PRIMUN SALMONELLA E

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain CAL10 Sm+/Rif+/Ssq-, Live

Product identification

Medicine name:
PRIMUN SALMONELLA E

Active substance:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain CAL10 Sm+/Rif+/Ssq-, 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 Enteritidis, strain CAL10 Sm+/Rif+/Ssq-, Live
600000000.00 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

Meat and offal: 21 days after 1st, 2nd and 3rd vaccination. 14 days after 4th vaccination; Eggs: Zero days after 4th vaccination.

  - Eggs. no withdrawal period

Meat and offal: 21 days after 1st, 2nd and 3rd vaccination. 14 days after 4th vaccination; Eggs: Zero days after 4th vaccination.

- **Future breeder pullet**

  - Meat and offal. no withdrawal period

Meat and offal: 21 days after 1st, 2nd and 3rd vaccination. 14 days after 4th vaccination; Eggs: Zero days after 4th vaccination.

  - Eggs. no withdrawal period

Meat and offal: 21 days after 1st, 2nd and 3rd vaccination. 14 days after 4th vaccination; Eggs: Zero days after 4th vaccination.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AE01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Package description:
cardboard box containing 10 vials (20 ml) of 2.000 doses

cardboard box containing 10 vials (20 ml) of 1.000 doses
cardboard box containing 1 vial (20 ml) of 2.000 doses
cardboard box containing 1 vial (20 ml) of 1.000 doses
Cardboard box containing 10 vials (20 ml) of 4000 doses
Cardboard box containing 1 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:
22/05/2023

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8799020 9/2023

Date of authorisation status change:
22/05/2023

Reference member state:
Spain

Procedure number:
ES/V/0218/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia France Germany Greece Hungary Ireland 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

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Package Leaflet

Labelling