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# 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 60000000.00 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:

**OI01AE01** 

#### 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.