AviPro SALMONELLA DUO Lyophilisate for suspension

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain Sm24/Rif12/Ssq, Live
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain Nal2/Rif9/Rtt, Live

Product identification

Medicine name:
AviPro SALMONELLA DUO Lyophilisate for suspension
AVIPRO SALMONELLA DUO

Active substance:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain Sm24/Rif12/Ssq, Live
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain Nal2/Rif9/Rtt, Live

Target species:
Turkey
Chicken (for reproduction)
Duck

Route of administration:
In drinking water/milk 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

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain Nal2/Rif9/Rtt, Live 600000000.00 Colony forming unit / 1.00 Dose

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Pharmaceutical form:

Lyophilisate for use in drinking water

Withdrawal period by route of administration:

In drinking water/milk use:

- **Turkey**

  - Meat and offal. 70 day

70 days after first vaccination, 49 days after further vaccination

- **Chicken (for reproduction)**

  - Meat and offal. 21 day

  - Egg. 21 day

- **Duck**

  - Meat and offal. 21 day

  - Egg. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

Package description:

(ID4): 1 Box with 10 Vials (Glass) with 2000 vaccine doses (20000 vaccine doses)

(ID3): 1 Box with 10 Vials (Glass) with 1000 vaccine doses (10000 vaccine doses)
(ID2): 1 Box with 1 Vial (Glass) with 2000 vaccine doses (2000 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:
22/03/2012

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
104300

Date of authorisation status change:
22/03/2012

Reference member state:
Germany

Procedure number:
DE/V/0249/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia France Greece Hungary

Italy Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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Combined File of all Documents

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