NEMOVAC liofilizat pentru suspensie pentru administrare oculonazală/ în apa de băut pentru găini

Authorised

- Turkey rhinotracheitis virus, strain PL21, Live

Product identification

Medicine name:
NEMOVAC liofilizat pentru suspensie pentru administrare oculonazală/ în apa de băut pentru găini

Active substance:
Turkey rhinotracheitis virus, strain PL21, Live

Target species:
Chicken (broiler)
Chicken (for reproduction)
Chicken (pullet for egg production, future layer)

Route of administration:
Oral use
Nebulisation use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain PL21, Live
2.30 log10 50% cell culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:
Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:
Oral use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

- **Chicken (for reproduction)**
  - Meat and offal. 0 day

- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day
  - Eggs. 0 day

Nebulisation use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day
  - Eggs. 0 day

- **Chicken (for reproduction)**

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD01

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
25/07/2006

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
120092

Date of authorisation status change:
17/12/2025

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

Documents

Combined File of all Documents

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