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MOXAPULVIS 500 mg/g powder for use in drinking water

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
MOXAPULVIS 500 mg/g powder for use in drinking water

Active substance:
Amoxicillin trihydrate

Target species:
Turkey
Chicken
Duck
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Amoxicillin trihydrate
574.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:
In drinking water use:

- **Turkey**
  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 5 day

- **Chicken**
  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 1 day

- **Duck**
  - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - 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:
Netherlands

Package description:
Round, white HDPE jars that are closed by a polypropylene lid with acardboard/aluminium/PE inner-layer.Pack size: 1 kg jar
Multi-layer laminated bag (Polyester/aluminium foil/polyethylene).Pack size: 1 kg bag
Round, white HDPE jars that are closed by a polypropylene lid with acardboard/aluminium/PE inner-layer.Pack size: 100 g jar

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
19/03/2018

Manufacturing sites for batch release:
V.M.D.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 120573

Date of authorisation status change:
1/02/2022

Reference member state:
Ireland

Procedure number:
IE/V/0541/001

Concerned member states:
Belgium Bulgaria Croatia Estonia France Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics