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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

AMOXICILLIN GLOBAL VET HEALTH 500 mg/g, pulbere pentru utilizare în apa de băut pentru pui de găină, curci, rațe și porci

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:

Available in:

Romania

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:

11/11/2015

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

200218

Date of authorisation status change:

R	efe	ran	6	m	am	har	cta	ta:
П	CIC	CII	CC	1117		NCI	3 LC	ILC.

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

Combined File of all Documents

English (PDF)

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