PRIMUN SALMONELLA E

Authorised

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

Product identification

Medicine name:
PRIMUN SALMONELLA E
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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:
Germany

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:
7/09/2015

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11784.01.1

Date of authorisation status change:
16/01/2020

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

Summary of Product Characteristics

English (PDF)
Published on: 11/04/2023
Download

Package Leaflet

Labelling