Tylosine 75 % Kela 750 mg/g Poeder voor gebruik in drinkwater

Authorised

- Tylosin tartrate

Product identification

Medicine name:
Tylosine 75 % Kela 750 mg/g Poeder voor gebruik in drinkwater
Tylosine 75 % Kela 750 mg/g Poudre pour administration dans l’eau de boisson
Tylosine 75 % Kela 750 mg/g Pulver zum Eingeben über das Trinkwasser

Active substance:
Tylosin tartrate

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Tylosin tartrate
825.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:
In drinking water use:

- **Chicken**
  - Meat and offal. 1 day
  - Egg. no withdrawal period

Do not use in animals producing eggs for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FA90

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Tylosine 75 % Kela 1 Bag with 2000 g of Powder for use in drinking water
Tylosine 75 % Kela 1 Bag with 1333 g of Powder for use in drinking water
Tylosine 75 % Kela 1 Bag with 1000 g of Powder for use in drinking water
Tylosine 75 % Kela 1 Bag with 667 g of Powder for use in drinking water
Tylosine 75 % Kela 1 Bag with 500 g of Powder for use in drinking water
Tylosine 75 % Kela 1 Bag with 133 g of Powder for use in drinking water
Tylosine 75 % Kela 1 Bag with 100 g of Powder for use in drinking water

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:
28/08/2000

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V216422

Date of authorisation status change:
15/12/2021

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

Documents

Summary of Product Characteristics

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

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.