AviPro SALMONELLA VAC E Lyophilisate for suspension for chickens

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain Sm24/Rif12/Ssq, Live

Product identification

Medicine name:
AviPro SALMONELLA VAC E Lyophilisate for suspension for chickens
AVIPRO SALMONELLA VAC E LYOPHILISAT POUR ADMINISTRATION DANS L'EAU DE BOISSON

Active substance:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain Sm24/Rif12/Ssq, Live

Target species:
Chicken
Chicken (chick)

Route of administration:
In drinking water use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain Sm24/Rif12/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**
  - Egg. 0 day
  - Meat and offal. 21 day
- **Chicken (chick)**
  - Egg. 0 day
  - Meat and offal. 21 day

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:
(ID6): 1 Box with 10 Vials (Glass) with 5000 vaccine doses (50000 vaccine doses)
(ID5): 1 Box with 1 Vial (Glass) with 5000 vaccine doses (5000 vaccine doses)
(ID4): 1 Box with 10 Vials (Glass) with 2000 vaccine doses (20000 vaccine doses)
(ID3): 1 Box with 1 Vial (Glass) with 2000 vaccine doses (2000 vaccine doses)
(ID2): 1 Box with 10 Vials (Glass) with 1000 vaccine doses (10000 vaccine doses)
(ID1): 1 Box with 1 Vial (Glass) with 1000 vaccine doses (1000 vaccine doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Elanco GmbH

Marketing authorisation date:
10/02/2009

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

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

Authorisation number:
FR/V/8083355 2/2009

Date of authorisation status change:
22/11/2017

Reference member state:
Germany

Procedure number:
DE/V/0215/001

Concerned member states:
Austria Belgium Estonia France Greece Italy Latvia Lithuania Netherlands Portugal Slovenia Spain

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/03/2022
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Package Leaflet and Labelling

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