

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:

Italy

Available in:

Italy

Package description:

(ID6) 50000 Dose: Box (cardboard) with 10 Vial (Glass type I) each with 5000 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)

(ID4) 20000 Dose: Box (cardboard) with 10 Vial (Glass type I) each with 2000 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)

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

(ID1) 1000 Dose: Box (cardboard) with 1 Vial (Glass type I) with 1000 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:

Elanco GmbH

Marketing authorisation date:

10/05/2007

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Ministry Of Health

Authorisation number:

103859

Date of authorisation status change:

10/05/2007

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

Summary of Product Characteristics

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

Combined File of all Documents

English (PDF)

Published on: 13/02/2026

[Download](#)