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Amix vet D 500 mg/g prášok na perorálny roztok

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
Amix vet D 500 mg/g prášok na perorálny roztok

Active substance:
Doxycycline hyclate

Target species:
Pig

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

Product details

Active substance and strength:
Doxycycline hyclate
570.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:

In drinking water use:

- **Pig**
  - Meat and offal. 8 day

In-feed use:

- **Pig**
  - Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
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:
Cymedica spol. s r.o.

Marketing authorisation date:
31/10/2007

Manufacturing sites for batch release:
Cymedica spol. s r.o.

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

Authorisation number:
96/061/07-S

Date of authorisation status change:
31/10/2007

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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