

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 liofilizāts suspensijas pagatavošanai vistām un tītariem

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 ocularnasal 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

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:

Latvia

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:

21/05/2008

Manufacturing sites for batch release:

Lohmann Animal Health GmbH

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/08/1562

Date of authorisation status change:

21/05/2008

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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