

AviPro IB – ND C131 Lyophilisate for ocular nasal 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 ocular nasal 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:

Ocular nasal 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:

Poland

Available in:

Poland

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:

22/08/2023

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3269

Date of authorisation status change:

22/08/2023

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

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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

Combined File of all Documents

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