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
Amoksycylina 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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