

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:

Hungary

Package description:

(ID4) 50000 Dose: Box (Cardboard) with 10 Bottle (Glass) each with 5000 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)

(ID2) 20000 Dose: Box (Cardboard) with 10 Bottle (Glass) each with 2000 Dose, closed with Lid and Stopper (Aluminium, Rubber)

(ID1) 2000 Dose: Box (Cardboard) with 1 Bottle (Glass) with 2000 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:

12/10/2023

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

4385/x/23 NÉBIH ÁTI

Date of authorisation status change:

12/10/2023

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