Amoxycilline 70 % Kela 700 mg/g Poeder voor toediening in het drinkwater/in de melk

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Amoxycilline 70 % Kela 700 mg/g Poeder voor toediening in het drinkwater/in de melk
Amoxycilline 70 % Kela 700 mg/g Poudre pour administration dans le lait ou l'eau de boisson
Amoxycilline 70 % Kela 700 mg/g Pulver zum Eingeben über das Trinkwasser/die Milch

Active substance:
Amoxicillin trihydrate

Target species:
Chicken
Pig
Cattle (calf)

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:

Amoxicillin trihydrate
817.70 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

Withdrawal period by route of administration:

In drinking water/milk use:

- **Chicken**

  - Meat and offal. 1 day

  - Egg. no withdrawal period

Not for use in chickens producing eggs for human consumption

- **Pig**

  - Meat and offal. 2 day

- **Cattle (calf)**

  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 2000 g powder
Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 1429 g powder

Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 1000 g powder
Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 714 g powder
Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 500 g powder
Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 143 g powder
Amoxycilline 70 % Kela 700 mg/g Powder for use in drinking water/milk Bag with 100 g powder

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Kela Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
2/04/2002

Manufacturing sites for batch release:
Kela - Kempisch Laboratorium - Kela Laboratoria

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V233764

Date of authorisation status change:
2/04/2002

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

Documents

Package Leaflet

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Summary of Product Characteristics

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