NOBILIS RHINO CV

Authorised

- Turkey rhinotracheitis virus, strain 11/94, Live

Product identification

Medicine name:
NOBILIS RHINO CV

Active substance:
Turkey rhinotracheitis virus, strain 11/94, Live

Target species:
Chicken (layer hen)
Chicken (for reproduction)
Chicken (broiler)

Route of administration:
Nasal use
Ocular use
Nebulisation use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain 11/94, Live
1.50 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension

Withdrawal period by route of administration:

Nasal use:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day

- **Chicken (for reproduction)**
  - All relevant tissues. 0 day

- **Chicken (broiler)**
  - All relevant tissues. 0 day

Ocular use:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day

- **Chicken (for reproduction)**
  - All relevant tissues. 0 day

- **Chicken (broiler)**
  - All relevant tissues. 0 day

Nebulisation use:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day

- **Chicken (for reproduction)**
  - All relevant tissues. 0 day

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Chicken (broiler)

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Cardboard box containing 20 glass (Type I) vials of 25000 doses
Cardboard box containing 10 glass (Type I) vials of 25000 doses
Cardboard box containing 5 glass (Type I) vials of 25000 doses
Cardboard box containing 2 glass (Type I) vials of 25000 doses
Cardboard box containing 1 glass (Type I) vial of 25000 doses
Cardboard box containing 50 glass (Type I) vials of 10000 doses
Cardboard box containing 20 glass (Type I) vials of 10000 doses
Cardboard box containing 10 glass (Type I) vials of 10000 doses
Cardboard box containing 5 glass (Type I) vials of 10000 doses
Cardboard box containing 2 glass (Type I) vials of 10000 doses
Cardboard box containing 1 glass (Type I) vial of 10000 doses
Cardboard box containing 50 glass (Type I) vials of 5000 doses
Cardboard box containing 20 glass (Type I) vials of 5000 doses
Cardboard box containing 10 glass (Type I) vials of 5000 doses
Cardboard box containing 5 glass (Type I) vials of 5000 doses
Cardboard box containing 2 glass (Type I) vials of 5000 doses
Cardboard box containing 1 glass (Type I) vial of 5000 doses
Cardboard box containing 50 glass (Type I) vials of 2500 doses
Cardboard box containing 20 glass (Type I) vials of 2500 doses

Cardboard box containing 10 glass (Type I) vials of 2500 doses
Cardboard box containing 5 glass (Type I) vials of 2500 doses
Cardboard box containing 1 glass (Type I) vial of 2500 doses
Cardboard box containing 50 glass (Type I) vials of 1000 doses
Cardboard box containing 20 glass (Type I) vials of 1000 doses
Cardboard box containing 10 glass (Type I) vials of 1000 doses
Cardboard box containing 5 glass (Type I) vials of 1000 doses
Cardboard box containing 2 glass (Type I) vials of 1000 doses
Cardboard box containing 1 glass (Type I) vial of 1000 doses
Cardboard box containing 50 glass (Type I) vials of 500 doses
Cardboard box containing 20 glass (Type I) vials of 500 doses
Cardboard box containing 10 glass (Type I) vials of 500 doses
Cardboard box containing 5 glass (Type I) vials of 500 doses
Cardboard box containing 2 glass (Type I) vials of 500 doses
Cardboard box containing 1 glass (Type I) vial of 500 doses
Cardboard box containing 20 glass (Type I) vials of 250 doses
Cardboard box containing 10 glass (Type I) vials of 250 doses
Cardboard box containing 5 glass (Type I) vials of 250 doses
Cardboard box containing 2 glass (Type I) vials of 250 doses
Cardboard box containing 1 glass (Type I) vial of 250 doses
Cardboard box containing 2 glass (Type I) vials of 2500 doses
Cardboard box containing 50 glass (Type I) vials of 25000 doses
Cardboard box containing 50 glass (Type I) vials of 250 doses

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
2/05/2005

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.03207.01.1

Date of authorisation status change:
9/03/2010

Reference member state:
France

Procedure number:
FR/V/0151/001

Concerned member states:
Austria Belgium Cyprus Czechia Denmark Estonia Germany Greece Hungary Ireland Latvia Lithuania Luxembourg Netherlands Poland Portugal Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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