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 liofilizat do podania w wodzie do picia dla kur
PRIMUN SALMONELLA T Lyophilisate for use in drinking water for chickens

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:
Poland

Package description:

Cardboard box with 10 vials (20 ml) of 2000 doses
Cardboard box with 1 vial (20 ml) of 2000 doses
Cardboard box with 1 vial (20 ml) of 2000 doses
Cardboard box with 1 vial (20 ml) of 1000 doses
Cardborard box with 1 vial (20 ml) of 4000 doses
Cardborard box with 10 vial (20 ml) of 4000 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:
25/11/2022

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3212

Date of authorisation status change:
25/11/2022

Reference member state:
Spain

Procedure number:
ES/V/0408/001

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

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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.