AviPro ND C131 Lyophilisate for suspension

Authorised

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

Product identification

Medicine name:
AVIPRO ND C131 ΚΟΝΙΣ ΓΙΑ ΕΝΑΙΩΡΗΜΑ
AviPro ND C131 Lyophilisate for suspension

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

Target species:
Turkey
Chicken

Route of administration:
In drinking water use
Nebulisation 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 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
- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

Nebulisation 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:
Valid

Authorised in:
Greece

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)
(ID12): 1 Box with 10 Bottle (Glass) with 2000 Dose (20000 Dose)
(ID11): 1 Box with 1 Bottle (Glass) with 2000 Dose (2000 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:
26/11/2007

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
National Organization For Medicines

Authorisation number:
36022/13-04-2022/K-0168301

Date of authorisation status change:
12/04/2022

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