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UNI-AMOXICILIN, 510.2mg/g, Prášek pro perorální roztok

Not authorised

- Amoxicillin

Product identification

Medicine name:
UNI-AMOXICILIN, 510.2mg/g, Prášek pro perorální roztok

Active substance:
Amoxicillin

Target species:
Pig (piglet)
Chicken (broiler)

Route of administration:
In drinking water use

Product details

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

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:

In drinking water use:

- **Pig (piglet)**
  - Meat and offal. 7 day
- **Chicken (broiler)**
  - Meat and offal. 1 day
  - Egg. no withdrawal period

Nepoužívat u nosnic, jejichž vejce jsou určena pro lidskou spotřebu.,

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Czechia

Package description:

Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

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:
Univit s.r.o.

Marketing authorisation date:
25/03/2003

Manufacturing sites for batch release:
Univit s.r.o.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/026/03-C

Date of authorisation status change:
3/01/2024

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.