

AMOXICILLIN GLOBAL VET HEALTH 500 mg/g, powder for use in drinking water for chickens, turkeys, ducks and pigs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

AMOXICILLIN GLOBAL VET HEALTH 500 mg/g, powder for use in drinking water for chickens, turkeys, ducks and pigs

Amoksyacylina Global Vet Health 500 mg/g Proszek do podania w wodzie do picia

Active substance:

Amoxicillin trihydrate

Target species:

Turkey

Chicken

Duck

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate

500.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

Oral use:

-

Turkey

- Meat and offal. 5 day

-

Chicken

- Meat and offal. 1 day

-

Duck

- Meat and offal. 9 day

-

Pig

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 100 g.

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 200 g.

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 500 g.

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 1 kg.

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:

Global Vet Health S.L.

Marketing authorisation date:

1/07/2016

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2544

Date of authorisation status change:

1/07/2016

Reference member state:

Ireland

Procedure number:

IE/V/0350/001

Concerned member states:

Bulgaria Cyprus France Greece Italy Malta Poland Portugal Romania Spain
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: 13/04/2025

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Package Leaflet

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Labelling

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