

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

Active substance:

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

Target species:

Chicken

Route of administration:

In drinking water use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain Sm24/Rif12/Ssq, 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:**

-

Chicken

- 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:

Germany

Available in:

Germany

Package description:

(ID1) 1000 Dose: Box (cardboard) with 1 Vial (Glass type I) with 1000 Dose, closed with Stopper (rubber) and Kappe (aluminium)

(ID2) 10000 Dose: Box (cardboard) with 10 Vial (Glass type I) each with 1000 Dose, closed with Stopper (rubber) and Kappe (aluminium)

(ID3) 2000 Dose: Box (cardboard) with 1 Vial (Glass type I) with 2000 Dose, closed with Stopper (rubber) and Kappe (aluminium)

(ID4) 20000 Dose: Box (cardboard) with 10 Vial (Glass type I) each with 2000 Dose, closed with Stopper (rubber) and Kappe (aluminium)

(ID5) 5000 Dose: Box (cardboard) with 1 Vial (Glass type I) with 5000 Dose, closed with Kappe (aluminium) and Stopper (rubber)

(ID6) 50000 Dose: Box (cardboard) with 10 Vial (Glass type I) each with 5000 Dose, closed with Stopper (rubber) and Kappe (aluminium)

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:

Lohmann Animal Health GmbH

Marketing authorisation date:

21/07/1999

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

85a/97

Date of authorisation status change:

31/05/2007

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

Combined File of all Documents

English (PDF)

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