

VETRIMOXIN 500 mg/g prášok na perorálny roztok

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

VETRIMOXIN 500 mg/g prášok na perorálny roztok

Active substance:

Amoxicillin trihydrate

Target species:

Pig

Chicken (broiler)

Cattle (pre-ruminant)

Route of administration:

In drinking water 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 use:

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Pig

- Meat and offal. 7 day

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Chicken (broiler)

- Meat and offal. 3 day

Do not feed in laying hens producing eggs for human consumption.

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Cattle (pre-ruminant)

- Meat and offal. 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:

Slovakia

Package description:

Available only in Slovak

Available only in Slovak

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Animal Health Slovakia s.r.o.

Marketing authorisation date:

30/04/2004

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/109/04-S

Date of authorisation status change:

30/04/2004

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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