AviPro IB – ND C131 Lyophilisate for oculonasal suspension/use in drinking water for chicken

Authorised

- Newcastle disease virus, strain Clone 13-1, Live
- Infectious bronchitis virus, type Massachusetts, strain H120, Live

Product identification

Medicine name:
AviPro IB – ND C131 Lyophilisate for oculonasal suspension/use in drinking water for chicken

Active substance:
Newcastle disease virus, strain Clone 13-1, Live
Infectious bronchitis virus, type Massachusetts, strain H120, Live

Target species:
Chicken
Chicken (layer hen)

Route of administration:
Oculonasal use
In drinking water use

Product details

Active substance and strength:

Newcastle disease virus, strain Clone 13-1, Live
316227.00 50% Embryo Infective Dose / 1.00 Dose

Infectious bronchitis virus, type Massachusetts, strain H120, Live
2511.00 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for suspension

Withdrawal period by route of administration:

Oculonasal use:

- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day
- **Chicken (layer hen)**
  - Egg. 0 day
  - Meat and offal. 0 day

In drinking water use:

- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day
- **Chicken (layer hen)**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD11

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
(ID1) 2000 Dose: Box (Cardboard) with 1 Bottle (Glass) with 2000 Dose, closed with Lid and Stopper (Aluminium, Rubber)
(ID2) 20000 Dose: Box (Cardboard) with 10 Bottle (Glass) each with 2000 Dose, closed with Lid and Stopper (Aluminium, Rubber)
(ID3) 5000 Dose: Box (Cardboard) with 1 Bottle (Glass) with 5000 Dose, closed with Lid and Stopper (Aluminium, Rubber)
(ID4) 50000 Dose: Box (Cardboard) with 10 Bottle (Glass) each with 5000 Dose, closed with Lid and Stopper (Aluminium, Rubber)

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:
28/06/2023

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 128447

Date of authorisation status change:
28/06/2023

Reference member state:
Germany

Procedure number:
DE/V/0291/001

Concerned member states:
Austria Czechia France Hungary Italy Netherlands Poland Portugal Romania 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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