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Powdox 500 mg/g

- Doxycycline hyclate

Authorised

Product identification

Medicine name:
Powdox 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
  - Chicken (for reproduction)
    - Meat and offal
      5

day
        - Eggs
          no withdrawal period
          Eggs: Not authorised for use in laying birds producing eggs for human consumption
    - Chicken (broiler)
        - Meat and offal
          5
          day
        - Eggs
          no withdrawal period
          Eggs: Not authorised for use in laying birds producing eggs for human consumption
    - Pig (for fattening)
        - Meat and offal
          4
          day
    - Turkey (for reproduction)
        - Meat and offal
          12
          day
        - Eggs
          no withdrawal period
          Eggs: Not authorised for use in laying birds producing eggs for human consumption
    - Turkey (for meat production)
        - Meat and offal
          12
          day
        - Eggs
          no withdrawal period
          Eggs: Not authorised for use in laying 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:

- Germany

Available in:

- Germany

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:

- Vetpharma Animal Health S.L.

Marketing authorisation date:

- 5/12/2012

Manufacturing sites for batch release:

- Laboratorios Karizoo S.A.

Responsible authority:

- Federal Office Of Consumer Protection And Food Safety

Authorisation number:

- 401566.00.00

Date of authorisation status change:

- 22/02/2018

Reference member state:

- Spain

Procedure number:

- ES/V/0177/001

Concerned member states:

- Austria
- Bulgaria
- Denmark

- Germany
- Greece
- Hungary
- Ireland
- Italy
- Poland
- Portugal
- United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Product information

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Other languages (1)
German (PDF)
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Package Leaflet

English (PDF)
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