Suramox 500 mg/g pulveris šķīduma iekšķīgai lietošanai pagatavošanai cūkām

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Suramox 500 mg/g pulveris šķīduma iekšķīgai lietošanai pagatavošanai cūkām

Active substance:
Amoxicillin trihydrate

Target species:
Pig

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

- **Pig**
  - Meat and offal. 14 day

In drinking water use:

- **Pig**
  - Meat and offal. 14 day

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
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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:
Virbac

Marketing authorisation date:
18/02/2008

Manufacturing sites for batch release:
Virbac

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/08/1716

Date of authorisation status change:
18/02/2008

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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