

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

English (PDF)

Published on: 6/04/2023

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Combined File of all Documents

Package Leaflet

English (PDF)

Published on: 5/06/2025

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Labelling

English (PDF)

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