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 POUR ADMINISTRATION DANS L'EAU DE BOISSON DES POULES

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:
France

Available in:
France

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:
Elanco GmbH

Marketing authorisation date:
13/01/2022

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6803906 6/2021

Date of authorisation status change:
13/01/2022

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

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

Package Leaflet and Labelling

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