# AviPro AE suspension for use in drinking water

Authorised

- Avian encephalomyelitis virus, strain Calnek 1143, Live

## Product identification

**Medicine name:**
AviPro AE suspension for use in drinking water
AviPro AE Suspension zur Anwendung über das Trinkwasser für Hühner

**Active substance:**
Avian encephalomyelitis virus, strain Calnek 1143, Live

**Target species:**
Chicken

**Route of administration:**
In drinking water use

## Product details

**Active substance and strength:**
Avian encephalomyelitis virus, strain Calnek 1143, Live
31622.80 50% Embryo Infective Dose / 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. 0 day
  - Egg. 0 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AD02

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**

Valid

**Authorised in:**

Germany

**Package description:**

(ID8): 1 Box with 10 Bottle (Glass) with 10000 Dose (100000.0 Dose)
(ID7): 1 Box with 1 Bottle (Glass) with 10000 Dose (10000.0 Dose)
(ID6): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000.0 Dose)
(ID5): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000.0 Dose)
(ID4): 1 Box with 10 Bottle (Glass) with 2500 Dose (25000.0 Dose)
(ID3): 1 Box with 1 Bottle (Glass) with 2500 Dose (2500.0 Dose)
(ID2): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000.0 Dose)
(ID1): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000.0 Dose)

## 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:**
94a/89

**Date of authorisation status change:**
8/06/2009

**Reference member state:**
Germany

**Procedure number:**
DE/V/0292/001

**Concerned member states:**
Austria France 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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