

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 liofilizado para suspensão para galinhas

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

6000000000.00 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

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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:

Portugal

Available in:

Portugal

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:

14/01/2002

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

R693/02 DGV

Date of authorisation status change:

19/01/2023

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

Combined File of all Documents

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