AviPro ND C131 Lyophilisate for suspension

Not authorised

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

Product identification

Medicine name:
AviPro ND C131 Lyophilisate for suspension
AviPro ND C131 Λυοφιλοποιημένο υλικό για εναιώρημα για όρνιθες και ινδόρνιθες

Active substance:
Newcastle disease virus, strain Clone 13-1, Live

Target species:
Turkey
Chicken

Route of administration:
In drinking water use
Topical use
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 suspension

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

- **Turkey**
  - Meat and offal. 0 day
  - Egg. 0 day
- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

Topical use:

- **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:
Surrendered

Authorised in:
Cyprus

Package description:
(ID2): 1 Box with 10 Bottle (Glass) with 500 Dose (5000 Dose)
(ID9): 1 Box with 1 Bottle (Glass) with 10000 Dose (10000 Dose)
(ID10): 1 Box with 10 Bottle (Glass) with 10000 Dose (100000 Dose)
(ID1): 1 Box with 1 Bottle (Glass) with 500 Dose (500 Dose)
(ID8): 1 Box with 10 Bottle (Glass) with 5000 Dose (50000 Dose)
(ID6): 1 Box with 10 Bottle (Glass) with 2500 Dose (25000 Dose)
(ID3): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 Dose)
(ID5): 1 Box with 1 Bottle (Glass) with 2500 Dose (2500 Dose)
(ID4): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)
(ID7): 1 Box with 1 Bottle (Glass) with 5000 Dose (5000 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:
18/01/2018

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00135V

Date of authorisation status change:
20/12/2021

Reference member state:
Germany

Procedure number:
DE/V/0239/001

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet