PRIMUN SALMONELLA T Lyophilisate for use in drinking water for chickens

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain ST CAL 16 Str+/Rif+/Enr-, Live

Product identification

Medicine name:
PRIMUN SALMONELLA T Lyophilisate for use in drinking water for chickens
Primun Salmonella T, Lyofilizát pro podání v pitné vodě

Active substance:
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain ST CAL 16 Str+/Rif+/Enr-, Live

Target species:
Chicken (pullet for egg production, future layer)
Future breeder pullet

Route of administration:
In drinking water use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain ST CAL 16 Str+/Rif+/Enr-, Live

100000000.00 Colony forming unit / 1.00 Dose

Pharmaceutical form:

Lyophilisate for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Chicken (pullet for egg production, future layer)**

  - Meat and offal. no withdrawal period

28 days after first and second vaccination and 14 days after third vaccination.

  - Eggs. no withdrawal period

Do not use in birds in lay and within 4 weeks before the start of the laying period.

- **Future breeder pullet**

  - Meat and offal. no withdrawal period

28 days after first and second vaccination and 14 days after third vaccination.

  - Eggs. no withdrawal period

Do not use in birds in lay and within 4 weeks before the start of the laying period.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Cardboard box with 1 vial (20 ml) of 1000 doses
Cardboard box with 10 vials (20 ml) of 1000 doses
Cardboard box with 1 vial (20 ml) of 2000 doses
Cardboard box with 10 vials (20 ml) of 2000 doses

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:
Laboratorios Calier S.A.

Marketing authorisation date:
23/02/2023

Manufacturing sites for batch release:
Laboratorios Calier S.A.

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

Authorisation number:
97/010/23-C

Date of authorisation status change:
23/02/2023

Reference member state:
Spain

Procedure number:
ES/V/0408/001

Concerned member states:
Belgium Czechia France Germany Greece Hungary Italy Lithuania Netherlands Poland Portugal Romania

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 24/03/2023
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Package Leaflet

English (PDF)
Published on: 24/03/2023
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Labelling

English (PDF)
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