

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:

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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.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

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 1 vial (20 ml) of 4000 doses

Cardboard 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:

2/04/2023

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

European Medicines Agency

Authorisation number:

LT/2/23/2741/001-004

Date of authorisation status change:

2/04/2023

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

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

Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

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