Source URL: https://medicines.health.europa.eu/veterinary/en/600000062394

AviPro ND C131 Lyophilisate for suspension

Authorised

• Newcastle disease virus, strain Clone 13-1, Live

Product identification

Medicine name:

AviPro ND C131 Lyophilisate for suspension AviPro ND C131 лиофилизат за суспензия за пилета и пуйки

Active substance:

Newcastle disease virus, strain Clone 13-1, Live

Target species:

Turkey

Chicken

Route of administration:

In drinking water use Topical Ocular use

Product details

Active substance and strength:

Newcastle disease virus, strain Clone 13-1, Live 15848900.00 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration: In drinking water use:

Turkey

- Meat and offal. 0 day
- Egg. 0 day

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Chicken

- Meat and offal. 0 day
- Egg. 0 day

Topical:

•

Chicken

- Meat and offal. 0 day
- Egg. 0 day

Ocular use:

•

Chicken

- Meat and offal. 0 day
- Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

(ID8): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000 Dose)

(ID7): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000 Dose)

(ID11): 1 Box with 1 Bottle (Glass) with 2000 Dose (2000 Dose)

(ID12): 1 Box with 10 Bottle (Glass) with 2000 Dose (20000 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:

31/10/2017

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2767

Date of authorisation status change:

14/03/2022

Reference member state:

Germany

Procedure number:

DE/V/0239/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Estonia France Greece Hungary Italy Latvia Lithuania Netherlands Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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