AMOXICILLINA TRIIDRATO 80%, 800 mg/g, milteliai naudoti su geriamuoju vandeniu vištoms, kalakutams ir kiaulėms

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
AMOXICILLINA TRIIDRATO 80%, 800 mg/g, milteliai naudoti su geriamuoju vandeniu vištoms, kalakutams ir kiaulėms

Active substance:
Amoxicillin trihydrate

Target species:
Chicken
Turkey
Pig

Route of administration:
In drinking water use
In-feed use

Product details

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

Pharmaceutical form:
Oral powder

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

- **Chicken**
  - Meat. 1 day

Not authorized for use in birds, whose eggs are intended for human consumption.

- **Turkey**
  - Meat. 2 day

Not authorized for use in birds, whose eggs are intended for human consumption.

- **Pig**
  - Meat. 1 day

In-feed use:

- **Chicken**
  - Meat. 1 day

Not authorized for use in birds, whose eggs are intended for human consumption.

- **Turkey**

- Meat. 2 day

Not authorized for use in birds, whose eggs are intended for human consumption.

- **Pig**
  - Meat. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Chemifarma S.p.A.

Marketing authorisation date:
29/06/2010

Manufacturing sites for batch release:
Chemifarma S.p.A.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/10/1938/001-005

Date of authorisation status change:
17/06/2015

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

Documents

Combined File of all Documents

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