Nobilis ND C2 lyophilisate for oculonasal suspension for chickens

Authorised

- Newcastle disease virus, strain C2, Live

Product identification

Medicine name:
Nobilis ND C2 lyophilisate for oculonasal suspension for chickens
Nobilis ND C2 liofilizat za okulonazalno suspenzijo za piščance

Active substance:
Newcastle disease virus, strain C2, Live

Target species:
Chicken

Route of administration:
Oculonasal use
Nebulisation use

Product details

Active substance and strength:
Newcastle disease virus, strain C2, Live
5.70 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oculonasal suspension

Withdrawal period by route of administration:

Oculonasal use:

- **Chicken**
  - All relevant tissues. no withdrawal period 0 days

Nebulisation use:

- **Chicken**
  - All relevant tissues. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Slovenia

Package description:
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian

Cardboard box with 10 vials of 1000 doses.
Cardboard box with 10 vials of 10000 doses.
Cardboard box with 10 vials of 2500 doses.
Cardboard box with 10 vials of 25000 doses.
Cardboard box with 10 vials of 5000 doses.
Cardboard box with 10 vials of 500 doses.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
10/01/2006

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0220/001

Date of authorisation status change:
10/01/2006

Reference member state:
Netherlands

Procedure number:
NL/V/0113/001

Concerned member states:
Belgium Cyprus Czechia Denmark Estonia Germany Greece Ireland Italy Latvia Lithuania Luxembourg Poland Portugal Slovakia Slovenia Spain Sweden 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.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents