VETRIMOXIN 500mg/g, pulveris iekšķīgi lietojama šķīduma pagatavošanai

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
VETRIMOXIN 500mg/g, pulveris iekšķīgi lietojama šķīduma pagatavošanai

Active substance:
Amoxicillin trihydrate

Target species:
Pig
Cattle (suckling calf)
Poultry

Route of administration:
In drinking water/milk use

Product details

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

Pharmaceutical form:
Powder for oral solution

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

- **Pig**
  - Meat and offal. 7 day
- **Cattle (suckling calf)**
  - Meat and offal. 1 day
- **Poultry**
  - Meat and offal. 3 day

Nav reģistrēts lietošanai putniem, no kuriem iegūst olas lietošanai pārtikā.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
7/11/2003

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/03/1588

Date of authorisation status change:
9/11/2003

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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