Vetrimoxin 50, 50 g/100 g, proszek do sporządzania roztworu doustnego dla świń, bydła, kur i indyków

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Vetrimoxin 50, 50 g/100 g, proszek do sporządzania roztworu doustnego dla świń, bydła, kur i indyków

Active substance:
Amoxicillin trihydrate

Target species:
Chicken (hen)
Cattle
Pig
Turkey

Route of administration:
Oral use

Product details

Active substance and strength:
Amoxicillin trihydrate
50.00 gram(s) / 100.00 gram(s)

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:
Oral use:

- **Chicken (hen)**
  - Meat and offal. 2 day
- **Cattle**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 7 day
- **Turkey**
  - Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:

Poland

Package description:
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
16/10/2007

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1761

Date of authorisation status change:
16/10/2007

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.