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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

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:
Germany

Package description:

Cardborard box with 1 vial (20 ml) of 4000 doses
Cardborard box with 10 vial (20 ml) of 4000 doses
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:
13/07/2022

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.12106.01.1

Date of authorisation status change:
13/07/2022

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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Documents

Labelling

English (PDF)
Published on: 14/03/2026
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Summary of Product Characteristics

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Package Leaflet

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

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