

# K-DOX 500 mg/g POLVO PARA ADMINISTRACION EN AGUA DE BEBIDA O EN LECHE PARA PORCINO, POLLOS Y TERNEROS

Authorised

- Doxycycline hyclate

## Product identification

### Medicine name:

K-DOX 500 mg/g POLVO PARA ADMINISTRACION EN AGUA DE BEBIDA O EN LECHE PARA PORCINO, POLLOS Y TERNEROS

### Active substance:

Doxycycline hyclate

### Target species:

Cattle (pre-ruminant)

Pig (for fattening)

Chicken (broiler)

### Route of administration:

In drinking water/milk use

## Product details

### **Active substance and strength:**

Doxycycline hyclate

500.00 milligram(s) / 1.00 gram(s)

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### **Pharmaceutical form:**

Powder for use in drinking water/milk

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### **Withdrawal period by route of administration:**

#### **In drinking water/milk use:**

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#### **Cattle (pre-ruminant)**

- Meat and offal. 7 day

- Milk. no withdrawal period

Leche: No autorizado en animales en lactación cuya leche se utiliza para el consumo humano

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#### **Pig (for fattening)**

- Meat and offal. 2 day

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#### **Chicken (broiler)**

- Meat and offal. 7 day

- Eggs. no withdrawal period

Huevos: No autorizado en aves ponedoras cuyos huevos se utilizan para el consumo humano. No usar en un plazo de 4 semanas desde el inicio de la puesta.

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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:**

Spain

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**Package description:**

Available only in [Spanish](#)

Available only in [Spanish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Karizoo S.A.

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**Marketing authorisation date:**

5/06/2013

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**Manufacturing sites for batch release:**

Laboratorios Karizoo S.A.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

2819 ESP

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**Date of authorisation status change:**

6/06/2013

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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

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Labelling

This document does not exist in this language (English). You can find it in another language below.

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