CENAMUTIN 125 mg/ml SOLUCION PARA ADMINISTRACION EN AGUA DE BEBIDA

Authorised

- Tiamulin hydrogen fumarate

Product identification

Medicine name:
CENAMUTIN 125 mg/ml SOLUCION PARA ADMINISTRACION EN AGUA DE BEBIDA

Active substance:
Tiamulin hydrogen fumarate

Target species:
Pig
Chicken (layer hen)
Chicken (for reproduction)
Chicken (chick, for replacement)
Chicken (broiler)
Turkey (for reproduction)
Turkey (for meat production)

Route of administration:
In drinking water use

Product details

Active substance and strength:
Tiamulin hydrogen fumarate
125.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

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

- **Pig**
  - Meat and offal. 4 day
- **Chicken (layer hen)**
  - Meat and offal. 6 day
  - Eggs. 0 day
- **Chicken (for reproduction)**
  - Meat and offal. 6 day
  - Eggs. 0 day
- **Chicken (chick, for replacement)**
  - Meat and offal. 6 day
  - Eggs. 0 day
- **Chicken (broiler)**
  - Meat and offal. 6 day
  - Eggs. 0 day
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Turkey (for reproduction)

- Meat and offal. 6 day
- Eggs. no withdrawal period

Huevos:Su uso no está autorizado en aves cuyos huevos se utilizan para el consumo humano

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Turkey (for meat production)

- Meat and offal. 6 day
- Eggs. no withdrawal period

Huevos:Su uso no está autorizado en aves cuyos huevos se utilizan para el consumo humano

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01XQ01

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Cenavisa S.L.

Marketing authorisation date:
3/02/2009

Manufacturing sites for batch release:
Cenavisa S.L.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
1979 ESP

Date of authorisation status change:
3/02/2009

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

Documents

Labelling

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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.

Package Leaflet

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