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

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
1000000.00 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:
Italy

Package description:
(ID11) 2000 Dose: Box with 1 Bottle (Glass type I) with 2000 Dose, closed with Stopper (rubber) and Kappe (aluminium)
(ID12) 20000 Dose: Box with 10 Bottle (Glass type I) each with 2000 Dose, closed with Stopper (rubber) and Kappe (aluminium)
(ID7) 5000 Dose: Box with 1 Bottle (Glass type I) with 5000 Dose, closed with Stopper (rubber) and Kappe (aluminium)
(ID8) 50000 Dose: Box with 10 Bottle (Glass type I) each with 5000 Dose, closed with Stopper (rubber) and Kappe (aluminium)

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:
19/01/2010

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
103923

Date of authorisation status change:
19/01/2010

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

Combined File of all Documents

English (PDF)
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