

# 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

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**Active substance:**

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

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**Target species:**

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

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**Route of administration:**

In drinking water use

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### 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

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**Pharmaceutical form:**

Lyophilisate for use in drinking water

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**Withdrawal period by route of administration:**

**In drinking water use:**

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**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.

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**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.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AE01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Germany

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**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 10 vials (20 ml) of 1000 doses

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

Cardboard box with 10 vial (20 ml) of 4000 doses

Cardboard box with 1 vial (20 ml) of 4000 doses

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## Additional information

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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### **Marketing authorisation holder:**

Laboratorios Calier S.A.

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### **Marketing authorisation date:**

13/07/2022

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### **Manufacturing sites for batch release:**

Laboratorios Calier S.A.

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### **Responsible authority:**

Paul-Ehrlich-Institut

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### **Authorisation number:**

PEI.V.12106.01.1

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### **Date of authorisation status change:**

13/07/2022

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### **Reference member state:**

Spain

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### **Procedure number:**

ES/V/0408/001

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**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](http://www.adrreports.eu/vet)

## Documents

### Labelling

English (PDF)

Published on: 24/03/2023

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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.

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