

# KARIDOX 500 mg/g

Authorised

- Doxycycline hyclate

## Product identification

**Medicine name:**

KARIDOX 500 mg/g

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**Active substance:**

Doxycycline hyclate

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**Target species:**

Chicken (for reproduction)

Chicken (broiler)

Pig (for fattening)

Turkey (for reproduction)

Turkey (for meat production)

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**Route of administration:**

In drinking water use

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## Product details

**Active substance and strength:**

Doxycycline hyclate

580.00 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Powder for use in drinking water

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01AA02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Lithuania

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**Package description:**

bag of 1 kg

bag of 200 g

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Karizoo S.A.

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**Marketing authorisation date:**

29/11/2012

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**Manufacturing sites for batch release:**

Laboratorios Karizoo S.A.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/12/2147/001-002

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**Date of authorisation status change:**

28/05/2025

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0178/001

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**Concerned member states:**

Germany Hungary Lithuania Netherlands Poland Portugal Romania

United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

English (PDF)

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