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KARIDOX 500 mg/g

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
KARIDOX 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 birds producing eggs for human consumption

- **Chicken (broiler)**
  - Meat and offal. 5 day
  - Eggs. no withdrawal period

Eggs: Not authorised for use in 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 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 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

Package description:
bag of 200 g
bag 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:
Laboratorios Karizoo S.A.

Marketing authorisation date:
13/12/2012

Manufacturing sites for batch release:
Laboratorios Karizoo S.A.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401567.00.00

Date of authorisation status change:
27/03/2018

Reference member state:

Spain

Procedure number:
ES/V/0178/001

Concerned member states:
Germany Hungary Lithuania Netherlands Poland Portugal Romania United Kingdom (Northern Ireland)

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

Documents

Labelling

Combined File of all Documents

English (PDF)
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