NEMOVAC, lyofilizát pro přípravu orální suspenze, Lyofilizát pro orální suspenzi

Authorised

- Turkey rhinotracheitis virus, strain PL21, Live

Product identification

Medicine name:
NEMOVAC, lyofilizát pro přípravu orální suspenze, Lyofilizát pro orální suspenzi

Active substance:
Turkey rhinotracheitis virus, strain PL21, Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain PL21, Live
4.00 log10 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oral suspension

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD01

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

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
11/11/1999

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/064/99-C

Date of authorisation status change:
11/09/2009

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

Documents

Summary of Product Characteristics

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Package Leaflet

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.